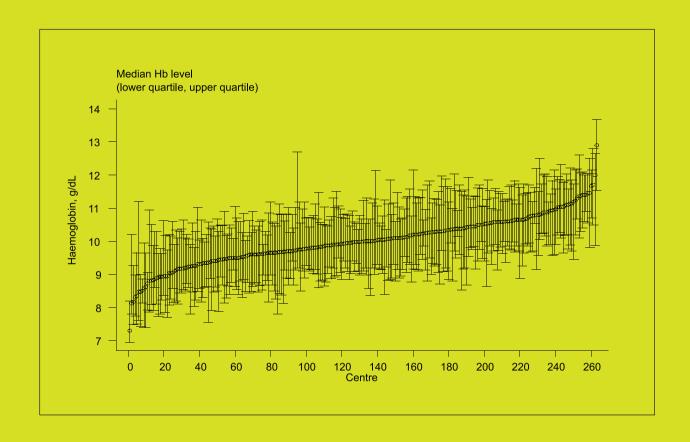
14th Report of The Malaysian Dialysis & Transplant Registry 2006



Edited by: Y N Lim, T O Lim

With contributions from:

Rozina G, Zaki Morad, Wong HS, Liu WJ, Lee ML, Philip N J, Ahmad Fauzi, Prasad M, Fan KS, Teo SM, Tan CC, Sunita B, Goh BL, Lee DG, Sharon Chen





14thReport OF THE MALAYSIAN DIALYSIS & TRANSPLANT REGISTRY 2006

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Malaysian Society of Nephrology Association of Dialysis Medical Assistants and Nurses

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Malaysian Society of Nephrology 2nd Floor, MMA House 124, Jalan Pahang 50286 Kuala Lumpur Malaysia

Telephone : (603) 4045 8636 Direct Fax : (603) 4042 7694 e-mail : nrr@msn.org.my

Web site : http://www.msn.org.my/nrr

Important information:

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Baxter Healthcare
Fresenius Medical Care
Roche

The staff of the Clinical Research Centre for IT and statistical support

The National Transplant Registry

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All who have in one way or another supported the National Renal Registry

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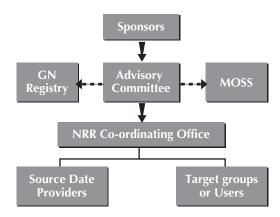
About the National Renal Registry

The National Renal Registry (NRR) has its origin in the Dialysis and Transplant Registry established by the Department of Nephrology in 1992. The sponsors of NRR are the Malaysian Society of Nephrology (MSN) and Association of Dialysis Medical Assistants and Nurses (ADMAN)

The objectives of NRR are to:

- 1. Determine the disease burden attributable to End Stage Renal Disease (ESRD), and its geographic and temporal trends in Malaysia.
- Determine the outcomes, and factors influencing outcomes of Renal Replacement Therapy.
- 3. Evaluate the RRT program.
- 4. Stimulate and facilitate research on RRT and ESRD.
- 5. Maintain the national renal transplant waiting list.

The NRR organization is as follows:



Sponsors

The Malaysian Society of Nephrology is the sponsor of the National Renal Registry (NRR) and Malaysian Organ Sharing System (MOSS) and the co-sponsor is the Association of Dialysis Medical Assistants and Nurses.

Advisory Committee

This is the committee established by the sponsors to oversee the operations of the registry and MOSS. Interested parties including source data producers, Renal Registry Unit and target groups or users are represented on this committee.

National Renal Registry Office

The NRR office is the coordinating center that collects and analyses the data. It publishes the annual report of Malaysian Dialysis & Transplant Registry and the Directory of Dialysis Centres in Malaysia. The Clinical Registry Manager (CRM) oversees the daily operation of the NRR. The Clinical Research Centre of Hospital Kuala Lumpur provides the epidemiological, statistical and information technological support to NRR.

Source Data Producers

These are the dialysis centres that collect the required data. It is the most critical and yet difficult element of the system. It has to be systematic and uniform, and producers of source data need to be trained and motivated to ensure high data quality.

Users or Target groups

These are the individuals or institutions to whom the regular registry reports are addressed. It is their need for information to assist in the planning and implementing disease treatment, control and prevention activity that justify the investment in the registry. They include:

- 1. the rnal community
- 2. the RRT provider
- 3. the public health practitioner
- 4. the decision maker in various government and non-government agencies who have responsibilities for any aspects of ESRD treatment, prevention and control
- the researcher with an interest in ESRD and RRT.
- 6. the press and the public.

About MOSS

Malaysian Organ Sharing System or MOSS has been upgraded to web application named e-MOSS. It was officially launched by Y. B. Dato Dr. Hj. Abd Latiff B Ahmad, the Deputy Minister of Health, Malaysia on 1st September 2006.

It is managed by the MOSS committee established under the Malaysian Society of Nephrology (MSN), with National Renal Registry (NRR) assisting in its daily operations.

The functions of the MOSS committee are:

- 1. Make policy decisions concerning MOSS.
- 2. Secure funding from various sources to support its operation.
- 3. Designate a place to be the coordinating centre.
- 4. Canvass the views of nephrologists and other clinical staff involved concerning its policies and operations.
- Appoint panel of nephrologists to assist in the potential recipient management.

The objectives of e-MOSS

- 1. To maintain a list of patients who have voluntarily enrolled as potential recipients in the cadaveric kidney transplantation program in Malaysia.
- 2. To prioritise the waiting list according to an agreed criteria and its scoring system.
- 3. To update the waiting lists according to the specified criteria.
- To provide a list of suitably matched potential recipients when a cadaver organ is available.
- 5. To facilitate centres to effectively manage the patients' listing.

The role of e-MOSS:

All patients registered with NRR will be included in the e-MOSS. However, the subsequent management of the patients' lists depends on the participating centres.

1. The doctor caring for dialysis patients who are potential recipients can now efficiently maintain their patients on the lists and update their patients' treatment information regularly.

2. The transplant coordinating centres can now access the potential recipients' listing that is ranked according to the pre-determined criteria. The patient could be easily contactable in the event of organ donation.

Participation in e-MOSS:

This system is located in a secured site; https://www.macr.org.my/emoss. There are link provided from http://msn.org.my or http://msn.org.my/nrr. All dialysis centres are welcome to be an e-MOSS user.

How to register with e-MOSS?

- The dialysis centre needs to register as an e-MOSS user. Registration instruction and its documents are available in this web application.
- 2. Registered centre can nominate more users. However, the authorization must be from the centre's doctor in-charge.
- 3. All e-MOSS users need to complete a user agreement form and submit it to NRR for processing.

Management of e-MOSS:

All patients registered with NRR shall be listed in the e-MOSS on the following day according to the criteria set in the e-MOSS. These are the listing where patients will be grouped:

- 1. SOS List
- 2. On Wait List
- 3. Auto Off List (Pending data update)
- 4. Temporary Off List
- 5. Pending Evaluation
- 6. Ineligible for transplant
- 7. Death and Transplanted

1. SOS List:

Patients on this list are given specially priority as they as expected to have lifespan of less than a year unless renal transplantations are performed. Only nephrologists can request placement of patients into this list and patient will only be placed into this list after approval is obtained from the MOSS Committee.

2. On Wait List

Patients listed here are patients who have met the criteria. These are the potential cadaver organ recipients.

3. Auto Off List (Pending data update)

If the viral serology results of a patient who is in the 'On Wait List' are not updated after 6 months, the system will automatically placed the patient into this list. The patient in this list will not be eligible for organ transplantation.

The patient will be placed back into the "On wait list" once the serology results have been updated and the patient will not be penalized.

4. Temporary Off List

Doctor in charge should place the patient who is temporarily unfit for a transplant into this list so that he/she will not be contacted in the event of organ donation.

5. Pending Evaluation

The potential eligible patients will be listed in the 'Pending List' upon register\ration with NRR. These patients' e-MOSS criteria have not been assessed by centre's doctor.

6. Ineligible for transplant

System auto list those patients who do not meet e-MOSS criteria.

7. Death and Transplanted

These are patients who had a transplant and the graft is still functioning and those patients who had passed away.

PARTICIPATING HAEMODIALYSIS CENTRES

JOHOR

- 1. Amitabha Haemodialysis Centre Johor Bahru
- 2. Batu Pahat Hospital
- 3. Batu Pahat Rotary
- 4. BP Renal Care (Batu Pahat)
- 5. BP Renal Care (Kluang)
- 6. BP Renal Care (Rengit)
- 7. BP Renal Care (Segamat)
- 8. BP Renalcare (Yong Peng)
- 9. Che Eng Khor Centre
- 10. Haemodialysis Mawar Gemas
- 11. Hospital Pakar Sultanah Fatimah Muar
- 12. JB Lions MAA-Medicare Charity Dialysis Centre (1)
- 13. JB Lions MAA-Medicare Charity Dialysis Centre (2)
- 14. JJ Lions Dialysis Centre
- 15. Johor Specialist Hospital
- 16. Kluang Hospital
- 17. Kota Tinggi Hospital
- 18. Mersing Hospital
- 19. Muar Dialysis
- 20. Muar Lions Renal Centre
- 21. Persatuan Membaiki Akhlak-Che Luan Khor NKF
- 22. Pertubuhan Hemodialisis Muhibbah (Labis)
- 23. Pertubuhan Hemodialisis Muhibbah (Segamat)
- 24. Pontian Hospital
- 25. Pontian Rotary Haemodialysis Centre
- 26. Premier Renal Care
- 27. Prima Dialysis Kluang
- 28. Pusat Dialisis & Kesihatan Masjid Bandar Baru Uda
- 29. Pusat Dialisis Nefro Utama (Kota Tinggi)
- 30. Pusat Dialisis Nefro Utama (Pontian)
- 31. Pusat Dialisis Perbadanan Islam (Pontian)
- 32. Pusat Dialisis Wagaf An-nur (Batu Pahat)
- 33. Pusat Dialisis Waqaf An-nur (Kota Raya)
- 34. Pusat Hemodialisis Darul Takzim
- 35. Pusat Hemodialisis Hidayah
- 36. Pusat Hemodialisis Rotary Kota Tinggi
- 37. Pusat Hemodialisis Rotary Kulai
- 38. Pusat Rawatan Perbadanan Islam (Kota Tinggi)
- 39. Puteri Specialist Hospital
- 40. Segamat Hospital
- 41. Sultan Ismail Hospital
- 42. Sultanah Aminah Hospital (Paed)
- 43. Sultanah Aminah Hospital
- 44. Systemic Dialysis Centre (1)
- 45. Systemic Dialysis Centre (2)
- 46. Tangkak Hospital
- 47. Temenggong Seri Maharaja Tun Ibrahim Hospital
- 48. The Rotary HD Centre (Johor Bahru)
- 49. Yayasan Pembangunan Keluarga Johor-NKF
- 50. Yayasan Rotary Kluang
- 51. Zhi En Dialysis Centre

KEDAH

- 52. 807 Rumah Sakit Angkatan Tentera (Sg. Petani)
- 53. Baling Hospital
- 54. Buddhist Tzu Chi (Jitra)
- 55. Kuala Nerang Hospital
- 56. Kulim Haemodialysis (CS Tan)
- 57. Kulim Hospital
- 58. Langkawi Hospital
- 59. Metro Specialist Hospital
- 60. Pertubuhan Bakti Fo En Bandar Kulim
- 61. Pusat Dialisis K K Tan (Kulim)
- 62. Pusat Dialysis K K Tan (Sg Petani)
- 63. Pusat Haemodialisis Dr. Ismail
- 64. Pusat Hemodialisis Beng Siew
- 65. Pusat Hemodialisis Mergong
- 66. Pusat Hemodialisis Seroja
- 67. Pusat Kesihatan Jitra
- 68. Pusat Rawatan Hemodialisis Yayasan Emkay & Sultanah Bahiyah
- 69. Putra Medical Centre
- 70. Rawatan Dialisis Amal Lion_NKF
- 71. Renal Care (Kedah)
- 72. Renal Medicare
- 73. Sik Hospital
- 74. Strand Specialist Hospital
- 75. Sultan Abdul Halim Hospital
- 76. Sultanah Bahiyah Hospital
- 77. Superkids Trinity-NKF Dialysis Centre
- 78. Yan Hospital

KELANTAN

- 79. Gua Musang Hospital
- 80. KB Rotary-MAA Charity Dialysis
- 81. Kuala Krai Hospital
- 82. Machang Hospital
- 83. Nephrolife Haemodialysis Centre
- 84. Pakar Perdana Hospital
- 85. Pasir Mas Hospital
- 86. Pusat Dialisis Yayasan Buah Pinggang Kebangsaan (Kota Bharu)
- 87. Pusat Perubatan Tentera (Kota Bharu)
- 88. Pusat Rawatan Dialisis Islah (Kota Bharu)
- 89. Raja Perempuan Zainab II Hospital
- 90. Renal-Link (Kelantan)
- 91. Tanah Merah Hospital
- 92. Tengku Anis Hospital
- 93. Tumpat Hospital
- 94. Universiti Sains Malaysia Hospital

MELAKA

- 95. 94 Hospital Angkatan Tentera (Terendak)
- 96 .Alor Gajah Dialysis Centre
- 97. Alor Gajah Hospital

MELAKA (con't)

- 98. Amitabha Centre (Melaka)
- 99. Damai Medical & Heart Clinic
- 100. Mahkota Medical Centre
- 101. Melaka Hospital
- 102. Pantai Air Keroh Hospital
- 103. Pusat HD SJAM Bacang Melaka
- 104. Pusat Hemodialisis Suria (Jasin)
- 105. Sinar Hemodialisis
- 106. Tenang Haemodialysis Centre
- 107. Tenang Haemodialysis Jasin
- 108. Yakin Jaya
- 109. Yayasan Kebajikan The Southern Melaka

NEGERI SEMBILAN

- 110. Jelebu Hospital
- 111. Port Dickson Hospital
- 112. Pusat Hemodialisis Mawar N. Sembilan (Bahau)
- 113. Pusat Hemodialisis Mawar N. Sembilan (Lukut)
- 114. Pusat Hemodialisis Mawar N. Sembilan (Seremban)
- 115. Pusat Waqaf An -nur (Senawang)
- 116. Seremban Specialist Hospital
- 117. Tampin Hospital
- 118. Tuanku Ampuan Najihah Hospital
- 119. Tuanku Jaafar Hospital (Paed)
- 120. Tuanku Jaafar Hospital

PAHANG

- 121. Bentong Hospital
- 122. Jerantut Hospital
- 123. Kuala Lipis Hospital
- 124. Kuantan Clinical Diagnostic Centre
- 125. MAA-Medicare Charity (Mentakab)
- 126. Mentakab Haemodialysis Unit
- 127. Muadzam Shah Hospital
- 128. Pahang Buddhist Association
- 129. Pekan Hospital
- 130. Pusat Hemodialisis Islam Makmur
- 131. Raub Hospital
- 132. SJAM-KPS Haemodialysis Centre 9 (Raub)
- 133. Sultan Haji Ahmad Shah Hospital
- 134. Tengku Ampuan Afzan Hospital (Paed)
- 135. Tengku Ampuan Afzan Hospital
- 136. 96 Hospital Angkatan Tentera (Lumut)

PERAK

- 137. Batu Gajah Hospital
- 138. Berchaam Dialysis Centre
- 139. C.S. Loo Kidney & Medical Specialist
- 140. Changkat Melintang Hospital
- 141. Emnur Teguh
- 142. Gerik Hospital
- 143. Hope Haemodialysis Society Ipoh
- 144. Ipoh Hospital

- 145. Ipoh Hospital (Home)
- 146. Kampar Hospital
- 147. Kuala Kangsar Hospital
- 148. MAA-Medicare Charity (Teluk Intan)
- 149. Parit Buntar Hospital
- 150. Persatuan Amal Chin Malaysia Barat
- 151. Pertubuhan Perkhidmatan Haemodialisis Ar-Ridzuan
- 152. Pertubuhan Perkhidmatan Hemodialisis AIXIN Kerian
- 153. Pulau Pangkor Hospital
- 154. Pusat Dialisis Darul Iltizam Taiping
- 155. Pusat Dialisis Ehsan Perak
- 156. Pusat Dialisis Intan
- 157. Pusat Dialisis Kuala Kangsar
- 158. Pusat Dialisis Mutiara
- 159. Pusat Dialisis Penawar Permai
- 160. Pusat Dialisis Taiping (Kamunting)
- 161. Pusat Dialisis Taiping (Kuala Kangsar)
- 162. Pusat Dialisis Taiping (Parit Buntar)
- 163. Pusat Dialisis Taiping
- 164. Pusat Dialysis Setia
- 165. Pusat Hemodialisis Darul Iltizam (Ipoh)
- 166. Pusat Hemodialisis Kampar, Yayasan Nanyang
- 167. Pusat Hemodialisis Manjung
- 168. Renal Care (Ipoh Specialist)
- 169. Selama Hospital
- 170. Seri Manjung Hospital
- 171. Sg Siput Hospital
- 172. Taiping Hospital
- 173. Tanjung Malim Hospital
- 174. Tapah Hospital
- 175. Teluk Intan Hospital
- 176. Who Peng Cheang Seah
- 177. Yayasan Akhlak-NKF Taiping
- 178. Yayasan Dialysis Pendidikan Akhlak Perak-NKF Ipoh

PERLIS

- 179. Tuanku Fauziah Hospital
- 180. Tuanku Syed Putra Haemodialysis Centre

PULAU PINANG

- 181. AMD Rotary (Penang)
- 182. Asia Renal Care (Penang)
- 183. Balik Pulau Hospital
- 184. Buddhist Tzu Chi Dialysis Centre (Butterworth)
- 185. Buddhist Tzu Chi HD Centre (Penang)
- 186. Bukit Mertajam Hospital
- 187. Bukit Mertajam Specialist Hospital
- 188. Fo Yi NKF Dialysis Centre
- 189. Gleneagles Medical Centre
- 190. Island Hospital
- 191. K K Tan Specialist (BM)
- 192. Kepala Batas Hospital
- 193. Lam Wah Ee Hospital
- 194. Loh Guan Lye Specialist Centre
- 195. MAA-Medicare Charity (Butterworth)

PULAU PINANG (con't)

- 196. Pantai Mutiara Hospital
- 197. Penang Adventist Hospital
- 198. Penang Caring Dialysis Society
- 199. Penang Hospital
- 200. Penang Hospital (Home)
- 201. Penang Hospital (Paed)
- 202. Pertubuhan Dialisis Rotary-Satu Hati
- 203. Pertubuhan Hemodialisis SPS
- 204. Province Wellesley Renal Medifund
- 205. Pusat Haemodialisis Zakat Jawi
- 206. Pusat Haemodialysis St Anne BM
- 207. Pusat Hemodialisis Zakat (Balik Pulau)
- 208. Pusat Hemodialisis Zakat (Bukit Mertajam)
- 209. Pusat Hemodialisis Zakat (Butterworth)
- 210. PWRM (BM) Dialysis Centre
- 211. Renal Link (Penang)
- 212. Seberang Jaya Hospital (Butterworth)
- 213. Seberang Perai (Bagan)
- 214. Sungai Bakap
- 215. The Penang Community HD Society
- 216. TSC Renal Care

SABAH

- 217. Beaufort Hospital
- 218. Beluran Hospital
- 219. Duchess of Kent Hospital
- 220. Keningau Hospital
- 221. Kota Belud Hospital
- 222. Kota Kinabatangan Hospital
- 223. Kota Marudu Hospital
- 224. Kudat Hospital
- 225. Lahad Datu Hospital
- 226. Likas Hospital
- 227. MAA-Medicare Charity (Kota Kinabalu)
- 228. Nobel Dialysis Centre
- 229. Papar Hospital
- 230. Persatuan Buah Pinggang Sabah
- 231. Pusat Hemodialisis Palmcare Malaysia
- 232. Queen Elizabeth Hospital
- 233. Ranau Hospital
- 234. Rotary Tawau Tanjung
- 235. Sabah Medical Centre
- 236. Semporna Hospital
- 237. Sipitang Hospital
- 238. SJAM-KPS Pusat Hemodialisis Centre 10 (Bintulu)
- 239. Tambunan Hospital
- 240. Tawau Hospital
- 241. Tenom Hospital

SARAWAK

- 242. 801 Rumah Sakit Angkatan Tentera (Kuching)
- 243. Bau Hospital
- 244. Betong Hospital

- 245. Bintulu Hospital
- 246. CHKMUS-MAA Medicare Charity
- 247. Kanowit Hospital
- 248. Kapit Hospital
- 249. KAS-Rotary-NKF
- 250. Kuching Specialist Hospital
- 251. Lawas Hospital
- 252. Limbang Hospital
- 253. Lundu Hospital
- 254. Marudi Hospital
- 255. Miri Hospital
- 256. Miri Red Crescent Dialysis Centre
- 257. Mukah Hospital
- 258. Normah Medical Specialist Centre
- 259. Rejang Medical Centre
- 260. Saratok Hospital
- 261. Sarawak General Hospital
- 262. Sarikei Hospital
- 263. Serian Hospital
- 264. Sibu Hospital
- 265. Sibu Kidney Foundation
- 266. Simunjan Hospital
- 267. SJAM-KPS Haemodialysis Centre 8 (Sibu)
- 268. Sri Aman Hospital
- 269. Timberland Medical Centre

SELANGOR

- 270. 819 Rumah Sakit Angkatan Tentera
- 271. Ampang Hospital
- 272. Ampang Puteri Specialist Hospital
- 273. Apex Club of Klang-NKF Charity Dialysis Centre
- 274. Assunta Hospital
- 275. Bakti-NKF Dialysis Centre
- 276. Bangi Dialysis Centre
- 277. Banting Hospital
- 278. Berjaya NKF Dialysis Centre
- 279. Caring Dialysis Centre (Tanjong Karang)
- 280. Damansara Specialist Hospital
- 281. Haemodialysis Association Klang
- 282. Haemodialysis Edina
- 283. Healthcare Dialysis Centre
- 284. Hemodialisis Yayasan Veteran ATM
- 285. Kajang Dialysis Centre
- 286. Kajang Hospital
- 287. Kelana Jaya Medical Centre
- 288. Kuala Kubu Bharu Hospital
- 289. MAA-Medicare Charity (Kajang)
- 290. Persatuan Dialisis Kurnia PJ
- 291. PingRong-NKF
- 292. Pusat Dialisis Aiman (Shah Alam)
- 293. Pusat Dialisis Mesra (Kuala Selangor)
- 294. Pusat Dialisis Sijangkang
- 295. Pusat Dialysis Mesra (Kapar)
- 296. Pusat Dialysis Mesra KKB
- 297. Pusat Dialysis Putra Jaya (Semenyih)

SELANGOR (con't)

- 298. Pusat Hemodialisis Fasa
- 299. Pusat Hemodialisis Kau Ong Yah Ampang
- 300. Pusat Hemodialisis Majlis Perbandaran Kelang
- 301. Pusat Hemodialisis Mawar N. Sembilan (Sepang)
- 302. Pusat Hemodialisis Mawar N. Sembilan (Seri Kembangan)
- 303. Pusat Perubatan Primier HUKM
- 304. Pusat Rawatan Dialisis Islah (Batu Caves)
- 305. Pusat Rawatan Hemodialisis Felina
- 306. Rawatan Dialysis Bukit Tinggi
- 307. Reddy Clinic
- 308. Renal Associates
- 309. S.P. Menon Dialysis Centre (Klang)
- 310. S.P. Menon Dialysis Centre (Petaling Jaya)
- 311. Selangor Medical Centre
- 312. Selayang Hospital
- 313. Serdang Hospital
- 314. SJAM-KPS Haemodialysis Centre 1 (Raja Muda Musa)
- 315. SJAM-KPS Haemodialysis Centre 11 (Shah Alam)
- 316. SJAM-KPS Haemodialysis Centre 2 (Klang)
- 317. SJAM-KPS Haemodialysis Centre 3 (Banting)
- 318. SJAM-KPS Haemodialysis Centre 5 (Rawang)
- 319. SJAM-KPS Haemodialysis Centre 6 (Kuala Selangor)
- 320. Smartcare Dialysis Centre (Subang Jaya)
- 321. Sri Kota Medical Centre
- 322. Subang Jaya Medical Centre
- 323. Sungai Buloh Hospital
- 324. Sunway Medical Centre (1)
- 325. Sunway Medical Centre (2)
- 326. Syukur Elit Sdn Bhd
- 327. Tanjung Karang Hospital
- 328. Tengku Ampuan Jemaah Hospital
- 329. Tengku Ampuan Rahimah Hospital
- 330. Universiti Kebangsaan Malaysia Bangi
- 331. Yayasan Kebajikan SSL (Puchong)
- 332. Yayasan Kebajikan SSL (Petaling Jaya)

TERENGGANU

- 333. Besut Hospital
- 334. Dungun Hospital
- 335. Hulu Terengganu Hospital
- 336. Kemaman Hospital
- 337. Pusat Dialisis Epic
- 338. Pusat Dialisis Terengganu / NKF
- 339. Pusat Hemodialisis Nabilah
- 340. Pusat Rawatan Dialisis Islah (Kuala Terengganu)
- 341. Sultanah Nur Zahirah Hospital

WILAYAH PERSEKUTUAN

- 342. Labuan Hospital
- 343. Aiman Dialysis Centre
- 344. Charis-NKF Dialysis Centre
- 345. Cheras Dialysis Centre
- 346. Kg Baru Medical Centre

- 347. Kuala Lumpur Hospital (Home)
- 348. Kuala Lumpur Hospital (Paed.)
- 349. Kuala Lumpur Hospital (Unit 1)
- 350. Kuala Lumpur Hospital (Unit 3)
- 351. Kuala Lumpur Hospital (Unit 4)
- 352. Kuala Lumpur Lions Renal Centre
- 353. Lifeline Dialysis Clinic
- 354. MAA-Medicare Charity (Cheras)
- 355. MAA-Medicare Charity (Kuala Lumpur)
- 356. National Kidney Foundation Dialysis Centre (Kuala Lumpur)
- 357. Pantai Indah Hospital
- 358. Pantai Medical Centre (1)
- 359. Pantai Medical Centre (2)
- 360. Poliklinik Komuniti Tanglin
- 361. Pusat Dialisis Falah
- 362. Pusat Dialisis Pusat Punggutan Zakat
- 363. Pusat Hemodialisis Dato' Lee Kok Chee
- 364. Pusat Hemodialisis Harmoni
- 365. Pusat Hemodialisis PUSRAWI
- 366. Pusat Hemodialisis Waz Lian
- 367. Pusat Hemodialisis Yayasan Felda
- 368. Pusat Pakar Tawakal
- 369. Pusat Rawatan Dialisis Nefro Utama (Setapak)
- 370. Rawatan Haemodialysis Koswip
- 371. Renal Dialysis Centre
- 372. Renal Healthcare
- 373. Renal-Link Sentosa (Sentosa Hospital)
- 374. Rotary Damansara-NKF Dialysis
- 375. S.P. Menon Dialysis Centre (Kuala Lumpur)
- 376. Smartcare Dialysis Clinic (Cheras)
- 377. The Kidney Dialysis Centre (Jalan Kelang Lama)
- 378. The Kidney Dialysis Centre (Jalan Ipoh)
- 379. The Nayang-NKF Dialysis Centre
- 380. Tung Shin Hospital & Yayasan Nanyang Press
- 381. Tung Shin Hospital
- 382. Universiti Kebangsaan Malaysia Hospital
- 383. University Malaya Medical Centre
- 384. University Malaya Specialist Centre
- 385. Putrajaya Hostpital

PARTICIPATING PERITONEAL DIALYSIS CENTRES

Centre Name		State
1.	BP Renal Care	Johor
2.	Sultanah Aminah Hospital (Paed)	Johor
3.	Sultanah Aminah Hospital	Johor
4.	Sultanah Bahiyah Hospital	Kedah
5.	Raja Perempuan Zainab II Hospital	Kelantan
6.	Universiti Sains Malaysia Hospital	Kelantan
7.	Damai Medical & Heart Clinic	Melaka
8.	Melaka Hospital	Melaka
9.	Tuanku Jaafar Hospital (Paed)	Negeri Sembilan
10.	Tuanku Jaafar Hospital	Negeri Sembilan
11.	Tengku Ampuan Afzan Hospital	Pahang
12.	96 Hospital Angkatan Tentera (Lumut)	Perak
13.	Ipoh Hospital	Perak
14.	Penang Hospital (Paed)	P. Pinang
15.	Penang Hospital	P. Pinang
16.	Queen Elizabeth Hospital	Sabah
17.	Sabah Medical Centre	Sabah
18.	Sarawak General Hospital	Sarawak
19.	Selayang Hospital	Selangor
20.	Serdang Hospital	Selangor
21.	Tengku Ampuan Rahimah Hospital	Selangor
22.	Sultanah Nur Zahirah Hospital	Terengganu
23.	Kuala Lumpur Hospital (Paed.)	W. P. Kuala Lumpur
24.	Kuala Lumpur Hospital	W. P. Kuala Lumpur
25.	Universiti Kebangsaan Malaysia Hospital	W. P. Kuala Lumpur
26.	University Malaya Medical Centre	W. P. Kuala Lumpur

PARTICIPATING TRANSPLANT FOLLOW-UP CENTRES

Centre Name		State
1.	Batu Pahat Hospital	Johor
2.	Hospital Sultan Ismail Pandan	Johor
3.	Hospital Sultanah Aminah (Paed)	Johor
4.	Kluang Hospital	Johor
5.	Pakar Sultanah Fatimah Muar Hospital	Johor
6.	Pontian Hospital	Johor
7.	Segamat Hospital	Johor
8.	Sultanah Aminah Hospital	Johor
9.	Alor Setar Hospital	Kedah
10.	Kota Bharu Hospital	Kelantan
11.	Universiti Sains Malaysia Hospital	Kelantan
12.	Damai Medical & Heart Clinic	Melaka
13.	Mahkota Medical Centre	Melaka
14.	Melaka Hospital	Melaka
15.	Seremban Hospital	Negeri Sembilan
16.	Tg. Ampuan Afzan Hospital	Pahang
17.	Ipoh Hospital	Perak
18.	Taiping Hospital	Perak
19.	Pulau Pinang Hospital	P. Pinang
20.	Duchess of Kent Hospital	Sabah
21.	Queen Elizabeth Hospital	Sabah
22.	Sabah Medical Centre	Sabah
23.	Tawau Hospital	Sabah
24.	Bintulu Hospital	Sarawak
25.	Miri Hospital	Sarawak
26.	Sarawak General Hospital	Sarawak
27.	Sibu Hospital	Sarawak
28.	Timberland Medical Centre	Sarawak
29.	Ampang Puteri Specialist Hospital	Selangor
30.	Selangor Medical Centre	Selangor
31.	Selayang Hospital	Selangor
32.	Serdang Hospital	Selangor
33.	Subang Jaya Medical Centre	Selangor
34.	Sunway Medical Centre	Selangor
35.	Tan Medical Renal Clinic	Selangor
36.	Tg. Ampuan Rahimah Hospital	Selangor
37.	Kemaman Hospital	Terengganu
38.	Kuala Terengganu Hospital	Terengganu
39.	Kuala Lumpur Hospital	W. P. Kuala Lumpur
40.	Kuala Lumpur Hospital (Paed)	W. P. Kuala Lumpur
41.	Renal Dialysis Centre	W. P. Kuala Lumpur
42.	Universiti Kebangsaan Malaysia Hospital	W. P. Kuala Lumpur
43.	University Malaya Medical Centre	W. P. Kuala Lumpur

CONTRIBUTING EDITORS

Chapter	Title	Editors	Institutions
1	All Renal Replacement Therapy in Malaysia	Lim Teck Onn Lim Yam Ngo	Clinical Research Centre, HKL Kuala Lumpur Hospital
2	Dialysis in Malaysia	Lim Teck Onn Lim Yam Ngo Lee Day Guat	Clinical Research Centre, HKL Kuala Lumpur Hospital Kuala Lumpur Hospital
3	Death & Survival on Dialysis	Wong Hin Seng Ong Loke Meng Wan Shaariah Md Yusuf	Selayang Hospital Penang Hospital Tuanku Jaafar Hospital
4	Quality of Life & Rehabilitation Outcomes	Liu Wen Jiun Chew Thian Fook Alinda Chiu Sze Fung Zaki Morad B Mohd Zaher	Sultanah Aminah Hospital Tuanku Jaafar Hospital University Putra Malaysia International Medical University
5	Paediatric Renal Replacement Therapy	Lee Ming Lee Lynster Liaw Susan Pee Wan Jazilah Wan Ismail Lim Yam Ngo	Tuanku Jaafar Hospital Penang Hospital Sultan Ismail Hospital Selayang Hospital Kuala Lumpur Hospital
6	Management of Anaemia in Dialysis Patients	Philip N. Jeremiah Bee Boon Cheak	Ampang Puteri Specialist Hospital Selayang Hospital
7	Nutrition Status on Dialysis	Ahmad Fauzi Abdul Rahman Tilakavati Karupaiah Winnie Chee Siew Swee	Puteri Specialist Hospital Faculty of Allied Health Sciences, UKM Faculty of Allied Health Sciences, UKM
8	Blood Pressure Control and Dyslipidemia	Prasad Menon Lee Wan Tin	Subang Jaya Medical Centre Subang Jaya Medical Centre
9	Management of Renal Bone Disease in Dialysis Patients	Fan Kin Sing Rozina Bt Ghazalli Ching Chen Hua Liew Yew Fong	Gleneagle Intan Medical Centre Penang Hospital Sultanah Bahiyah Hospital Penang Hospital
10	Hepatitis on Dialysis	Teo Sue Mei Claire Tan Hui Hong Foo Sui Mei	Ipoh Hospital Sarawak Hospital Ipoh Hospital
11	Haemodialysis Practices	Tan Chwee Choon Shahnaz Shah Firdaus Khan Rafidah Abdullah Norleen Bt Zulkarnain Sim	Tengku Ampuan Rahimah Hospital Tengku Ampuan Rahimah Hospital Selayang Hospital Tengku Ampuan Rahimah Hospital
12	Chronic Peritoneal Dialysis Practices	Sunita Bavanandan Lily Mushahar	Kuala Lumpur Hospital Kuala Lumpur Hospital
13	Renal Transplant	Goh Bak Leong Zaki Morad B Mohd Zaher Fan Kin Sing Lily Mushahar Rohan Malek Prasad Menon Tan Si Yen	Serdang Hospital International Medical University Gleneagle Intan Medical Centre Kuala Lumpur Hospital Selayang Hospital Subang Jaya Medical Centre University Malaya Medical Centre

FOREWORD

The Renal Replacement Therapy program in the country continues to grow and we achieved a treatment rate of 118 per million population in 2006. There were nearly 15,000 patients on dialysis at the end 2006. The growth has, as in previous years, been contributed by the combined efforts of the three provider groups – public, non-governmental and private sectors. This success is something we are justifiably proud of. It has attracted the attention of countries which faced similar challenges who want us to share our experience. Nonetheless a number of challenges identified in the previous reports remain. There is still the problem of equity in dialysis provision. Treatment rates have improved in states which were underprovided over the last few years. However the gap between these states and the economically developed ones remains. There is a lack of centres run by NGOs to complement the efforts of the Ministry of Health in the states with low provision rates. It would appear that charitable organizations that fund NGO dialysis centres flourish in areas with higher income and quite rightly so as they depend on population in the area for support.

Last year saw the implementation of the Regulations on Hemodialysis centres of the Private Healthcare Facilities Act. These regulations are meant to ensure that dialysis centres meet the minimum standards which are enforceable by law. It is a welcomed move and officers in the relevant department of the Ministry visited dialysis centres to ensure regulations in the act were enforced. While many may argue that certain provisions of the regulations need revision, the mere fact that a minimum set of standards can be enforced is a big and laudable step in the care of dialysis patients. But nephrologists and all others interested in the care of ESRD patients should set standards of care beyond that of the Act. Data from this report showed that we have done well in most aspects of dialysis practice. Death rate for Hemodialysis patients has remained unchanged over the last 10 years despite more older and diabetic patients being accepted for treatment. There is still considerable variation amongst centres in various parameters that are monitored by the registry. We should set targets for various aspects of dialysis practice that are consistent with those advocated by the international guidelines and work on minimising the variations.

The Registry in its 15th year of existence has collected enormous amount of data that has been useful to various people not the least the practising nephrologists. It has introduced innovations including the "centre report card" where individual centres can gauge its performance against the national average. The registry should now consider collecting data on the cost of dialysis care. The country spends hundreds of millions of ringgit on dialysis and it will help the government as well as the practitioners to have accurate data on the financing of the dialysis program.

The 14th report also shows an alarming trend in the incidence of diabetic nephropathy as a cause of ESRD. Fifty seven percent of new patients accepted for dialysis last year were diabetics. We undoubtedly have the dubious honour of being the number one country in the world when it comes to diabetes as a cause of ESRD. We should consider doing a more detailed study to find whether diabetes is the real cause of ESRD in these patients or present as a co-morbid illness in a patient with some other cause of ESRD. The current method of data collection does not distinguish diabetes as cause or a co-morbid illness.

Renal transplantation continues to stagnate in the country as evidenced by this and the previous years reports. There has been an attempt at revitalizing transplantation including the formulation of a National Organ and Tissue Transplantation policy, the proposed setting up of a Transplantation unit in the Ministry of health to catalyse the Transplantation process and the creation of a budget specific for Transplantation. Nephrologists must play a more active role to promote transplantation. The clamp down by the Chinese authorities on commercial cadaveric transplantation in hospitals in China should be taken as an opportunity to promote live related kidney transplantation.

Finally we thank all those who contributed to the success of this report - the indefatigable Ms Lee Day Guat and her staff, Chapter editors and of course the Report editors Drs Lim Yam Ngo and Lim Teck Onn whose untiring efforts at ensuring an accurate and readable document shows through in this 14th report.

Dr. Rozina Ghazalli Chairperson, National Renal Registry

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REPORT SUMMARY

- Intake of new dialysis patients showed a linear increase over the years -from 1136 in 1997 to 3152 in 2006 with corresponding treatment rates of 52 and 118 per million population.
- Prevalent dialysis patients increased from 3698 (171 per million) in 1997 to almost 15000 (550 per million) at year end 2006.
- Transplant rates remain at about 5-7 per million. Patients with functioning renal transplants increased from 1083 (50 per million) to 1725 (65 per million) over the same period.
- Dialysis treatment rates varied from about 56-104 per million state population in the economically underdeveloped states to >140 per million in the more economically advantaged states in 2006.
- From the centre survey carried out at the end of 2006, there were a total of 14946 dialysis patients, 33.7% in MOH centres, 31.1% in non-governmental organizations (NGO) centres and 32.8% in private sector. Almost all patients in NGO and private centres were on centre HD. In MOH, 24% were on chronic PD. HD capacity to patient ratio showed slight reduction in 2006 with the lowest in Kelantan and Terengganu.
- The treatment gap between men and women has remained consistent over the years.
- Dialysis treatment rates for those < 55 years of age had leveled while those >55 years continued to increase. 53% of new dialysis patients were at least 55 years old
- At least 87% of new patients were accepted into centre haemodialysis
- The government continued to fund about 53% of dialysis treatment, NGO funding was 14% in 2006, and self funding had decreased to 27%.
- Proportion of new ESRD patients due to diabetes mellitus increased further to 57% in 2006, followed by hypertension at 6%.
- The annual death rate for those on hemodialysis remained relatively unchanged while the annual death rate on CAPD showed a 15% reduction over the last 10 years.
- Cardiovascular disease and death at home remained the commonest cause of death in 2006 at 28 and 21% respectively; sepsis was third at 13%.
- The overall unadjusted 5 years and 10 years patient survival on dialysis were 57% and 35% respectively. HD patient survival was superior to those on CAPD even after adjusting for age and diabetes. There was wide centre variation with regards to HD patient survival at one year which was more apparent at 5 years. Adjusted patient survival varied widely between CAPD centres at 5-years but not at 1-year.
- Older and diabetic patients had poorer survival on dialysis.
- After adjustment for multiple risk factors, younger patients, lower diastolic BP, lower calcium and phosphate were associated with lower mortality. Low serum albumin, Kt/V, haemoglobin and calcium phosphate product were associated with higher mortality.
- Median QoL index scores on dialysis were satisfactory and HD patients achieved a lower score than CAPD patients. Diabetes Mellitus and older age group were associated with lower median QoL index scores. Higher employment rate amongst HD and CAPD patients who started dialysis earlier
- In 2006, 84% of HD and 74% of CAPD patients were on erythropoietin (EPO). Blood transfusion rate in dialysis patients increase to 18% in 2006. Use of IV Iron has increased. The median weekly EPO dose has increased to 8000 units, in both HD and CAPD patients. Median haemoglobin level increased to 10.5g/L in 2006. Variations were seen in the use of EPO, blood transfusion rates, measures of iron stores and hemoglobin levels in HD and CAPD centres

REPORT SUMMARY (con't)

- Serum albumin levels remained at mean and median of about 40g/L for HD and about 33.5g/L in CAPD patients in 2006. There were wide variations in the proportion of patients with serum albumin >40g/L in both HD and CAPD centres.
- BMI for HD patients has stabilized around 24, but was still increasing for patients on CAPD patients improved. There was some variation in proportion of patients with BMI ≥ 18.5 in both HD and PD centres.
- In 2006, there was a trend noted towards increasing predialysis systolic but not diastolic BP in HD patients. BP control in CAPD patients improved slightly over the years. The variation noted among the various HD and PD centres in median systolic or diastolic BP was not wide but there was wide variation in the proportion of patients achieving BP of <140/90 mmHg. BP control in CAPD was much better than in haemodialysis patients
- Improving cholesterol levels were seen in both HD and CAPD patients with lower levels seen in HD patients. Serum triglyceride levels did not show much change over the years and was lower in HD patients. There was some variation in lipid control between dialysis centres.
- In 2006 calcium carbonate remained the major phosphate binder in both HD and CAPD patients. Phosphate control was better in CAPD patients. The target of calcium phosphate product of less than 4.5 mmol2/L2 was achieved more by CAPD patients than HD. Mean iPTH levels was about 220 ng/ml for both HD and CAPD patients in 2006. There was wide variation in serum calcium, phosphate, calcium phosphate product and iPTH among both hemodialysis and CAPD centres.
- The prevalence of Hepatitis B infection has remained unchanged over the years, and was quite similar between HD and CAPD patients. HCV prevalence showed a declining trend to a median of 8% in 2006. The proportion of HCV infected patients varied widely between HD centers. Previous renal transplant and history of blood transfusion were associated with a significantly higher risk of HCV seroconversion Completely assisted HD patients had a significantly lower risk of acquiring HCV infection
- Haemodialysis practices: There was increased use of brachiocephalic fistulae, higher blood flow rates, increased usage of synthetic membranes, increased number of reuse and almost universal use of bicarbonate buffer. Although the prescribed median KT/V was 1.6 in 2006, the delivered median KT/V was only1.4. The percentage of patients with a delivered KT/V > 1.2 and KT/V > 1.3 was 82% and 66% respectively. In 2006, the median urea reduction ratio was 71.9% and the percentage of patients with URR > 65% was 79%. There was wide variation in the proportion of patients with blood flow rates of >250 ml/min, prescribed KT/V of >1.3 and delivered KT/V of >1.2 but less variation in urea reduction ratio among HD centres. Technique survival was better in HD compared to PD, in the younger age groups and the non-diabetics but was not related to the year of starting dialysis.
- Chronic PD practices In 2006, CAPD remained the commonest mode of PD at 90% but APD use increased to 6%. 92% were on the Baxter disconnect system. 91% were on 4 exchanges a day, 82% used a fill volume of 2 L. The median delivered weekly Kt/V was 2.1, 59% achieved target Kt/V of >2.0 with a 3-fold variation between the highest- and the lowest-performing centres. Technique survival was better for younger patients and non-diabetics but was not related to the year of starting dialysis or gender. Commonest cause of technique failure was peritonitis and membrane failure.
- In 2006, median peritonitis rate was 33 patient-months but varied between 16 and 103 patient-months/episode among centres. Gram positive and Gram negative organisms each accounted for 32% and 22% of peritonitis episodes.

REPORT SUMMARY (con't)

Renal Transplantation

- There were 161 new renal transplant recipients in 2005 and 1725 with functioning transplants at the end of 2006.
- Mean age of new transplant patients in 2006 was 37 years; 68% were male, 21% diabetic, 7% HbsAg positive and 7% anti-HCV positive at the time of transplantation.
- Commonest known primary renal disease was chronic glomerulonephritis followed by hypertension and diabetes mellitus.
- In 2006, commercial transplants from China constituted 57% of all new renal transplantation, live donor transplantation 21% and local cadaveric transplants contributed only 19%.
- 76% of renal transplant recipients were on cyclosporine, 97% on prednisolone, and 17% were on tacrolimus. 48% were on MMF and 33% on azathioprine
- 34% of the prevalent renal transplant recipients had diabetes mellitus before transplantation, another 8% developed diabetes mellitus post transplantation
- In 2006, 49 (3%) of transplant recipients died and 30 (2%) lost their grafts. Infection, cardiovascular disease and death at home were the commonest causes of death for the last decade and accounted for 41%, 19% and 13% respectively. Renal allograft rejection accounted for 50-78% of graft losses for the last 10 years
- The overall transplant patient survival rate from 1993 to 2006 was 95%, 91%, 88% and 80% at 1 year, 3 years, 5 years and 10 years respectively, while the overall graft survival rate was 92%, 85%, 79% and 63% respectively. Living donor and commercial cadaver grafts had the best patient and graft survival rates.
- In 2006, 85.5% of recipients were hypertensive, 22.9% had diabetes and 57.1% had renal insufficiency fulfilling the criteria for CKD III and above, 13.1% were obese with BMI above 30.

Paediatric Renal Replacement Therapy

- Intake of new paediatric dialysis patients increased from 41 in 1997 to 88 in 2006 giving a dialysis acceptance rate of 4 per million age related population (pmarp) to 8 pmarp respectively.
- New renal transplant rate was 2 pmarp in 2005 and 2006.
- At the end of 2006 there were a total of 468 patients under 20 on dialysis giving a dialysis prevalence rate 42 pmarp.
- The number of patients with functioning transplants in 2006 was 139 giving a prevalence rate of 12 pmarp.
- Dialysis treatment rates were higher in the economically advantaged states of Malaysia.
- The number of 0-4 year olds provided RRT remained very low.
- Chronic PD was the initial dialysis modality in about 50% of patients. Of this 10-20% were on automated PD..
- About 90% received dialysis in government centres.
- Glomerulonephritis accounted for 24% of ESRD, focal segmental glomerulosclerosis 10%, and SLE 8%. 39% of patients had unknown primary renal disease.
- Patient survival for HD was 95% at 1 year, 84% at 5 years. CAPD patient survival was 95% at 1 year and 79% at 5 years.
- In the last 10 years, live related transplantation constituted slightly more than 50%, and cadaveric transplantation a quarter with another quarter from overseas cadaveric transplantation.
- Transplant patient survival was 98% at 1 year and 94% at 5 years; graft survival was 89% at 1 year and 81% at 5 years.

ABBREVIATIONS

APD Automated peritoneal dialysis

CAPD Continuous ambulatory peritoneal dialysis

CCPD Continuous cycling peritoneal dialysis

CKD Chronic Kidney Disease
CRC Clinical Research Centre
ESRD End Stage Renal Disease

HD Haemodialysis

JNC IV Joint National Council IV

MOH Ministry of Health

MOSS Malaysian Organ Sharing System NGO Non-governmental organization

NRR National Renal Registry

PD Peritoneal dialysis

pmarp Per million age related population

pmp Per million population

PTDM Post transplant diabetes mellitus

RRT Renal replacement therapy

SDP Source data producer

TX Transplant

CHAPTER 1

All Renal Replacement Therapy In Malaysia

Lim Yam Ngo Lim Teck Onn

1.1 Stock and Flow

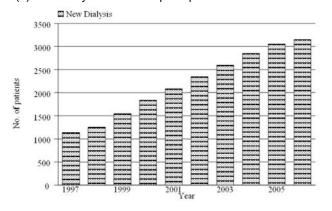
The intake of new dialysis patients continued to increase linearly over the years - from 1136 in 1997 to at least 3152 in 2006. The number of prevalent dialysis patients also increased linearly from 3698 in 1996 to almost 15000 at year end 2006.

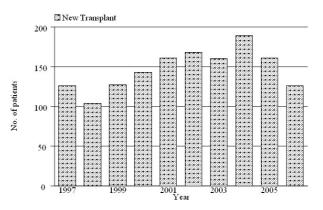
The number of new transplant had been increasing since the late 1990s, but with increasing proscription against commercial transplant, is beginning to how a downward trend from the year 2004. Patients with functioning renal transplants showed a moderate increase - from 1083 to 1725 over the same period. (table and figure 1.01)

Table 1.01: Stock and	Flow of RRT, Malay	ysia 1997 – 2006
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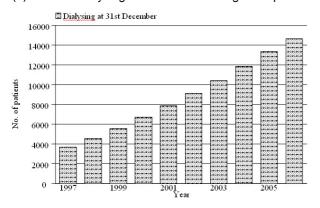
Year	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006
New Dialysis patients	1136	1252	1544	1840	2085	2348	2597	2858	3054	3152
New Transplants	126	104	127	143	161	168	160	189	161	126
Dialysis deaths	316	376	492	594	816	925	1153	1265	1403	1575
Transplant deaths	31	26	25	30	37	32	37	41	41	49
Dialysing at 31st December	3698	4539	5538	6691	7839	9114	10429	11855	13337	14647
Functioning transplant at 31st December	1083	1111	1176	1248	1330	1425	1502	1590	1683	1725

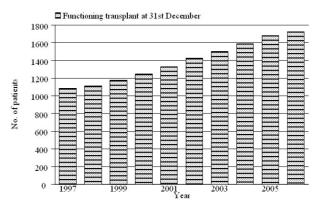
Figure 1.01: Stock and Flow of RRT, Malaysia 1997 – 2006 (a) New Dialysis and Transplant patients





(b) Patients Dialysing and with Functioning Transplant at 31st December 1997 – 2006





1.2 Treatment Provision Rate

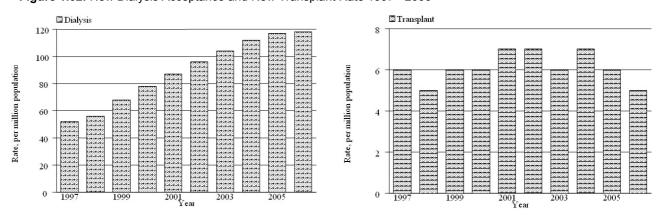
Dialysis acceptance rates doubled from 52 in 1997 to 104 in 2003. Since then, the dialysis acceptance rate has exceeded 110 per million population and was 117 in 2005 and at least 118 in 2006. (Data for 2006 are preliminary since at the time preparation of this report there were still many new cases yet to be notified to registry.)

New transplant rates remained low over the years fluctuating between 5-7 per million population per year. (table and figure 1.02)

Table 1.02: New Dialysis Acceptance Rate and New Transplant Rate per million population 1997 – 2006

Acceptance rate	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006
New Dialysis	52	56	68	78	87	96	104	112	117	118
New Transplant	6	5	6	6	7	7	6	7	6	5

Figure 1.02: New Dialysis Acceptance and New Transplant Rate 1997 - 2006

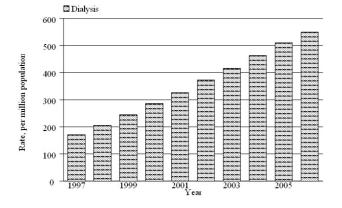


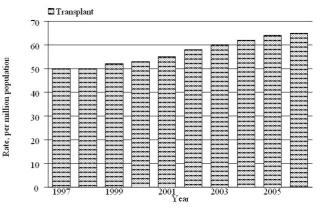
Dialysis prevalence rate increased linearly over the last 10 years, from 171 per million population in 1997 to 510 in 2005 and at least 550 in 2006. The transplant prevalence rate however only showed a slight increase from 50 to 65 per million in 2006. (table and figure 1.03)

Table 1.03: RRT Prevalence Rate per million population 1997 – 2006

Prevalence rate	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006
Dialysis	171	205	244	285	326	372	416	463	510	550
Transplant	50	50	52	53	55	58	60	62	64	65

Figure 1.03: Dialysis and Transplant Prevalence Rate per million population 1997 - 2006





CHAPTER 2

Dialysis In Malaysia

Lim Yam Ngo Lim Teck Onn Lee Day Guat

SECTION 2.1: PROVISION OF DIALYSIS IN MALAYSIA (registry report)

Information on provision of dialysis was obtained from data on individual patients reported to the registry shown in section 2.1 as well as from the centre survey carried out at the end of each year shown in section 2.2.

2.1.1 Dialysis treatment provision

In 2005, 3054 patients commenced dialysis, giving a treatment rate of 117 per million population. There were at least 3152 patients accepted for dialysis in 2006 with corresponding treatment rate of 118 per million population. At year end 2006, a total of 14647 patients were reported to the registry as being on dialysis treatment giving a prevalence rate of 550 per million per year. Dialysis patients lost to follow-up which was negligible in the earlier years almost reached 1% in 2006.

Table 2.1.1: Stock and flow – Dialysis Patients 1997 – 2006

Year	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006
New Dialysis patients	1136	1252	1544	1840	2085	2348	2597	2858	3054	3152
Died	316	376	492	594	816	925	1153	1265	1403	1575
Transplanted	59	61	69	106	132	143	122	149	120	115
Lost to Follow-up	3	8	7	7	8	17	26	36	54	136
Dialysing at 31st December	3698	4539	5538	6691	7839	9114	10429	11855	13337	14647

Table 2.1.2: Dialysis Treatment Rate per million population 1997 – 2006

Year	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006
Acceptance rate	52	56	68	78	87	96	104	112	117	118
Prevalence rate	171	205	244	285	326	372	416	463	510	550

2.1.2. Geographic distribution

The economically advantaged states of Malaysia –Pulau Pinang, Melaka, Johor, Perak, Selangor and W. Persekutuan of Kuala Lumpur, and Negeri Sembilan – continued to have high dialysis treatment rates although the rates appear to be plateauing in the last few years. In 2006, except for Sabah, Sarawak and Kelantan, the other less economically developed states have treatment provision rates touching 100 per million. Pulau Pinang continued to have the highest treatment rate at 181 and Sabah the lowest at 56 per million in 2006.

Table 2.1.3: Dialysis Treatment Rate by State, per million state population 1997 – 2006

State	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006
P Pinang	85	113	124	110	126	160	145	212	197	181
Melaka	95	107	88	150	156	171	185	210	168	171
Johor	79	71	104	131	138	148	146	156	165	180
Perak n	61	64	75	105	103	114	129	144	164	162
Selangor & Kuala Lumpur	76	91	102	121	119	126	137	142	147	144
Negeri Sembilan	73	95	97	116	110	131	147	158	146	139
Kedah & Perlis	55	47	59	69	68	89	106	96	106	96
Terengganu	36	34	36	37	75	89	67	80	99	97
Pahang	46	36	47	49	52	53	66	72	90	104
Kelantan	12	15	27	31	60	62	74	64	79	72
Sarawak	46	33	44	50	67	59	62	73	72	85
Sabah	16	24	32	26	35	37	43	48	46	56

SECTION 2.2: DIALYSIS PROVISION IN MALAYSIA (Centre survey report)

Data submission of individual dialysis and transplant patients to the National Renal Registry which was entirely voluntary prior to 2006 is now made compulsory by the Private Health Care Facilities and Services Act 1996 and its Regulations 2006 which was implemented on 1st May 2006. This however only applies to private and NGO centres and data submission from centres managed by the Ministry of Health, Defence or the Universities is still voluntary.

Dialysis centre surveys have been conducted in December of each year since 1999. This annual cross-sectional survey was carried out to describe the most current level and distribution of dialysis provision for both hemodialysis and peritoneal dialysis at the end of each year. This section reports the results of the centre survey carried out in December 2006. Dialysis provision is expressed in terms of number of centres, HD machines, treatment capacity (one HD machine to 5 patients) and patients.

At the end of 2006, 412 hemodialysis centres and 31 CAPD centres provided dialysis care to 14946 patients. (Data on 14647 inidividual dialysis patients were reported to the Registry giving a dialysis patient ascertainment rate of almost 98%) The Ministry of Health (MOH) provided dialysis to 33.7% of patients, non-governmental organizations (NGO) 31.1% and the private sector at 32.8%. Almost all private dialysis patients received centre haemodialysis treatment compared to the MOH sector where patients on chronic peritoneal dialysis (PD) made up 24% of all dialysis patients. There were no PD patients in NGO centres. (table 2.2.1)

Of the 3 main sectors, the private sector had the largest number of dialysis centres but the NGO centres had the largest HD capacity. (fig 2.2.1 a & b) The Ministry of Health had the lowest HD treatment capacity to patient ratio at 1.47 in 2006 compared to 1.52 in 2005. The HD treatment to patient ratio had increased further in the NGO sector from 1.70 in 2005 to 1.72 in 2006. (fig 2.2.1d)

Table 2.2.1: Number of dialysis centres, HD machines and treatment capacity by sector, December 2006

Sector	HD Centre (No.)	Centre HD machines (No.)	Centre HD capacity (No.)	Centre HD patients (No.)	Centre HD capacity: patient ratio	CAPD Centre (No.)	CAPD patients (No.)	All dialysis patients (No.)
МОН	131	1133	5665	3841	1.47	20	1200	5041
NGO	111	1603	8015	4648	1.72	0	0	4648
Private (PRV))	158	1533	7665	4882	1.57	7	19	4901
University (UNI)	5	41	205	128	1.6	3	129	257
Armed Forces (AF)	7	44	220	96	2.29	1	3	99
TOTAL	412	4354	21770	13595		31	1351	14946

Figure 2.2.1(a): Distribution of dialysis centres by Sector, December 2006

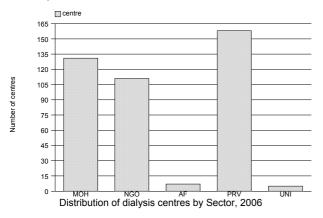


Figure 2.2.1(c): Distribution of dialysis patients by Sector, December 2006

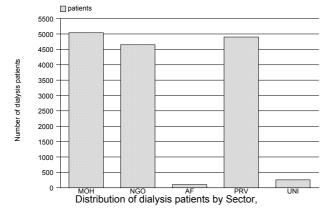


Figure 2.2.1(b): Distribution of HD capacity by Sector, December 2006

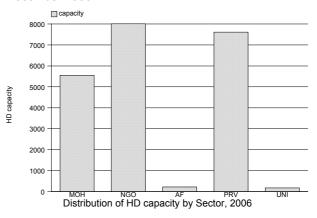
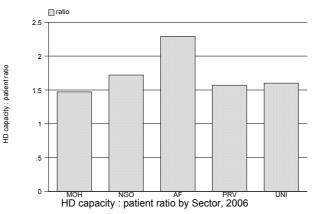


Figure 2.2.1(d): HD capacity: patient ratio by Sector, December 2006



2.2.2. Geographic distribution

compared to the economically disadvantaged states where the rate varied between 61 to 114 per million state population. At the end of 2006 dialysis The economically advantaged states except for Negeri Sembilan have centre HD machine to state population ratio above 200 per million state population treatment rates ranged from a high of 917 for Pulau Pinang to a low of 202 per million state population for Sabah. The HD capacity to patient ratio did not vary too widely between the states except for the economically disadvantaged states of Kelantan and Terengganu where the ratio was 1.21 and 1.36 respectively. (table and figures 2.2.2.). This is unlike previous years when HD capacity to patient ratio was higher in the economically disadvantaged states compared to the advanced states. The intake of patients has obviously outstripped the increase in number of new machines. These states need more HD machines and other resources to enable them to provide dialysis at rates similar to the more economically advantaged states

.⊑ Table 2.2.2: Number of dialysis centres, number of HD machines and treatment capacity, HD capacity to patients ratio and number of dialysis patients by state December 2006

	Centre HD (No.)	Centre HD machines	Centre HD machines pmp	Centre HD capacity (No.)	Centre HD capacity pmp	Centre HD patients (No.)	Centre HD patients pmp	HD capacity: patient ratio	All dialysis patients (No.)	Dialysis treatment rate pmp
Pulau Pinang (Pe)	36	424	284	2120	1421	1247	836	1.7	1369	917
Johor (Jo)	28	299	210	3335	1052	2332	736	1.43	2585	815
Melaka (Me)	15	189	261	945	1303	532	733	1.78	222	292
Selangor & W.Persekutuan (SF)	118	1349	210	6745	1049	3990	621	1.69	4500	200
Perak (Pr)	47	476	208	2380	1042	1487	651	1.6	1541	675
Negeri Sembilan (Ne)	14	154	160	770	801	493	513	1.56	256	218
Kedah & Perlis (KP)	30	302	143	1510	716	904	428	1.67	929	440
Trengganu (Tr)	6	85	82	425	408	350	336	1.21	425	408
Sarawak (Sw)	28	268	411	1340	268	884	375	1.52	942	400
Pahang (Pa)	16	147	101	735	202	427	293	1.72	487	335
Kelantan (Ke)	15	105	69	525	343	386	252	1.36	433	283
Sabah (Sa)	26	188	61	940	305	563	183	1.67	622	202
Malaysia	412	4354	167	21770	833	13595	520	1.6	14946	572

Figure 2.2.2(a): Distribution of dialysis centres by State, December 2006

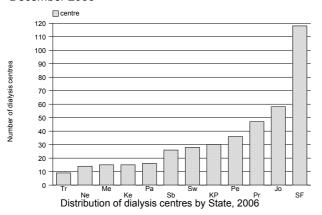


Figure 2.2.2(c): Distribution of dialysis treatment by State, December 2006

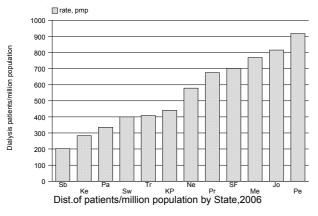


Figure 2.2.2(b): Distribution of dialysis patients by State, December 2006

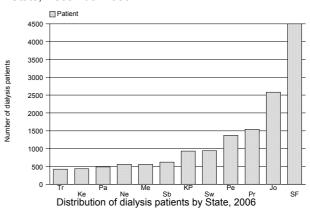
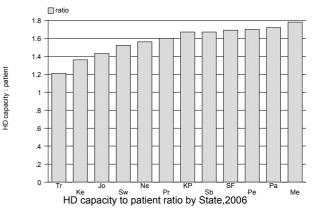


Figure 2.2.2(d): HD capacity to patient ratio by State, December 2006



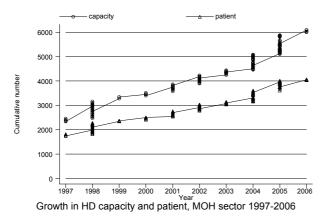
2.2.3 Growth in dialysis provision by sector

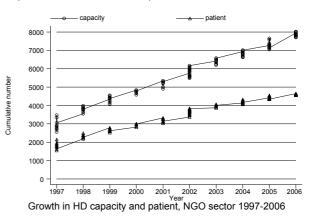
The number of HD patients more than doubled in the last 10 years in all 3 major sectors providing dialysis.(table 2.2.3). The increase in HD capacity almost paralleled that of increase in number of HD patients for MOH and the private sector but showed a divergence in the NGO sector indicating that gap between HD capacity and patient intake is widening. (figures 2.2.3a-c)

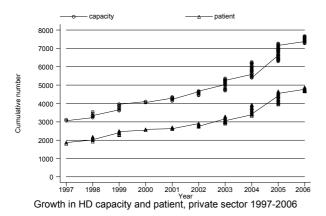
Table 2.2.3: Growth in HD capacity and HD patients in Private, NGO and MOH sectors, 1997 - 2006

Castan	Priv	/ate	NO	30	MC	ЭH
Sector	Cumulative HD capacity	Cumulative HD patients	Cumulative HD capacity	Cumulative HD patients	Cumulative HD capacity	Cumulative HD patients
1997	3115	1872	3475	2148	2460	1817
1998	3545	2188	3985	2490	3160	2281
1999	3960	2502	4545	2792	3340	2365
2000	4105	2585	4835	3012	3510	2505
2001	4345	2697	5345	3328	3865	2758
2002	4670	2902	6165	3837	4225	3036
2003	5360	3292	6570	4063	4440	3148
2004	6260	3928	7005	4324	5085	3607
2005	7260	4655	7650	4544	5880	4009
2006	7665	4882	8015	4648	6090	4065

Figure 2.2.3: Growth in HD capacity and HD patients in Private, NGO and MOH sectors, 1997 - 2006







SECTION 2.3: DISTRIBUTION OF DIALYSIS TREATMENT

2.3.1 Gender distribution

The treatment gap between men and women accepted for dialysis has remained consistent over the years, suggesting this is a true reflection of the difference in ESRD incidence between the 2 sexes rather than any conscious or unconscious bias in treatment allocation. However, figure 2.3.1(ii) shows a convergence in the proportion of prevalent male and female patients. This is probably because of the survival advantage in female patients.

Table 2.3.1(a): Dialysis Treatment Rate by Gender, per million male or female population 1997 – 2006

Gender	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006
Male	51	63	63	81	92	97	110	123	128	137
Female	45	50	57	61	73	89	95	96	110	110

Figure 2.3.1: Dialysis Treatment by Gender 1997 - 2006

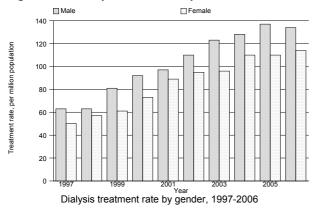
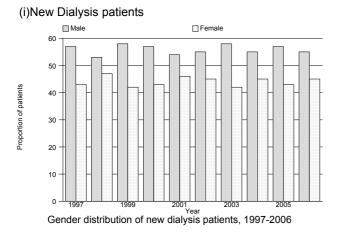


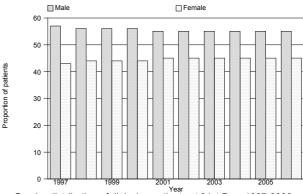
Table 2.3.1(b): Gender distribution of Dialysis Patients 1997 – 2006

Year	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006
New Dialysis patients	1136	1252	1544	1840	2085	2348	2597	2858	3054	3152
% Male	57	53	58	57	54	55	58	55	57	55
% Female	43	47	42	43	46	45	42	45	43	45
Dialysing at 31st December	3698	4539	5538	6691	7839	9114	10429	11855	13337	14647
% Male	57	56	56	56	55	55	55	55	55	55
% Female	43	44	44	44	45	45	45	45	45	45

Figure 2.3.1(b): Gender Distribution of Dialysis patients 1997 – 2006







Gender distribution of dialysing patients at 31st Dec, 1997-2006

2.3.2 Age distribution

Except for those aged 55 years or older which continued to register increase in treatment rates, dialysis treatment rates in the other age groups have plateaued in the last few years, suggesting that almost all patients with ESRD in those age groups who were in need of dialysis were able to access treatment. The treatment rate for patients 65 years and older had exceeded 700 per million in 2006. 53% of new dialysis patients were at least 55 years old.

Table 2.3.2(a): Dialysis Treatment Rate by Age Group, per million age group population 1997 – 2006

Age groups (years)	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006
1-14	3	3	3	4	4	5	4	5	5	5
15-24	15	15	16	18	22	28	26	27	30	26
25-34	39	41	42	46	47	55	51	51	56	53
35-44	80	81	85	98	103	100	102	114	110	104
45-54	166	173	225	249	252	275	278	308	293	313
55-64	291	313	30	432	508	534	586	588	641	595
>=65	213	228	301	347	438	501	584	651	652	710

Figure 2.3.2(a): Dialysis Treatment Rate by Age Group 1997 - 2006

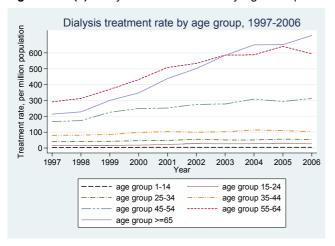
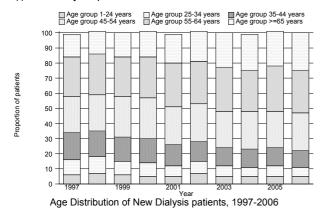


Table 2.3.2(b): Percentage Age Distribution of Dialysis Patients 1997 – 2006

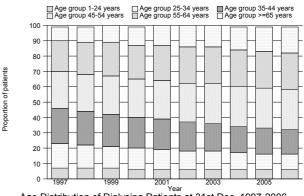
Year	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006
New Dialysis patients	1136	1252	1544	1840	2085	2348	2597	2858	3054	3152
% 1-14 years	1	2	2	1	1	2	1	1	1	1
% 15-24 years	5	5	4	4	4	5	4	4	4	4
% 25-34 years	10	11	9	9	7	8	7	6	7	6
% 35-44 years	18	17	16	16	14	13	12	12	12	11
% 45-54 years	24	24	27	27	25	25	24	25	24	25
% 55-64 years	26	27	26	27	29	28	29	27	30	28
% >=65 years	15	15	16	17	19	20	23	24	23	25
Dialysing at 31st December	3698	4539	5538	6691	7839	9114	10429	11855	13337	14647
% 1-14 years	2	2	2	1	1	1	1	1	1	1
% 15-24 years	5	5	5	5	5	5	5	5	5	5
% 25-34 years	16	15	14	14	13	12	12	11	10	10
% 35-44 years	23	22	21	20	20	19	18	17	17	16
% 45-54 years	24	24	25	25	25	25	26	26	26	26
% 55-64 years	20	21	22	22	23	24	24	24	25	24
% >=65 years	9	10	11	12	13	14	14	15	16	17

Figure 2.3.2(b): Age Distribution of New Dialysis patients 1997 - 2006

(i)New Dialysis patients



(ii) Dialysing patients at 31st December



Age Distribution of Dialysing Patients at 31st Dec, 1997-2006

2.3.3 Method and Location of dialysis

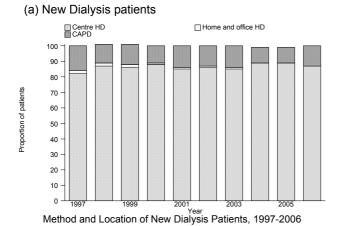
87% of new patients were accepted into centre haemodialysis in 2006. Home/office HD which accounted for a significant proportion of patients in the earlier years of dialysis therapy has died a natural death. Perhaps the future will see home HD being re-introduced as diurnal or nocturnal HD as practiced by the more wealthy nations. Chronic PD accounted for about 12% of new dialysis patients but only 10% of prevalent dialysis patients in 2006. (table & fig 2.3.3)

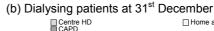
Table 2.3.3: Method and Location of Dialysis 1997 – 2006

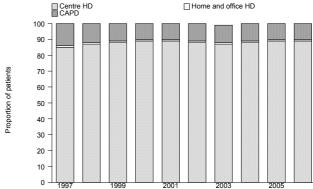
			-							
Year	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006
New Dialysis patients	1136	1252	1544	1840	2085	2348	2597	2858	3054	3152
% Centre HD	82	87	86	88	85	86	85	89	89	87
% Home and office HD	2	2	2	1	1	1	1	0	0	0
% CAPD	16	12	13	11	14	13	14	10	10	12
Dialysing at 31st December	3181	3908	4785	5817	6869	8072	9243	10477	11739	12566
% Centre HD	85	87	88	89	89	88	87	88	89	89
% Home and office HD	1	1	1	1	1	1	1	1	1	1
% CAPD	14	12	11	10	10	11	11	11	10	10

 $^{^{\}star}$ Analysis of $\,$ Dialysing at 31st December is analyzed base on patient annual return

Figure 2.3.3: Method and Location of Dialysis Patients 1997–2006







Method and Location of Dialysing Patients at 31st Dec, 1997-2006

2.3.4 Funding for Dialysis Treatment

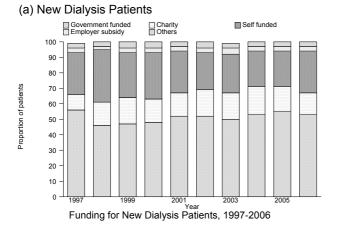
Funding for dialysis treatment in Malaysia is a complicated process. A patient may need to obtain funds from multiple sources for his dialysis treatment. In the initial years of the registry, data for funding for dialysis treatment were mainly from the initial notification of the patient. In 2006, data on funding was included in the annual returns.

The government continues to be the main payer for dialysis therapy. A quarter of patients paid for their dialysis treatment. The proportion of new patients funded by charity organizations ranged between 14-18% over the last 10 years. (table & fig 2.3.4)

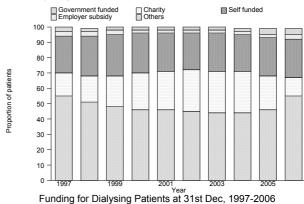
Table 2.3.4: Funding for Dialysis Treatment 1997 – 2006

Year	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006
New Dialysis patients	1136	1252	1544	1840	2085	2348	2597	2858	3054	3152
% by Government	56	46	47	48	52	52	50	53	55	53
% by Charity	10	15	17	15	15	17	17	18	16	14
% self funded	27	34	29	30	27	24	25	23	23	27
% subsidized by Employer	3	2	3	3	3	3	4	3	3	3
% Others	3	3	4	4	3	4	3	3	3	3
Dialysing at 31st December	3389	4194	5151	6246	7305	8497	9766	11136	12693	14062
% by Government	55	51	48	46	46	45	44	44	46	55
% by Charity	15	17	20	24	25	27	27	27	22	12
% self funded	24	26	27	26	25	24	25	24	25	25
% subsidized by Employer	3	3	3	2	2	2	2	2	2	3
% Others	3	2	2	2	2	2	2	2	4	4

Figure 2.3.4: Funding for Dialysis Treatment 1997 – 2006







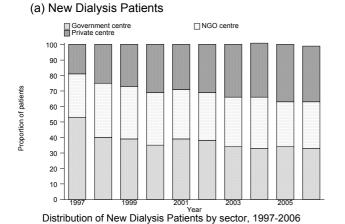
2.3.5 Distribution of dialysis patients by sector

The MOH continued to provide dialysis treatment to one third of new patients, the private sector 36% and the NGO sector 31% in 2006. There was a steady reduction in the proportion of prevalent patients in the MOH from 56% in 1997 to 37% in 2006 with a corresponding increase in the private sector. The proportion of prevalent patients in the NGO sector has remained fairly constant over the years at about 32-33%.

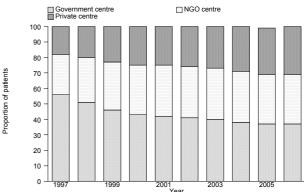
Table 2.3.5: Distribution of Dialysis Patients by Sector 1997 – 2006

Year	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006
New Dialysis patients	1136	1252	1544	1840	2085	2348	2597	2858	3054	3151
% Government centre	53	40	39	35	39	38	34	33	34	33
% NGO centre	28	35	34	34	32	31	32	33	29	31
% Private centre	19	25	27	31	29	31	34	35	37	36
Dialysing at 31st December	3698	4539	5538	6691	7839	9114	10429	11855	13337	14646
% Government centre	56	51	46	43	42	41	40	38	37	37
% NGO centre	26	29	31	32	33	33	33	33	32	32
% Private centre	18	20	23	25	25	26	27	29	30	31

Figure 2.3.5: Distribution of Dialysis Patients by Sector 1997 – 2006



(b) Dialysing patients at 31st December



Distribution of Dialysing Patients at 31st Dec by sector, 1997-2006

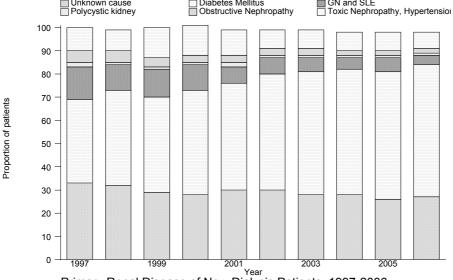
SECTION 2.4: PRIMARY RENAL DISEASE

Diabetes mellitus continues to be the commonest cause of ESRD and the proportion unfortunately has continued to increase and accounted for 57% of new ESRD in 2006. Hypertension was the second commonest known cause of ESRD at about 6%. The proportion of patients with unknown primary renal disease has slowly decreased and was 27% in 2006. Glomerulonephritis as a cause of ESRD has decreased from 13% in 1997 to only 3% in 2006. Systemic lupus erythematosus (SLE) continue to contribute 1% of new ESRD patients.

Table 2.4.1: Primary Renal Disease 1997 – 2006

Year	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006
New Dialysis patients	1136	1252	1544	1840	2085	2348	2597	2858	3054	3152
% Unknown cause	33	32	29	28	30	30	28	28	26	27
% Diabetes Mellitus	36	41	41	45	46	50	53	54	55	57
% GN	13	10	10	9	6	6	5	4	5	3
% SLE	1	1	2	2	1	1	1	1	1	1
% Polycystic kidney	2	1	1	1	2	1	1	1	1	1
% Obstructive Nephropathy	5	5	4	3	3	3	3	2	2	2
% Toxic Nephropathy	0	0	1	0	1	0	0	0	0	0
% Hypertension	9	8	11	12	9	7	7	7	7	6
% Others	1	1	1	1	1	1	1	1	1	1

Figure 2.4.1: Primary Renal Disease for New Dialysis Patients 1997 – 2006 Unknown cause
Polycystic kidney ☐ Diabetes Mellitus
☐ Obstructive Nephropathy GN and SLE
Toxic Nephropathy, Hypertension 100



Primary Renal Disease of New Dialysis Patients, 1997-2006

CHAPTER 3

Death and Survival on Dialysis

Wong Hin Seng
Ong Loke Meng
Wan Shaariah Md Yusuf

SECTION 3.1: DEATH ON DIALYSIS

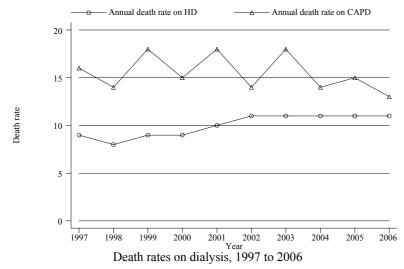
The number of death in dialysis patients for 2006 was 1575 (annual death rate of 11.3%). One thousand four hundred and eleven haemodialysis patients died in 2006 (annual rate of 11.1%) while 164 died on continuous ambulatory peritoneal dialysis (annual death rate of 13.3%).

Year	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006
No. of dialysis patients at risk	3309	4119	5039	6115	7265	8477	9772	11142	12596	13992
Dialysis deaths	316	376	492	594	816	925	1153	1265	1403	1575
Dialysis death rate %	10	9	10	10	11	11	12	11	11	11
No. of HD patients at risk	2838	3599	4471	5487	6552	7635	8784	10056	11460	12762
HD deaths	242	302	392	502	686	810	975	1113	1232	1411
HD death rate %	9	8	9	9	10	11	11	11	11	11
No. of CAPD patients at risk	471	520	568	628	713	842	988	1086	1137	1230
CAPD deaths	74	74	100	92	130	115	178	152	171	164
CAPD death rate %	16	14	18	15	18	14	18	14	15	13

Figure 3.1.1 shows the annual death rate on dialysis from 1997 till 2006. Despite a higher percentage of diabetics (27% in 1997 to 45% in 2006) and elderly patients (in 1997, 41% were aged more than 55 years compared with 53% in 2006) on dialysis in recent years, the overall annual death rate of patients on dialysis remained remarkable unchanged over the last 10 years.

The annual death rate for those on CAPD showed a downward trend in recent years while the annual death rate for those on haemodialysis remained unchanged over the last 5 years. The annual death rate for those on CAPD has decreased by 15% over the last 10 years (from 15.7% in 1997 to 13.3% in 2006) and this has narrowed the difference in the annual death rate between the two modalities of dialysis (from 7.2% in 1997 to 2.3% in 2006).

Figure 3.1.1: Death Rates on Dialysis 1997 – 2006



The causes of death on dialysis are showed in Table 3.1.2. Cardiovascular disease remained the main cause of death in 2006; accounting for 28%. This has remained unchanged over the last 10 years. Death at home accounted for another 21% and a majority of these deaths were probably secondary to cardiovascular events. Death due to sepsis has decreased by 23% over the last 10 years and now accounting for only 13%.

Table 3.1.2: Causes of Death on Dialysis 1997 – 2006

Year	199	97	199	98	199	99	200	00	200)1
	No.	%								
Cardiovascular	85	27	110	29	129	26	177	30	210	26
Died at home	52	16	72	19	107	22	135	23	228	28
Sepsis	53	17	66	18	84	17	85	14	128	16
CAPD peritonitis	5	2	2	1	11	2	21	4	29	4
GIT bleed	4	1	7	2	18	4	18	3	18	2
Cancer	9	3	8	2	6	1	8	1	18	2
Liver disease	3	1	5	1	7	1	14	2	11	1
Others	32	10	55	15	75	15	91	15	108	13
Unknown	73	23	51	14	55	11	45	8	66	8
TOTAL	316	100	376	100	492	100	594	100	816	100
Year	200	02	200)3	200	04	200)5	200)6
	No.	%								
Cardiovascular	307	33	323	28	332	26	350	25	434	28
Died at home	212	23	290	25	304	24	313	22	327	21
Sepsis	141	15	183	16	154	12	159	11	199	13
CAPD peritonitis	16	2	11	1	13	1	18	1	21	1
GIT bleed	24	3	28	2	24	2	28	2	24	2
Cancer	18	2	27	2	19	2	28	2	34	2
Liver disease	16	2	23	2	29	2	25	2	31	2
Others	122	13	186	16	326	26	406	29	385	24
Unknown	69	7	82	7	64	5	76	5	120	8

3.2: PATIENT SURVIVAL ON DIALYSIS

3.2.1 Patient survival by type of dialysis modality

Patient survival by dialysis modalities is showed in Table 3.2.1(a) and Figure 3.2.1(a). The overall unadjusted 5 years and 10 years patient survival on dialysis were 57% and 35% respectively. The unadjusted patient survival was better for those on haemodialysis compared to those on CAPD and this survival difference progressively widened up to 3 years. At 3 years the unadjusted patient survival on haemodialysis was 72% compared 62% in those on CAPD. Beyond 3 years the difference in survival remained at about 10%. Despite adjustment for age and diabetes mellitus [Table 3.2.1(b) and Figure 3.2.1 (b)], patient survival among haemodialysis patients remained superior over CAPD patients. These data contrast with those from the USRDS, Canadian, Australian and the UK registries where patients on PD appeared to have a better survival compared to haemodialysis.

Table 3.2.1(a): Unadjusted patient survival by Dialysis modality, 1997-2006

Dialysis modality		CAPD			HD		All Dialysis			
Interval (months)	No.	% Survival	SE	No.	% Survival	SE	No.	% Survival	SE	
6	2537	93	0	17560	95	0	20097	94	0	
12	2087	88	1	14950	89	0	17037	89	0	
24	1376	76	1	10955	80	0	12331	80	0	
36	847	62	1	7895	72	0	8742	71	0	
48	497	53	1	5592	65	0	6089	64	0	
60	293	47	1	3809	58	0	4102	57	0	
72	160	40	2	2536	52	1	2695	51	1	
84	88	37	2	1578	47	1	1666	46	1	
96	44	31	2	884	43	1	927	42	1	
108	19	29	3	372	39	1	390	38	1	
120	3	27	3	27	35	1	29	35	1	

^{*} No. = Number at risk

SE=standard error

Figure 3.2.1(a): Unadjusted patient survival by Dialysis modality, 1997-2006

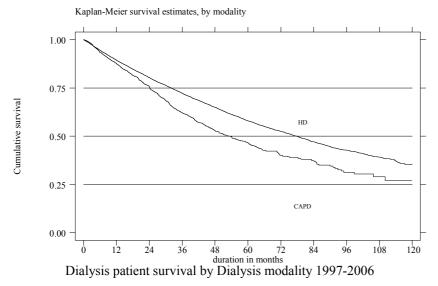


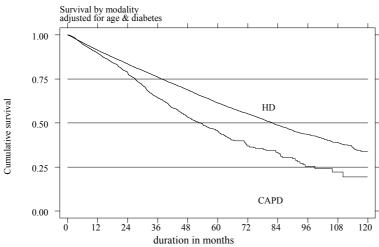
Table 3.2.1(b): Adjusted patient survival with age and diabetes by Dialysis modality, 1997-2006

Dialysis modality		CAPD			HD			All Dialysis	
Interval (months)	No.	% Survival	SE	No.	% Survival	SE	No.	% Survival	SE
6	2537	95	0	17560	96	0	20097	95	0
12	2087	90	1	14950	91	0	17037	91	0
24	1376	79	1	10955	84	0	12331	83	0
36	847	64	1	7895	76	0	8742	75	0
48	497	53	1	5592	69	0	6089	67	0
60	293	46	1	3809	61	0	4102	60	0
72	160	37	2	2536	55	1	2695	54	1
84	88	33	2	1578	49	1	1666	48	1
96	44	25	2	884	44	1	927	43	1
108	19	22	3	372	39	1	390	38	1
120	3	19	3	27	34	1	29	33	1

^{*} No. = Number at risk

SE=standard error

Figure 3.2.1(b): Adjusted patient survival for age and diabetes by Dialysis modality, 1997-2006



Dialysis patient survival by Dialysis modality 1997-2006

3.2.2 Patient survival by year of starting dialysis

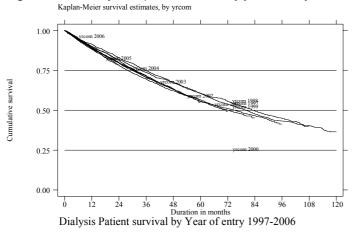
Table 3.2.2 and Fig 3.2.2 show the unadjusted patient survival by year of entry. The unadjusted 6 months survival of those starting dialysis in 2006 was 94%. Despite a progressive increase in the number of diabetic patients and older people starting dialysis in recent years, the unadjusted patient survival remained constant over the last 10 years with a 1-year and 5-year survival of 88-90% and 56-61% respectively.

Table 3.2.2: Unadjusted patient survival by year of entry, 1997-2006

Year		1997			1998			1999			2000	
Interval (months)	No.	% Survival	SE	No.	% Survival	SE	No.	% Survival	SE	No.	% Survival	SE
6	1133	94	1	1244	95	1	1510	95	1	1807	95	1
12	1060	90	1	1178	91	1	1410	90	1	1666	90	1
24	950	82	1	1037	83	1	1213	81	1	1414	80	1
36	835	74	1	913	75	1	1038	72	1	1224	71	1
48	735	67	1	802	68	1	895	63	1	1059	63	1
60	643	61	1	708	61	1	791	57	1	919	56	1
72	555	54	2	637	56	1	706	51	1	802	50	1
84	481	48	2	558	50	1	628	46	1	-	-	-
96	429	43	2	499	45	1	-	-	-	-	-	-
108	390	40	2	-	-	-	-	-	-	-	-	-
120	29	36	2	-	-	-	-	-	-	-	-	-
Year		2001			2002			2003			2004	
Interval (months)	No.	% Survival	SE	No.	% Survival	SE	No.	% Survival	SE	No.	% Surviv al	SE
6	2068	94	1	2350	95	0	2526	94	0	2835	95	0
12	1884	89	1	2175	90	1	2330	89	1	2606	89	1
24	1601	78	1	1842	80	1	2016	80	1	2263	80	1
36	1385	70	1	1598	71	1	1747	70	1	-	-	-
48	1204	62	1	1398	63	1	-	-	-	-	-	-
60	1044	56	1	-	-	-	-	-	-	-	-	-
Year			200)5					200	6		
Interval (months)		lo.	% Sur		SE		No		% Sur		SE	
6 12		968 732	94 88		0 1		166 -	5	94 -		0	_

^{*} No. = Number at risk

Figure 3.2.2: Unadjusted patient survival by year of entry, 1997-2006



SE=standard error

3.2.3 Patient survival by Age at starting dialysis

The unadjusted survival for age groups \leq 14 years, 15-24 years and 25-34 years at the start of dialysis were similar, with a 5-year survival of more than 80% as shown in Table 3.2.3.. Beyond the age of 34 years old the unadjusted survival progressively worsened with increasing age. The 9-year unadjusted survival for those who started dialysis at the age of less than 15 years was 72 % compared with 15% in those more than 64 years of age at the time of initiation of dialysis.

Table 3.2.3: Unadjusted patient survival by age, 1997-2006

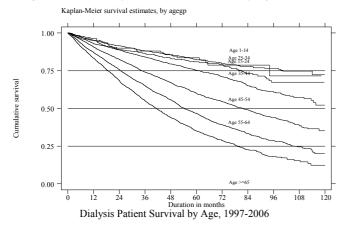
Age group (years)		<=14		15-24				25-34		35-44			
Interval (months)	No.	% Survival	SE	No.	% Survival	SE	No.	% Survival	SE	No.	% Survival	SE	
6	294	98	1	928	97	1	1649	97	0	2832	96	0	
12	258	96	1	793	95	1	1444	95	1	2467	93	0	
24	182	90	2	568	89	1	1111	92	1	1934	89	1	
36	133	88	2	428	86	1	881	88	1	1502	84	1	
48	94	86	3	305	83	1	677	85	1	1146	79	1	
60	58	84	3	219	81	2	493	82	1	850	75	1	
72	39	79	4	157	79	2	373	80	1	601	71	1	
84	23	79	4	103	73	3	253	78	1	381	65	1	
96	11	72	8	59	67	3	155	76	2	233	61	2	
108	6	72	8	32	67	3	69	75	2	101	57	2	
120	-	-	-	4	67	3	4	72	3	6	52	3	

Age group (years)		45-54			55-64			>=65	
Interval (months)	No.	% Survival	SE	No.	% Survival	SE	No.	% Survival	SE
6	5083	96	0	5431	93	0	3881	91	0
12	4329	91	0	4582	87	0	3166	83	1
24	3218	83	1	3231	76	1	2092	69	1
36	2303	74	1	2222	65	1	1279	55	1
48	1643	67	1	1475	56	1	750	44	1
60	1125	60	1	928	47	1	431	35	1
72	750	54	1	554	39	1	230	29	1
84	464	49	1	333	33	1	116	23	1
96	247	44	1	171	29	1	57	18	1
108	99	39	2	70	24	1	19	15	2
120	9	35	2	6	20	2	3	12	2

^{*} No. = Number at risk

SE=standard error

Figure 3.2.3: Unadjusted patient survival by age, 1997-2006



3.2.4 Patient survival by Diabetic status

The unadjusted patient survival among diabetic and non-diabetic patients is shown in Table 3.2.4 and Figure 3.2.4. The presence of diabetes mellitus has major impact on patient survival. The difference in the unadjusted patient survival appeared as early as 6 months after initiation of dialysis and increased with the time on dialysis. The 10 years unadjusted patient survival among diabetics and non-diabetics were 49% and 15% respectively, a three fold difference.

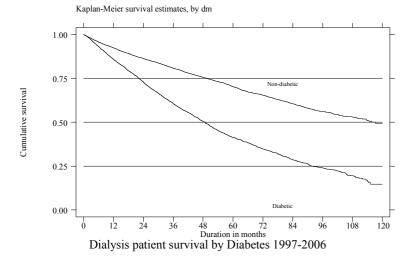
Table 3.2.4: Unadjusted patient survival by Diabetes status, 1997-2006

Diabetes status		Non-Diabetic			Diabetic			
Interval (months)	No.	% Survival	SE	No.	% Survival	SE		
6	10287	96	0	9810	93	0		
12	8967	92	0	8070	86	0		
24	6887	86	0	5444	73	0		
36	5258	81	0	3484	61	1		
48	3908	76	0	2181	51	1		
60	2805	70	1	1297	41	1		
72	1938	65	1	759	35	1		
84	1256	61	1	410	29	1		
96	718	56	1	210	24	1		
108	328	53	1	63	20	1		
120	25	49	1	5	15	2		

^{*} No. = Number at risk

SE=standard error

Figure 3.2.4: Unadjusted patient survival by Diabetes status, 1997-2006



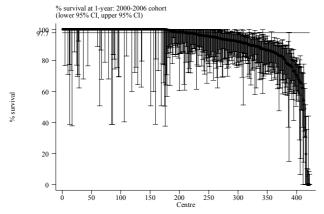
3.3 SURVIVAL OF INCIDENT PATIENTS 2000 – 2006 BY CENTRE

3.3.1. Survival of incident haemodialysis patients 2000 – 2006 by centre

Figure 3.3.1(a) and Figure 3.3.1(b) show the patient survival (adjusted to age and diabetes) by haemodialysis centres at 1 year and at 5 years respectively. The median adjusted patient survival among haemodialysis centres at 1 year and 5 years for the 2000-2006 cohort were 97.7% and 71.9% respectively. There was wide centre variation with regards to patient survival at one year and this became more apparent at 5 years (more than 10 fold different).

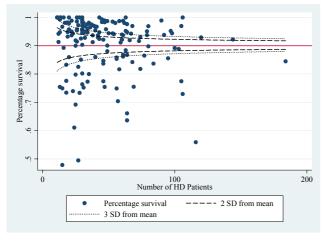
As smaller centres and newer centres tend to have wider confidence interval, data on survival at 1 year and 5 years adjusted for age and diabetes are also shown in funnel plots (Figure 3.3.1(c) and Figure 3.3.1 (d) respectively) to identify outliers. For 1 year survival, 39 (25%) centres lie below 3SD while for 5 years survival 69 (33%) centres are below the 3SD.

Figure 3.3.1(a): Variation in % Survival at 1-year adjusted to age and diabetes, 2000-2006



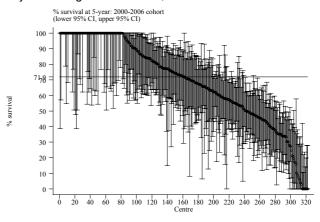
^{*} Horizontal line represents the median % survival among HD centres

Figure 3.3.1(c): Funnel plot for adjusted age at 60 and diabetes at 1 year after 90 days survival; 1997-2002 cohorts (HD centres)



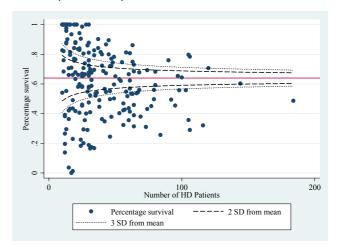
^{*}Centres with new patients <10 were excluded from this analysis.

Figure 3.3.1(b): Variation in % Survival at 5-years adjusted to age and diabetes, 2000-2006



^{*} Horizontal line represents the median % survival among HD centres

Figure 3.3.1(d): Funnel plot for adjusted age at 60 and diabetes at 5 years after 90 days survival; 1997-2002 cohorts (HD centres)



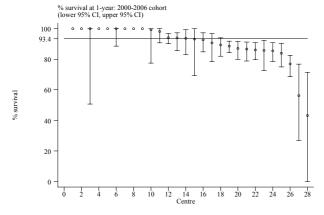
^{*}Centres with new patients <10 were excluded from this analysis.

3.3.2. Survival of incidence CAPD patients 2000 – 2006 by centre

The adjusted patient survival (adjusted to age and diabetes) at 1 year and at 5 years according to CAPD centres are shown in Figure 3.3.2(a) and Figure 3.3.2.(b). The median adjusted patient survival among CAPD centres at one year and 5 years for the 2000-2006 cohort were 93.4% and 51.9% respectively. There was no overt centre variation with regards to patient survival at one year. However the adjusted CAPD patient survival at 5 years demonstrated marked centre variation with a 5 fold difference.

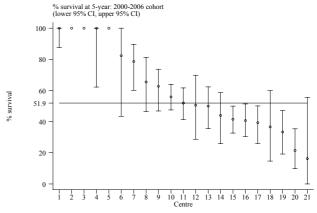
Figure 3.3.2(c) and Figure 3.3.2(d) show the funnel plot for 1 year and 5 years adjusted patient survival among CAPD centres respectively. In both the 1 year and 5 years survival funnel plots, there are 5 centres which lie below the 3SD.

Figure 3.3.2(a): Variation in % Survival at 1-year adjusted to age and diabetes, 2000-2006



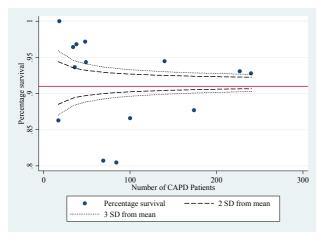
^{*} Horizontal line represents the median % survival among CAPD centres

Figure 3.3.2(b): Variation in % Survival at 5-years adjusted to age and diabetes, 2000-2006



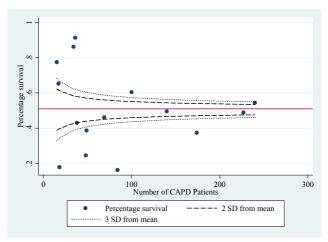
* Horizontal line represents the median % survival among CAPD centres

Figure 3.3.2(c): Funnel plot for adjusted age at 60 and diabetes at 1 year after 90 days survival; 1997-2002 cohorts (CAPD centres)



^{*}Centres with new patients <10 were excluded from this analysis.

Figure 3.3.2(d): Funnel plot for adjusted age at 60 and diabetes at 5 years after 90 days survival; 1997-2002 cohorts (CAPD centres)



*Centres with new patients <10 were excluded from this analysis.

3.4 Adjusted Mortality of dialysis patient

Table 3.4.1 shows the adjusted hazard ratio for mortality of dialysis patients (1997-2006). The 1997-2006 cohort was adjusted for age, gender, primary diagnosis, year commencing dialysis, dialysis modality, body mass index (BMI), serum albumin, serum cholesterol, adequacy of dialysis (KT/V), diastolic blood pressure, haemoglobin, serum calcium, calcium phosphate product, serum phosphate, viral hepatitis status and presence of cardiovascular disease.

Patient characteristics that had significant impact on mortality were age, gender, year commencing dialysis, dialysis modality, BMI, KT/V, diastolic blood pressure and the presence of diabetes or cardiovascular disease. The significant biochemical risk factors for mortality were serum albumin, serum cholesterol, haemoglobin, calcium, calcium phosphate product, phosphate and hepatitis C status.

There were positive correlation between age of patient, diastolic blood pressure (Figure 3.4.1a), serum calcium and serum phosphate (Figure 3.4.1b) with mortality while negative correlation was noted between serum albumin, KT/V (Figure 3.4.1c), haemoglobin concentration (Figure 3.4.1d) and calcium phosphate product with mortality.

Table 3.4.1: Adjusted hazard ratio for mortality of dialysis patients (1997-2006 cohort)

Factors	N	Hazard ratio	95% CI	P value
Age (years):				
0-14(ref.*)	331	1.00		
15-24	1045	1.63	(1.13, 2.34)	0.01
25-34	1856	1.47	(1.03, 2.09)	0.03
35-44	3172	2.06	(1.46, 2.90)	0.00
45-54	5797	2.81	(2.00, 3.95)	0.00
55-64	6301	3.73	(2.66, 5.24)	0.00
<u>></u> 65	4709	5.06	(3.60, 7.12)	0.00
Gender:				
Male (ref.*)	12897	1.00		
Female	10314	0.89	(0.84, 0.94)	0.00
Primary diagnosis:				
Unknown/ Uncertain (ref.*)	6622	1.00		
Diabetes mellitus	11285	1.60	(1.50, 1.70)	0.00
GN/ SLE	1952	0.92	(0.81, 1.03)	0.16
Polycystic kidney	315	1.14	(0.91, 1.45)	0.26
Obstructive Nephropathy	763	1.09	(0.94, 1.25)	0.26
Others	2274	1.06	(0.96, 1.17)	0.24
Year start dialysis:				
1997-8 (ref.*)	2546	1.00		
1999-2000	3563	1.06	(0.98, 1.14)	0.14
2001-2002	4752	1.10	(1.02, 1.19)	0.01
2003-2006	12350	1.10	(1.01, 1.18)	0.02
Modality:				
CAPD	3069	1.35	(1.24, 1.48)	0.00
HD (ref*)	20142	1.00		
BMI:				
<18.5	2257	1.52	(1.37, 1.70)	0.00
18.5-<25	15615	1.34	(1.26, 1.43)	0.00
≥25(ref.*)	5339	1.00		

Table 3.4.1: Adjusted hazard ratio for mortality of dialysis patients (1997-2006 cohort) - cont'd

N	Hazard ratio	95% CI	P value
			
			0.00
			0.00
11046	1.70	(1.59, 1.82)	0.00
7866	1.00		
749	1.11	(0.96, 1.28)	0.17
16703	1.16	(1.10, 1.24)	0.00
5759	1.00		
358	1.55	(1.30, 1.85)	0.00
1486	1.04	(0.94, 1.16)	0.44
4783	1.00		
		(0.83, 0.95)	0.00
		• • •	0.00
		(- ,)	
2548	0.89	(0.01, 0.82)	0.96
		• • •	0.07
		(0.00, 1.00)	0.07
		(1 00 1 21)	0.04
			0.04
707	1.30	(1.07, 2.31)	0.00
2379	3.07	(2.74. 3.45)	0.00
			0.00
			0.00
		(0.97, 1.22)	0.14
		(0.70, 4.00)	2.62
1105	0.92	(0.78, 1.09)	0.32
2225		(0.00.0.55)	•
		(0.80, 0.90)	0.00
			_
646	1.63	(1.43, 1.85)	0.00
7163	1.00	(0.91, 1.10)	0.98
10759	1.00		
3668	0.69	(0.62, 0.77)	0.00
1621	0.57	(0.47, 0.69)	0.00
		•	
7879	0.90	(0.82, 0.98)	0.02
10244	1.00	,	
2190		(0.87, 1.10)	0.76
			0.02
			0.05
		• • •	0.00
• • • • • • • • • • • • • • • • • • • •		(1.3., 2.30)	2.50
22250	1 00		
		(0.08.1.22)	0.12
902	1.08	(0.30, 1.22)	0.12
22424	1.00		
		(0.770.4)	0.00
1090	0.85	(0.77, 0.94)	0.00
18599	1.00		
	7866 749 16703 5759 358 1486 4783 6126 10458 2548 8465 9358 2353 487 2378 3567 10019 4084 2058 1105 6325 16240 646 7163 10759 3668 1621 7879 10244	2952 2.33 11046 1.70 7866 1.00 749 1.11 16703 1.16 5759 1.00 358 1.55 1486 1.04 4783 1.00 6126 0.89 10458 0.80 2548 0.89 8465 0.95 9358 1.00 2353 1.10 487 1.96 10019 2.13 4084 1.09 2058 1.00 1105 0.92 6325 0.85 16240 1.00 646 1.63 7163 1.00 10759 1.00 3668 0.69 1621 0.57 7879 0.90 10244 1.00 2190 0.98 1303 1.19 794 1.23 801 2.05 22259 1.00 952 <td>2952 2.33 (2.14, 2.54) 11046 1.70 (1.59, 1.82) 7866 1.00 (1.59, 1.82) 749 1.11 (0.96, 1.28) 16703 1.16 (1.10, 1.24) 5759 1.00 (1.10, 1.24) 358 1.55 (1.30, 1.85) 1486 1.04 (0.94, 1.16) 4783 1.00 (0.83, 0.95) 6126 0.89 (0.83, 0.95) 10458 0.80 (0.74, 0.86) 2548 0.89 (0.90, 1.00) 9358 1.00 (1.00, 1.21) 487 1.96 (1.67, 2.31) 2378 3.07 (2.74, 3.45) 3567 1.96 (1.75, 2.19) 10019 2.13 (1.92, 2.36) 4084 1.09 (0.97, 1.22) 2058 1.00 1105 0.92 (0.78, 1.09) 6325 0.85 (0.80, 0.90) 16240 1.00 (0.646 1.63 (1.43, 1.85) 7163 1.00 (0.62, 0.77) (0.47, 0.69)</td>	2952 2.33 (2.14, 2.54) 11046 1.70 (1.59, 1.82) 7866 1.00 (1.59, 1.82) 749 1.11 (0.96, 1.28) 16703 1.16 (1.10, 1.24) 5759 1.00 (1.10, 1.24) 358 1.55 (1.30, 1.85) 1486 1.04 (0.94, 1.16) 4783 1.00 (0.83, 0.95) 6126 0.89 (0.83, 0.95) 10458 0.80 (0.74, 0.86) 2548 0.89 (0.90, 1.00) 9358 1.00 (1.00, 1.21) 487 1.96 (1.67, 2.31) 2378 3.07 (2.74, 3.45) 3567 1.96 (1.75, 2.19) 10019 2.13 (1.92, 2.36) 4084 1.09 (0.97, 1.22) 2058 1.00 1105 0.92 (0.78, 1.09) 6325 0.85 (0.80, 0.90) 16240 1.00 (0.646 1.63 (1.43, 1.85) 7163 1.00 (0.62, 0.77) (0.47, 0.69)

ref: Reference group

Figure 3.4.1(a): Adjusted hazard ratio for mortality of dialysis patients by diastolic blood pressure (1997-2006 cohort)

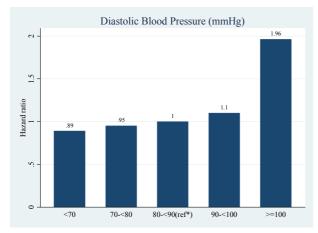


Figure 3.4.1(c): Adjusted hazard ratio for mortality of dialysis patients by KT/V (1997-2006 cohort)

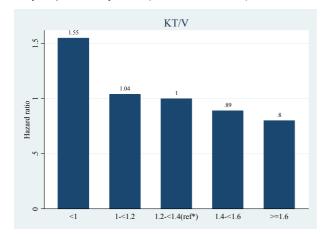


Figure 3.4.1(b): Adjusted hazard ratio for mortality of dialysis patients by serum phosphate (1997-2006 cohort)

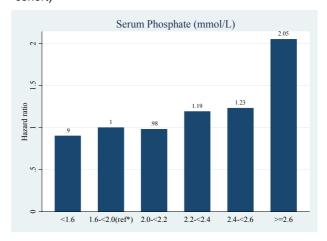
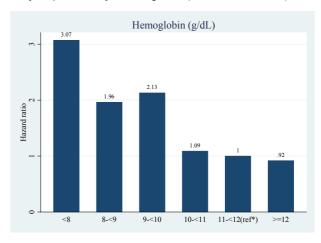


Figure 3.4.1(d): Adjusted hazard ratio for mortality of dialysis patients by haemoglobin (1997-2006 cohort)



CHAPTER 4

Quality of Life & Rehabilitation Outcomes

Liu Wen Jiun Chew Thian Fook Alinda Chiu Sze Fung Zaki Morad B Mohd Zaher

SECTION A: QoL index score

16476 patients who entered dialysis between 1997-2006 were analysed. 13872 HD patients and 2604 CAPD patients reported median QoL index score of 9 and 10 respectively (Table 4.1, Figure 4.1) Diabetics have a lower median QoL index score (8 versus 10) than non diabetics (Table 4.2, Figure 4.2) whilst there was no difference seen between gender (Table 4.3, Figure 4.3). There is a trend of lower median QoL index score being associated with older dialysis patients (Table 4.4, Figure 4.4). There are no obvious trends in QoL index seen either in the HD or CAPD cohort over the last 10 years. (Table 4.5, Table 4.6, Fig 4.5 and Figure 4.6)

Table 4.1: Cumulative distribution of QoL-Index score in relation to Dialysis modality, All Dialysis patients 1997-2006

Dialysis modality	CAPD	HD
Number of patients	2604	13872
Centile		
0	0	0
0.05	5	4
0.10	6	5
0.25 (LQ)	8	7
0.5 (median)	10	9
0.75 (UQ)	10	10
0.90	10	10
0.95	10	10
_1	10	10

Table 4.2: Cumulative distribution of QoL-Index score in relation to Diabetes mellitus, All Dialysis patients 1997-2006

·		
Diabetes mellitus	No	Yes
Number of patients	8764	7712
Centile		
0	0	0
0.05	6	4
0.10	7	5
0.25 (LQ)	9	6
0.5 (median)	10	8
0.75 (UQ)	10	10
0.90	10	10
0.95	10	10
1	10	10

Figure 4.1: Cumulative distribution of QoL-Index score in relation to Dialysis modality, All Dialysis patients 1997-2006

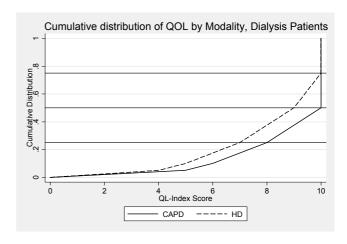


Figure 4.2: Cumulative distribution of QoL-Index score in relation to Diabetes mellitus, All Dialysis patients 1997-2006

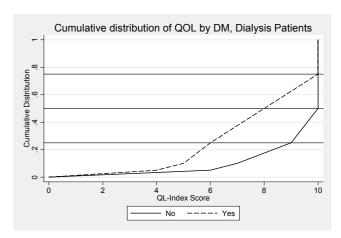


Table 4.3: Cumulative distribution of QoL-Index score in relation to Gender, All Dialysis patients 1997-2006

Gender Female Male Number of patients 9128 7348 Centile 0 0 0 0.05 5 4 0.10 6 5 0.25 (LQ) 8 7 9 0.5 (median) 9 0.75 (UQ) 10 10 0.90 10 10 0.95 10 10 10 10 1

Figure 4.3: Cumulative distribution of QoL-Index score in relation to Gender, All Dialysis patients 1997-2006

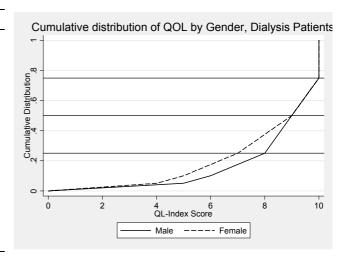


Table 4.4: Cumulative distribution of QoL-Index score in relation to Age, All Dialysis patients 1997-2006

Age group (years)	<20	20-39	40-59	>=60
Number of patients	672	2918	7960	4926
Centile				
0	0	0	0	0
0.05	6	7	5	4
0.10	7	8	6	5
0.25 (LQ)	9	9	8	6
0.5 (median)	10	10	9	8
0.75 (UQ)	10	10	10	9
0.90	10	10	10	10
0.95	10	10	10	10
1	10	10	10	10

Figure 4.4: Cumulative distribution of QoL-Index score in relation to Age, All Dialysis patients 1997-2006

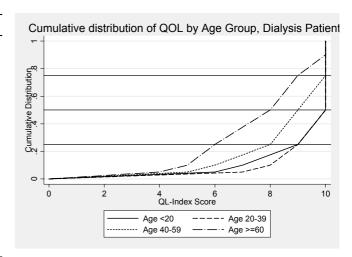


Table 4.5: Cumulative distribution of QoL-Index score in relation to Year of entry, HD patients 1997-2006

Year of Entry	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006
	730	826	1027	1248	1407	1600	1626	1895	1883	1630
Centile										
0	0	0	0	0	0	0	0	0	0	0
0.05	5	5	5	5	5	4	5	4	4	4
0.10	6	6	6	6	5	5	5	5	5	5
0.25 (LQ)	8	8	7	7	7	7	7	7	7	7
0.5 (median)	9	9	9	9	9	9	9	9	9	9
0.75 (UQ)	10	10	10	10	10	10	10	10	10	10
0.90	10	10	10	10	10	10	10	10	10	10
0.95	10	10	10	10	10	10	10	10	10	10
1	10	10	10	10	10	10	10	10	10	10

Figure 4.5: Cumulative distribution of QoL-Index score in relation to Year of entry, HD patients 1997-2006

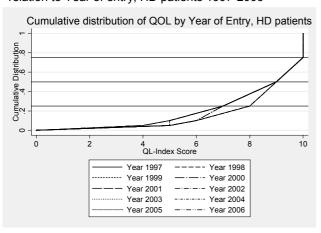


Figure 4.6: Cumulative distribution of QoL-Index score in relation to Year of entry, CAPD patients 1997-2006

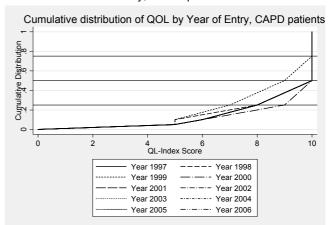


Table 4.6: Cumulative distribution of QoL-Index score in relation to Year of entry, CAPD patients 1997-2006

Year of Entry	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006
Number of patient	162	117	167	188	269	319	369	306	316	391
Centile										
0	0	0	0	0	0	0	0	0	0	0
0.05	5	5	5	5	5	5	5	5	5	5
0.10	6	5	5	6	6	6	6	6	6	6
0.25 (LQ)	8	8	7	9	8	8	8	8	8	8
0.5 (median)	10	10	9	10	10	10	10	10	10	10
0.75 (UQ)	10	10	10	10	10	10	10	10	10	10
0.90	10	10	10	10	10	10	10	10	10	10
0.95	10	10	10	10	10	10	10	10	10	10
1	10	10	10	10	10	10	10	10	10	10

SECTION B: WORK RELATED REHABILITATION

Analysis was done on HD patients (n=5680) and CAPD patients (n=861) who entered dialysis between 1997 -2006, (Table 4.7). Only patients who are working for pay and those who are unable to work for pay due to health reasons are included. The proportion of patients on employment are similar in both modalities (HD = 72% vs CAPD 72%)

Amongst HD as well as CAPD patients, the proportion on employment increases with longer duration on dialysis. (Table 4.8 and Table 4.9) This may be confounded by the healthier individuals who survived longer in the earlier cohort and therefore spuriously increased the proportion on employment

Table 4.7: Work related rehabilitation in relation to Modality, Dialysis patients 1997-2006

Modality	CA	.PD	HD		
	N	%	N	%	
Number of patients	861		5680		
Able to return for Full or Part time for pay	623	72	4077	72	
Unable to work for pay*	238	28	1603	28	

^{*} Exclude patients unable to find employment for non-health related reasons

Table 4.8: Work related rehabilitation in relation to Year of Entry, HD patients 1997-2006

Year		1997	1998	1999	2000	2001	2002	2003	2004	2005	2006
Number of pati	ients	373	421	520	571	574	648	658	708	650	557
Able to return for	No.	310	337	395	443	414	477	461	469	438	333
Full or Part time for pay %	%	83	80	76	78	72	74	70	66	67	60
Unable to	No	63	84	125	128	160	171	197	239	212	224
work for pay*	%	17	20	24	22	28	26	30	34	33	40

^{*} Exclude patients unable to find employment for non-health related reasons

Table 4.9: Work related rehabilitation in relation to Year of Entry, CAPD patients 1997-2006

Year		1997	1998	1999	2000	2001	2002	2003	2004	2005	2006
Number of pat	ients	69	39	47	62	82	116	136	94	109	107
Able to return for	No.	50	31	35	41	67	89	105	68	73	64
Full or Part time for pay	%	72	79	74	66	82	77	77	72	67	60
Unable to	No	19	8	78	21	15	27	31	26	36	43
work for pay*	%	28	21	26	34	27	23	23	28	33	40

^{*} Exclude patients unable to find employment for non-health related reasons

CHAPTER 5

Paediatric Renal Replacement Therapy

Lee Ming Lee
Lynster Liaw
Susan Pee
Wan Jazilah Wan Ismail
Lim Yam Ngo

SECTION A: RRT PROVISION FOR PAEDIATRIC PATIENTS

The paediatric RRT population in this report is defined as patients less than 20 years of age. This report describes the trend of paediatric RRT over the last 10 years.

The number of new patients commencing on dialysis continued to show an increasing trend from 41 in 1997 to 88 in 2006. The dialysis acceptance rate doubled from 4 per million age related population (pmarp) in 1997 to 8 pmarp in 2006 but had plateau at around 7-8 pmarp over the last 5 years. There has been only a modest increase in the number of new transplants in 2006 and the transplant rate remained low at 2 pmarp. The total replacement treatment rate was 8 pmarp.

The number of prevalent dialysis patients continued to rise and by the end of 2006 there were a total of 468 children under 20 on dialysis. The equivalent dialysis prevalence rate increased from 13 pmarp in 1997 to 42 pmarp in 2006. The number of patients with functioning transplants has doubled from 71 in 1997 to 139 in 2006 giving a prevalence rate of 7 and 12 pmarp respectively.

Table 5.01: Stock and Flow of Paediatric Renal Replacement Therapy 1997-2006

Year	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006
New HD patients	21	21	23	12	24	28	33	39	34	46
New CAPD patients	20	28	29	37	39	54	39	41	47	42
New Transplants	14	6	11	14	8	11	11	9	17	19
HD deaths	3	3	2	4	1	11	6	10	9	7
CAPD deaths	3	7	2	3	8	8	9	5	9	15
Transplant deaths	0	0	0	1	0	1	1	0	1	1
On HD at 31st Dec	70	90	106	120	144	162	186	218	241	279
On CAPD at 31st Dec	62	73	91	109	123	152	163	176	192	189
Functioning transplant at 31st December	71	74	83	90	94	102	107	114	125	139

Figure 5.01a: Incident cases of RRT by modality in children under 20 years old, 1997-2006

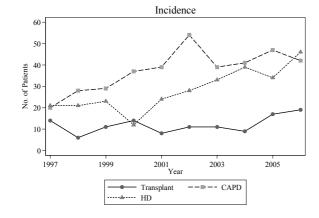


Figure 5.01b: Prevalent cases of RRT by modality in children under 20 years old, 1997-2006

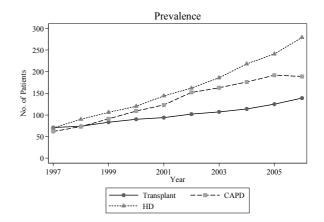
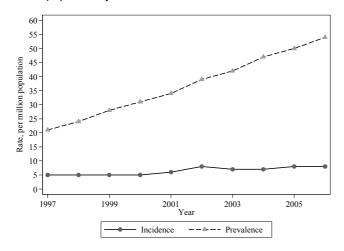


Table 5.02: Paediatric Dialysis and Transplant Rates per million age-group population 1997-2006

Year	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006
Incidence Rate										
New HD	2	2	2	1	2	3	3	4	3	4
New CAPD	2	3	3	4	4	5	4	4	4	4
New Transplant	1	1	1	1	1	1	1	1	2	2
All RRT	5	5	5	5	6	8	7	7	8	8
Prevalence Rate at 31 st December										
On HD	7	9	11	12	14	15	17	20	22	25
On CAPD	6	7	9	11	12	14	15	16	17	17
Functioning Graft	7	8	8	9	9	10	10	10	11	12
All RRT	20	24	28	32	35	39	42	46	50	54

Figure 5.02: Incidence and prevalence rate per million age related population years old on RRT, 1997-2006



SECTION B: DISTRIBUTION OF PAEDIATRIC DIALYSIS

Table 5.03 shows that the treatment rate is consistently higher for states in the west coast of West Malaysia compared to the east coast or East Malaysia.

Table 5.03a: Dialysis Treatment Rate by State, per million state age group population, 1997-2006

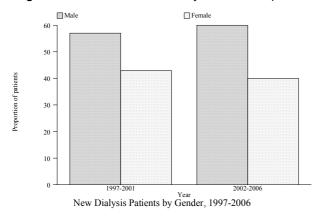
State	1997-2001	2002-2006
Melaka	7	15
Pulau Pinang	6	15
Negeri Sembilan	10	11
Johor Takzim	8	10
Perak Darul Redzuan	3	10
Terengganu Darul Iman	7	9
Kedah & Perlis	8	8
Kelantan Darul Naim	2	8
Pahang Darul Makmur	6	8
Selangor & W. Persekutuan	8	8
Sarawak	5	7
Sabah	2	6

Table 5.03b: New Dialysis Patients by State, 1997-2006

State	1997-2001	2002-2006
Selangor &	80	95
W. Persekutuan	00	00
Johor Darul Takzim	45	60
Perak Darul Redzuan	15	47
Pulau Pinang	15	40
Sabah	13	38
Kedah & Perlis	34	36
Sarawak	21	36
Kelantan Darul Naim	9	31
Pahang Darul Makmur	18	25
Negeri Melaka	9	23
Negeri Sembilan	18	22
Terengganu Darul Iman	17	21

Figure 5.04 shows persistent trend of male predominance amongst the new dialysis and transplant patients. This may be consistent with higher incidence of ESRD among males. However this trend appears more marked among the transplant patients

Figure 5.04: Number of New Dialysis and Transplant Patients by gender 1997-2006



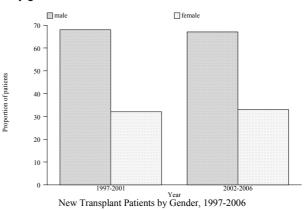


Figure 5.05 shows that except for the age group 15-19 years old which still showed an increasing trend; the treatment rate has begun to level off for all the other age groups. The number of 0-4 year olds provided chronic dialysis treatment remained very low at around 1 pmarp.

Figure 5.05: Dialysis and Transplant Treatment Rate by Age group 1997-2006

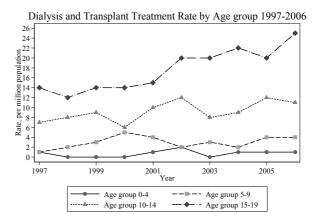


Figure 5.06 shows that CAPD was the preferred mode of initial dialysis modality; however over the last 2 years a significant proportion of children were also started on automated PD as the first modality of dialysis.

Figure 5.06: New Dialysis by treatment modality 1997-2006

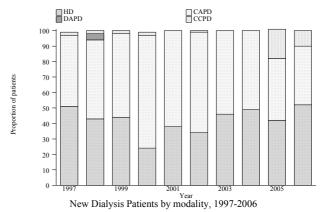
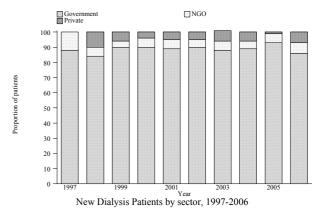


Figure 5.07 shows that up to 90% of children less than 20 years of age received their dialysis treatment from government centres and hence were government funded.

Figure 5.07: New Dialysis by sector 1997-2006



SECTION C: PRIMARY RENAL DIASEASE AND TYPES OF RENAL TRANSPLANTATION

Figure 5.08 shows that glomerulonephritis was the commonest known cause of ESRD accounting for 24% of patients. FSGS on its own accounted for 10% of ESRD. The number of children presenting with ESRD of unknown aetiology was still high at 39%.

Table 5.08: Primary renal disease by sex, 1997-2006

Primary Renal Disease	M	ale	Fen	nale	A	dl .
	N	%	N	%	N	%
Glomerulonephritis	99	25	64	23	163	24
FSGS	38	10	27	10	65	10
Refux nephropathy	23	6	7	3	30	4
SLE	13	3	39	14	52	8
Obstructive uropathy	33	8	9	3	42	6
Renal dysplasia	12	3	6	2	18	3
Others	9	2	10	4	19	3
Hereditary nephritis	11	3	1	0	12	2
Cystic kidney disease	4	1	4	1	8	1
Drug induced nephropathy	0	0	1	0	1	0
Metabolic	3	1	1	0	4	1
Unknown	154	39	108	39	262	39
Total	399	100	277	100	676	100

Table 5.09 shows that living related renal transplantation was still the commonest type of transplantation done accounting for 53% of the transplant done among children under 20 years of age. The second commonest source was cadaveric transplant; up to 24%. Another quarter (23%) of renal transplantation was done overseas under the commercial cadaver and living donor programs.

Table 5.09: Types of Renal Transplant 1997-2006

Year	1997	-2001	2002	-2006
	No.	%	No.	%
Commercial Cadaver	8	15	17	20
Commercial Living donor	3	6	2	3
Living related donor	29	55	42	53
Cadaver	13	25	19	24
Living emotionally related	0	0	0	0
TOTAL	53	100	67	100

SECTION D: SURVIVAL ANALYSIS

Table and figure 5.10 show that renal transplantation has the best patient survival; 98%, 94% and 92% at 1 year, 5 years and 10 years respectively. Patient survival for HD was 95% at 1 year, 84% at 5 years and 75% at 10 years. CAPD patients showed the worst survival; 95% at 1 year and 79% at 5 years. There were too few CAPD patients at 10 years for meaningful analysis. Figure 5.10 shows that patient survival for CAPD and HD were quite comparable up till 3-5 years into dialysis.

Table 5.10: Patient Survival by Modality of RRT, 1997-2006

Modality		Transplant			CAPD			HD		
Interval (years)	No.	% survival	SE	No.	% survival	SE	No.	% survival	SE	
1	91	98	1	306	95	1	272	95	1	
5	41	94	3	64	79	3	81	84	2	
10	2	92	4	-	-	-	2	75	5	

^{*} No. = Number at risk SE = Standard Error

Figure 5.10: Patient Survival by Modality of RRT, 1997-2006

Kaplan-Meier survival estimates, by modality

1.00

0.75

0.75

0.50

0.25

0.00

12

Duration in months
Patient survival by Modality 1997-2006

Table and figure 5.11 show comparable technique survival for both HD and CAPD in the first 2 years of dialysis. After that CAPD showed a progressive deterioration in technique survival compared to HD.

Figure 5.11: Dialysis Technique Survival by Modality, 1997-2006

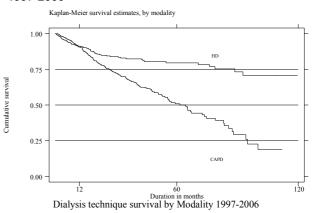


Table 5.11: Dialysis Technique Survival by Modality, 1997-2006

	_ · <i>J</i> · · ·			<u> </u>	j ,	
Modality		CAPD			HD	
Interval (years)	No.	% survival	SE	No.	% survival	SE
1	306	91	2	272	91	2
5	64	51	3	81	80	3
10	-	-	-	2	71	5

^{*} No. = Number at risk SE = Standard Error

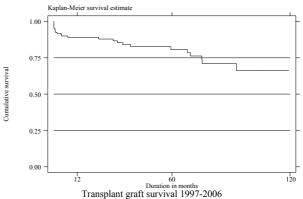
Table and Figure 5.12 below show that the graft survival was 89% at 1 year, 81% at 5 years and 66% at 10 years.

Table 5.12: Transplant Graft Survival 1997-2006

Interval (years)	No.	% survival	SE
1	91	89	3
5	41	81	4
10	2	66	7

^{*} No. = Number at risk SE = Standard Error

Figure 5.12: Transplant Graft Survival 1997-2006



CHAPTER 6

Management of Anaemia In Dialysis Patients

Philip N. Jeremiah Bee Boon Cheak

SECTION 6.1: TREATMENT FOR ANAEMIA IN DIALYSIS

From 1997 – 2006, there was an increasing percentage of patients on erythropoietin (EPO); there were more haemodialysis patients on EPO; 84% compared to 74% in CAPD. The percentage of patients requiring blood transfusion however has also increased.

There were a decreasing number of patients on oral iron, with correspondingly significant increase of patients on parenteral iron.

Table 6.1.1: Treatment for Anaemia, HD patients 1997-2006

Year	No. of subjects	% on Erythropoietin	% received blood transfusion	% on oral Iron	% received parenteral Iron
1997	1695	46	8	92	4
1998	2141	46	13	92	4
1999	2996	51	15	90	5
2000	4392	56	15	88	5
2001	5194	62	13	88	5
2002	6108	67	10	85	7
2003	7017	71	12	83	8
2004	8064	74	11	80	10
2005	9344	81	14	74	11
2006	11494	84	18	76	16

Table 6.1.2: Treatment for Anaemia, CAPD patients 1997-2006

Year	No. of subjects	% on Erythropoietin	% received blood transfusion	% on oral Iron	% received parenteral Iron
1997	476	37	12	96	3
1998	541	44	16	96	3
1999	610	44	14	94	0
2000	662	46	11	92	4
2001	781	45	11	91	2
2002	891	49	11	93	2
2003	1230	53	14	87	4
2004	1312	63	15	85	7
2005	1390	72	12	87	8
2006	1550	74	16	83	13

In 2006, the percentage of patients on EPO among the HD centres varied significantly from 0% to 100%. The median usage of EPO was 88% compared to 45.5% 10 years ago.

Table 6.1.3: Variation in Erythropoietin utilization (% patients) among HD centres, 2006

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	44	7	20	37.5	45.5	62.5	72	92
1998	50	0	4	33	49	57	78	86
1999	76	6	15	42	52	68	82	90
2000	109	0	20	45	56	69	90	100
2001	125	0	19	50	62	75	89	100
2002	156	14	25	55.5	68.5	78	92	100
2003	178	18	35	60	73	82	95	100
2004	207	7	35	65	77	86	96	100
2005	236	4	56	74	84	91	100	100
2006	285	0	58	79	88	93	100	100

% Erythropoietin utilization (lower 95% CI, upper 95% CI)

100

80

40

20

0 20 40 60 80 100 120 140 160 180 200 220 240 260 280 Centre

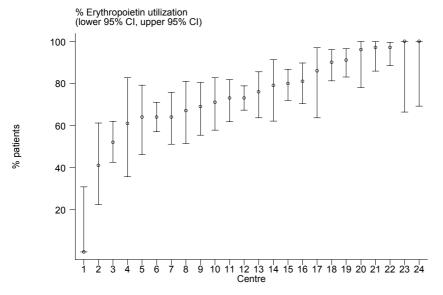
Figure 6.1.3: Variation in Erythropoietin utilization (% patients) among HD centres, 2006

Surprisingly, in CAPD centres too, the variation in EPO utilization was from 0 to 100 %. This was not so in 2005. The median usage of EPO was 74.5% in 2006

Table 6.1.4: Variation in Erythropoietin utilization (% patients) among CAPD centres, 2006

		,	'		` '	,		,
Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	7	19	19	21	41	49	53	53
1998	9	15	15	30	46	55	64	64
1999	10	22	22	32	40.5	54	79	79
2000	11	26	26	33	47	56	67	67
2001	12	25	25	33.5	46.5	56.5	84	84
2002	15	25	25	41	53	56	68	68
2003	19	26	26	38	51	73	92	92
2004	19	5	5	55	63	83	97	97
2005	21	42	56	59	67	81	97	100
2006	24	0	41	64	74.5	90.5	100	100

Figure 6.1.4: Variation in Erythropoietin utilization (%patients) among CAPD centres, 2006



The median weekly EPO dose has increased significantly to 8000 units, in both haemodialysis and CAPD. In HD centres at the 5th and 95th centiles, 5% of centres have their weekly dose at 2000 units and 18,000 units respectively. (Table and Figure 6.1.5 and 6.1.6)

Table 6.1.5: Variation in median weekly Erythropoietin dose (u/week) among HD centres 2006

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	30	2000	2000	4000	4000	4000	6000	8000
1998	34	2000	2000	4000	4000	4000	4000	4000
1999	51	2000	2000	2000	4000	4000	4000	4000
2000	78	2000	2000	2000	4000	4000	4000	6000
2001	93	2000	2000	2000	4000	4000	5000	8000
2002	116	2000	2000	4000	4000	4000	5000	6000
2003	137	2000	2000	4000	4000	4000	6000	8000
2004	171	2000	2000	4000	4000	4000	6000	8000
2005	197	2000	2000	4000	4000	8000	8000	18000
2006	238	2000	2000	8000	8000	8000	18000	18000

Figure 6.1.5: Variation in median weekly Erythropoietin dose (u/week) among HD centres 2006

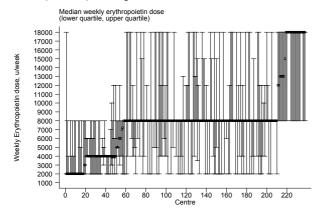


Figure 6.1.6: Variation in median weekly Erythropoietin dose (u/week) among CAPD centres 2006

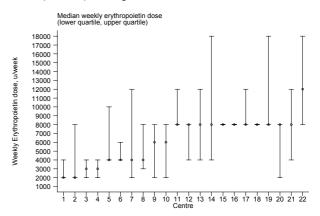


Table 6.1.6: Variation in median weekly Erythropoietin dose (u/week) among CAPD centres 2006

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Cen- tile	Max
1997	6	4000	4000	4000	4000	4000	4000	4000
1998	6	3000	3000	4000	4000	4000	4000	4000
1999	7	2000	2000	2000	4000	4000	4000	4000
2000	9	2000	2000	4000	4000	4000	4000	4000
2001	11	2000	2000	4000	4000	4000	4000	4000
2002	12	2000	2000	3000	4000	4000	4000	4000
2003	14	2000	2000	4000	4000	4000	5000	5000
2004	13	2000	2000	4000	4000	4000	4000	4000
2005	20	2000	2000	3000	4000	4000	8000	8000
2006	22	2000	2000	4000	8000	8000	8000	12000

In HD and CAPD patients, the median requirement of blood transfusion has steadily increased over the last 2 years. (Table and Figure 6.1.7, 6.1.8)

Table 6.1.7: Variation in use of blood transfusion (% patients) among HD centres, 2006

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	44	0	0	0	6.5	13.5	37	71
1998	50	0	0	4	9	16	36	48
1999	76	0	0	4	10	21	41	56
2000	109	0	0	4	11	21	48	75
2001	125	0	0	5	12	20	38	50
2002	156	0	0	2.5	7	14	36	67
2003	178	0	0	3	9	19	36	63
2004	207	0	0	2	8	17	36	50
2005	236	0	0	5	11	20	40	78
2006	285	0	0	10	17	30	50	90

Figure 6.1.7: Variation in use of blood transfusion (% patients) among HD centres, 2006

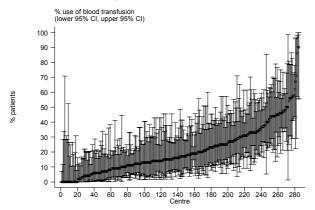


Figure 6.1.8: Variation in use of blood transfusion (% patients) among CAPD centres, 2006

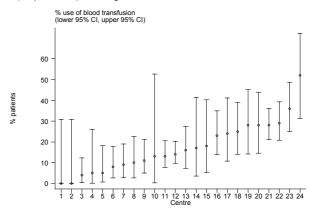


Table 6.1.8: Variation in use of blood transfusion (% patients) among CAPD centres, 2006

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	7	1	1	4	6	29	47	47
1998	9	0	0	7	11	17	47	47
1999	10	0	0	0	6.5	23	47	47
2000	11	0	0	0	8	16	43	43
2001	12	0	0	0	3.5	15.5	38	38
2002	15	0	0	4	7	21	44	44
2003	19	0	0	3	13	21	59	59
2004	19	0	0	5	15	20	37	37
2005	21	0	0	4	10	17	41	43
2006	24	0	0	8.5	15	26.5	36	52

SECTION 6.2: IRON STATUS ON DIALYSIS

In HD and CAPD patients with or without EPO, the mean and median serum Ferritin was steadily increasing over the years except in 2006, when it decreased. This corresponded to an increase in the use of EPO. Up to 98% of patients have serum ferritin of 100 ng/ml. (Table and Figure 6.2.1 to 6.2.4)

Table 6.2.	1: Distribution of Serum Femuli without Erythropoletin, Fib patients 1997 –2006	
	Ni- of	

Year	No of subjects	Mean	Std Dev	Median	LQ	UQ	% Patients ≤100 ng/ml
1997	280	493.1	349.3	435.5	162.5	850.5	86
1998	224	430.8	383.2	297.5	128.4	636.5	80
1999	337	517.9	424.3	402.8	162.8	809.5	86
2000	571	487.5	416.8	363.2	152.5	741	83
2001	758	537.6	453.9	383.5	172	828	87
2002	803	519.5	447.3	373	168.5	781	85
2003	916	551.6	434.2	456.7	190	827.7	87
2004	1044	590.1	463.4	473	218	908.5	89
2005	1012	616.5	498.4	482.5	224.5	901	90
2006	1127	562.7	488.7	407	190	819.5	87

Figure 6.2.1: Cumulative distribution of Serum Ferritin without Erythropoietin, HD patients 1997-2006

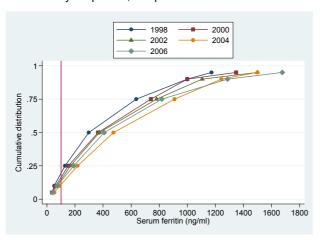


Figure 6.2.2: Cumulative distribution of Serum Ferritin without Erythropoietin, CAPD patients 1997-2006

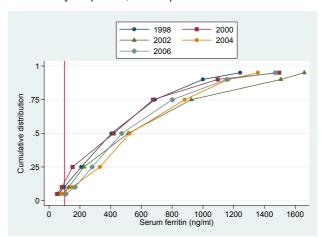


Table 6.2.2: Distribution of Serum Ferritin without Erythropoietin, CAPD patients 1997–2006

Year	No of subjects	Mean	Std Dev	Median	LQ	UQ	% Patients ≤100 ng/ml
1997	133	469	333.5	392	198	718	88
1998	92	492.4	368.3	405	208.2	687.5	87
1999	124	553.7	400.1	499.3	255.3	686.8	94
2000	144	505.9	433.8	420	152.3	675.5	88
2001	223	543.8	417.5	440	216.9	754	91
2002	236	634.8	491.2	514.9	226	924.6	93
2003	329	602.5	429.2	503.7	269	834	93
2004	303	608.4	385.7	522.7	330	882	94
2005	225	651.4	397.8	609	324	913.3	96
2006	262	582.4	407.1	471.7	279.1	801	95

Table 6.2.3: Distribution of Serum Ferritin on Erythropoietin, HD patients 1997 – 2006

Year	No of subjects	Mean	Std Dev	Median	LQ	UQ	% Patients ≤100 ng/ml
1997	471	543.3	347	495.5	219	973	90
1998	328	549.9	382.4	476.5	248	809.8	91
1999	586	560.4	418.6	453	225	829	93
2000	1174	588.3	456.6	475.5	219	860	91
2001	1637	597.5	444.2	491	236	894.2	91
2002	2224	593.1	459.3	464.8	231.3	878.2	91
2003	3134	640.8	428.1	562.8	298	931	94
2004	3902	669.9	460.5	571	306	977	94
2005	5114	683.2	471	599.8	316	972.5	93
2006	6714	641.6	459.3	545	293.8	882	93

Figure 6.2.3: Cumulative distribution of Serum Ferritin on Erythropoietin, HD patients 1997-2006

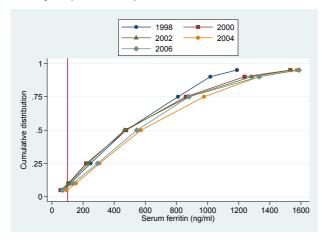


Figure 6.2.4: Cumulative distribution of Serum Ferritin on Erythropoietin, CAPD patients 1997-2006

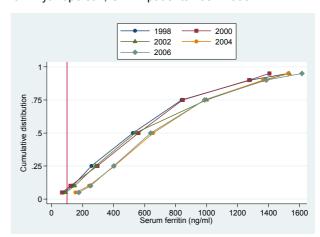


Table 6.2.4: Distribution of Serum Ferritin on Erythropoietin, CAPD patients 1997 – 2006

Year	No of subjects	Mean	Std Dev	Median	LQ	UQ	% Patients ≤100 ng/ml
1997	129	550.8	323.7	496	256	862	93
1998	135	611.2	438.3	524.7	257	839.5	93
1999	136	604.8	436.3	540.6	264.6	870.1	93
2000	180	608.2	416.7	560	295.2	846.3	92
2001	261	645.9	449.2	557.5	275.7	885.4	93
2002	345	666.8	462.4	538.5	284	999.5	94
2003	517	689.9	459.9	589	304	993.2	96
2004	540	728.8	427.2	655.6	406.3	986.7	98
2005	767	732.9	433.6	659	403.6	997.5	97
2006	888	730.5	434.4	639.6	401.3	986.9	98

The median transferrin saturation has remained the same over the last 10 years, with the mean and median always greater than 30%. Up to 95% of all patients have transferrin saturation greater than 20% (Table and Figure 6.2.5 to 6.2.8)

Table 6.2.5: Distribution of transferrin saturation without Erythropoietin, HD patients 1997 -2006

Year	No of subjects	Mean	Std Dev	Median	LQ	UQ	% Patients ≤20%
1997	723	34.1	16.6	29.8	22.7	40.4	84
1998	599	33.3	16.2	29.5	22.1	41.7	82
1999	654	32.9	16.3	29.9	20.9	42.4	78
2000	800	32.7	16.9	28.6	20.9	41.4	78
2001	836	36.9	18.5	32.5	23.9	45.8	84
2002	811	36.5	18.9	32	22.9	45.7	83
2003	921	40.3	18.6	36	27.2	51.1	91
2004	1031	41.2	18.1	37.5	28.5	50.1	92
2005	1110	37.7	17.7	34.4	25.6	46.2	87
2006	1116	36.2	16.8	33	24.6	44.1	87

Figure 6.2.5: Cumulative distribution of transferrin saturation without Erythropoietin, HD patients 1997-2006

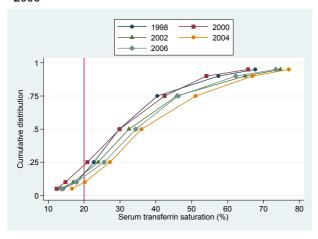


Figure 6.2.6: Cumulative distribution of transferrin saturation without Erythropoietin, CAPD patients 1997-2006

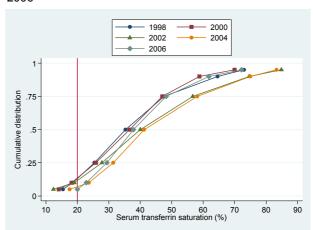


Table 6.2.6: Distribution of transferrin saturation without Erythropoietin, CAPD patients 1997–2006

Year	No of subjects	Mean	Std Dev	Median	LQ	UQ	% Patients ≤20%
1997	246	38.7	17.9	35.3	25.4	47.6	88
1998	184	37.7	15.7	37.3	25.6	47	85
1999	194	37.7	16.2	36.6	25.9	47	88
2000	237	37.9	18.5	34.2	25	48	86
2001	279	43.2	20.8	40	27.8	56.7	89
2002	332	42.7	19.1	38.1	28.3	54.5	92
2003	397	45.2	19.7	41.2	31.4	58.1	93
2004	379	44.5	18.2	41.6	30.9	55.5	98
2005	287	40.6	16.2	37.8	29.4	48.2	95
2006	298	40.5	17.3	38	27.3	46.8	95

Table 6.2.7: Distribution of transferrin saturation on Erythropoietin, HD patients 1997 – 2006

Year	No of subjects	Mean	Std Dev	Median	LQ	UQ	% Patients ≤20%
1997	636	35.9	17.3	31.4	24.2	43.3	87
1998	549	34.9	15.5	32	24.4	42.5	86
1999	703	34.5	16	31.6	23.2	42	85
2000	1247	34.9	16.7	30.4	23	44	84
2001	1634	36.2	17.9	32.3	23.6	45	84
2002	1995	34.6	17.6	30.6	22.2	43.6	81
2003	2642	39.6	18.4	35.9	26.6	48.9	90
2004	3269	39.6	17	36.1	27.8	48.1	93
2005	4804	36.6	17.3	32.8	24.6	45	87
2006	6393	35.2	16.4	31.7	24.1	42.1	87

Figure 6.2.7: Cumulative distribution of transferrin saturation on Erythropoietin, HD patients 1997-2006

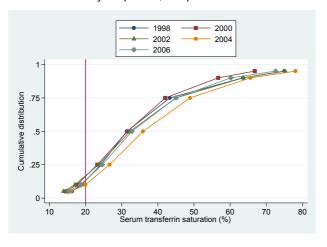


Figure 6.2.8: Cumulative distribution of transferrin saturation on Erythropoietin, CAPD patients 1997-2006

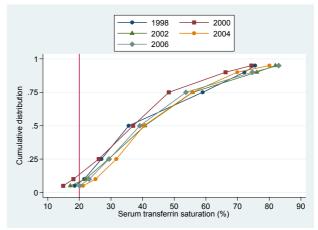


Table 6.2.8: Distribution of transferrin saturation on Erythropoietin, CAPD patients 1997 – 2006

Year	No of subjects	Mean	Std Dev	Median	LQ	UQ	% Patients ≤20%
1997	147	42.2	19.7	35.6	27	59	91
1998	111	39.4	13.8	38.5	28.8	47.4	94
1999	137	38.9	17	37	26.1	48.3	86
2000	238	38.9	18.7	36	24.5	51.1	86
2001	292	44.1	19.6	40.7	29.2	55.8	94
2002	363	43.6	18.6	39.7	30	54.3	94
2003	460	44.6	17.8	40.4	31.7	55.7	96
2004	697	44.7	18.7	40.8	30.8	54.5	96
2005	820	43.5	19.3	39.1	29.4	53.7	95
2006	916	41.7	17.5	38	29.3	50.8	95

From 1997 to 2005, the median for both ferritin and transferrin saturation for all HD centres had increased, except in 2006 when it reduced. There was a wide variation in median ferritin level between HD centres in 2006, ranging from 236 to more than 800 ng/ml. At the median, more than 90% of patients on EPO have serum ferritin levels greater 100 ng/ml and 89% have transferrin saturation greater than 20%. (Table and Figure 6.2.9)

A similar trend, but with higher level of ferritin and transferrin saturation was seen in the CAPD centres. (Table and Figure 6.2.10)

Table 6.2.9: Variation in iron status outcomes among HD centres 2006

a) Median serum ferritin among patients on erythropoietin

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	21	220.5	291.5	414.5	495.5	623.5	792	809.3
1998	13	205	205	432	468.5	560.3	722.8	722.8
1999	22	189.5	202	355.5	419	569	898.8	949.5
2000	41	154	229	387.5	548.3	683.8	813.5	1232
2001	52	217	238.3	393.3	510.4	661.3	883.3	1191.3
2002	71	106.6	192	364.5	475	611.6	875.3	1070.8
2003	98	138	303.8	458.5	542.5	708.5	997	1742.8
2004	123	112	327	463.1	566	735	1001.5	2000
2005	167	1.6	249	458.3	617.5	727.5	965.8	2000
2006	206	1.5	236	417	553.6	688.8	872.6	2000

Figure 6.2.9(a): Variation in median serum ferritin among patients on erythropoietin, HD centres 2006

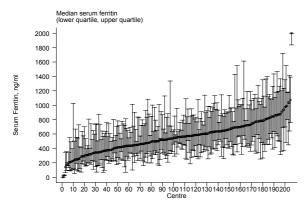
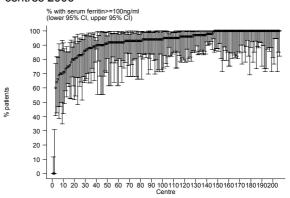


Figure 6.2.9(b): Variation in proportion of patients on erythropoietin with serum ferritin ≥100 ng/ml, HD centres 2006



(b) Proportion of patients on erythropoietin with serum ferritin \geq 100 ng/ml

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	21	73	77	87	91	93	100	100
1998	13	75	75	90	91	95	100	100
1999	22	73	76	92	96	100	100	100
2000	41	70	73	88	94	97	100	100
2001	52	71	73	86.5	93	97	100	100
2002	71	55	73	88	93	97	100	100
2003	98	60	75	91	95	100	100	100
2004	123	52	84	92	96	100	100	100
2005	167	7	79	90	95	100	100	100
2006	206	0	71	92	95	100	100	100

-	(C)	Median	traneferrin	eaturation	amona	nationte	on an	ythropoietin	
- 1	(U)	IVICUIAII	lialisiellili	Saturation	annong	palients	OH CI	y ii ii OpOi c iii i	

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Cen- tile	Max
1997	26	22.6	25.7	29.3	32	37.1	68.5	69.2
1998	22	22.8	24.1	27.8	32.3	35.6	44.4	51.9
1999	26	16.4	20.7	26.9	31.8	34	41.8	44.8
2000	42	16	23.2	27.6	31.5	37.2	44.1	57.5
2001	54	21	22.5	27.1	31.3	37	48.1	76.6
2002	61	14.7	21	25.3	29.8	36	47.5	59.7
2003	91	19.2	24.2	30.9	34.7	41.6	57.3	70.7
2004	114	22.7	27.2	32.7	36.6	41.1	52.7	67.6
2005	150	15.2	23.6	29.1	32.1	37.1	49.5	71.7
2006	191	14.1	23	27.5	31.6	35.7	46.6	77.2

Figure 6.2.9(c): Variation in median transferrin saturation among patients on erythropoietin, HD centres 2006

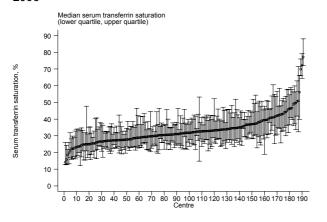
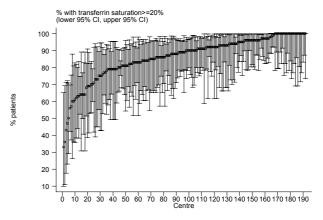


Figure 6.2.9(d): Variation in proportion of patients on erythropoietin with transferrin saturation $\geq 20\%$, HD centres 2006



(d) Proportion of patients on erythropoietin with transferrin saturation >20%

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	26	69	75	82	90	93	100	100
1998	22	57	64	78	88	97	100	100
1999	26	30	57	83	87	94	100	100
2000	42	20	60	77	86	94	100	100
2001	54	53	59	74	87.5	94	100	100
2002	61	33	56	70	82	92	100	100
2003	91	47	69	85	92	100	100	100
2004	114	57	71	91	94	100	100	100
2005	151	29	69	83	91	95	100	100
2006	192	33	61	81	89	96	100	100

Table 6.2.10: Variation in iron status outcomes among CAPD centres 2006

a) Median serum ferritin among patients on erythropoietin

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	4	377.5	377.5	404.8	457.3	530	577.5	577.5
1998	4	418.4	418.4	468.7	534.3	606.3	663	663
1999	5	302.8	302.8	320.4	470	495.6	719.5	719.5
2000	6	335	335	437.3	653.8	745	773	773
2001	9	275.7	275.7	508	581	623	908	908
2002	10	372.2	372.2	437.4	475.6	618	826.5	826.5
2003	12	307.6	307.6	441	508.5	708.3	963.6	963.6
2004	14	312.4	312.4	518	608.3	701.3	1011	1011
2005	18	358.1	358.1	538.2	696.5	807.9	873.8	873.8
2006	20	371	384.1	496.9	623.8	795	885.5	917.6

Figure 6.2.10(a): Variation in median serum ferritin among patients on erythropoietin, CAPD centres 2006

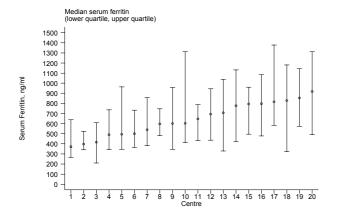
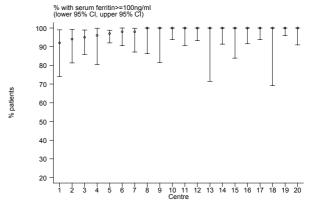


Figure 6.2.10(b): Variation in proportion of patients on erythropoietin with serum ferritin ≥100 ng/ml, CAPD centres 2006



(b) Proportion of patients on erythropoietin with serum ferritin ≥100 ng/ml

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	4	84	84	88.5	93.5	97	100	100
1998	4	83	83	89	97.5	100	100	100
1999	5	84	84	92	95	100	100	100
2000	6	87	87	88	92.5	100	100	100
2001	9	80	80	84	95	100	100	100
2002	10	91	91	92	95	100	100	100
2003	12	86	86	95.5	96	98	100	100
2004	14	90	90	94	99.5	100	100	100
2005	18	85	85	94	97.5	100	100	100
2006	20	92	93	97.5	100	100	100	100

(c) Median transferrin saturation among patients on erythropoietin

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	6	26.7	26.7	27.6	33.6	42.5	70.5	70.5
1998	4	34.2	34.2	35.5	37	41.6	46.2	46.2
1999	6	24	24	27.2	33.6	39.4	42.4	42.4
2000	6	23.1	23.1	26.5	36.3	37.6	52.5	52.5
2001	8	28.4	28.4	31.3	35.9	47.5	79.8	79.8
2002	9	30.5	30.5	36.5	38	40.3	61	61
2003	13	31.9	31.9	35.7	41.9	47.5	64	64
2004	17	29.1	29.1	36.1	40.8	42.8	82.3	82.3
2005	19	31.1	31.1	35.6	38.8	46.2	74.9	74.9
2006	18	32.3	32.3	35.9	37.5	40.5	79.5	79.5

Figure 6.2.10(c): Variation in median transferrin saturation among patients on erythropoietin, CAPD centres 2006

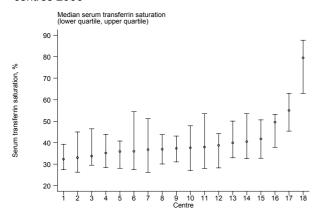
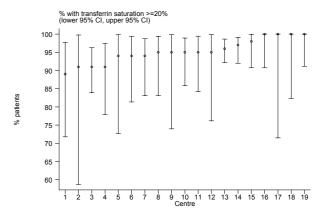


Figure 6.2.10(d): Variation in proportion of patients on erythropoietin with transferrin saturation ≥20%, CAPD centres 2006



(d) Proportion of patients on erythropoietin with transferrin saturation \geq 20%

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	6	70	70	88	90.5	100	100	100
1998	4	81	81	88	95.5	96.5	97	97
1999	6	53	53	84	87.5	94	100	100
2000	6	68	68	74	90	100	100	100
2001	8	85	85	92	93.5	95.5	97	97
2002	9	78	78	92	92	98	100	100
2003	13	90	90	95	96	100	100	100
2004	17	88	88	95	97	100	100	100
2005	19	89	89	94	97	100	100	100
2006	19	89	89	94	95	98	100	100

SECTION 6.3: HAEMOGLOBIN OUTCOMES ON DIALYSIS

The mean and median haemoglobin(Hb) concentrations in all dialysis patients with or without EPO were steadily increasing; in 2006 the mean and median haemoglobin ranged from 10 to 10.9g/dl. The percentage of patients with Hb >10 or >11 g/dl was also steadily increasing for patients not on EPO. However the percentage of patients on EPO, with haemoglobin >10 or >11 g/dl were essentially static.

For dialysis patients without EPO the percentage of patients with the Hb > 10 g/dl ranged from 59% to 70%; whereas the percentage of patient with Hb > 11 gm/dl ranged from 42% to 46%.

For dialysis patients on EPO the percentage of patients with the Hb >10 g/dl was at 50%; whereas the percentage of patient with Hb >11 g/dl was around 27%. (Table and Figure 6.3.1 – 6.3.4)

Table 6.3.1: Distribution of Haemoglobin Concentration without Erythropoietin, HD patients 1997 - 2006

Year	No. of subjects	Mean	Std Dev	Median	LQ	UQ	% Patients ≤ 10 g/dL	% Patients >10 g/dL	% Patients ≤ 11 g/dL	% Patients >11 g/dL
1997	896	9.3	1.9	9	8	10.5	69	31	82	18
1998	1119	9.1	1.9	8.9	7.8	10.3	71	29	83	17
1999	1400	9.1	1.9	8.9	7.8	10.3	70	30	85	15
2000	1754	9.4	2.1	9.1	7.9	10.6	67	33	80	20
2001	1809	9.4	1.9	9.3	8	10.6	64	36	81	19
2002	1795	9.6	2.1	9.4	8.1	10.9	62	38	76	24
2003	1802	9.7	2.1	9.5	8.3	11	60	40	75	25
2004	1927	10.1	2.2	9.9	8.6	11.5	53	47	68	32
2005	1672	10.5	2.3	10.3	8.9	12.1	46	54	62	38
2006	1697	10.6	2.2	10.5	9	12.1	41	59	58	42

Figure 6.3.1: Cumulative distribution of haemoglobin Concentration without Erythropoietin, HD patients 1997-2006

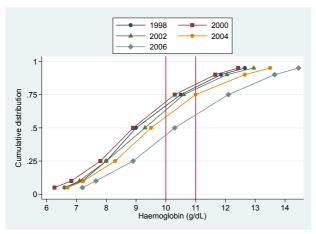


Figure 6.3.2: Cumulative distribution of haemoglobin concentration without Erythropoietin, CAPD patients 1997-2006

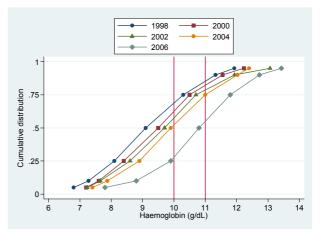


Table 6.3.2: Distribution of Haemoglobin Concentration without Erythropoietin, CAPD patients 1997–2006

Year	No. of subjects	Mean	Std Dev	Median	LQ	UQ	% Patients ≤ 10 g/dL	% Patients >10 g/dL	% Patients ≤ 11 g/dL	% Patients >11 g/dL
1997	297	9.2	1.6	9.1	8.1	10.3	72	28	86	14
1998	301	9.3	1.8	9.2	8.1	10.3	68	32	84	16
1999	336	9.5	1.6	9.5	8.4	10.5	66	34	84	16
2000	342	9.8	1.7	9.7	8.7	10.9	58	42	79	21
2001	405	9.8	1.8	9.7	8.6	10.7	59	41	78	22
2002	434	10	1.8	9.9	8.8	11	54	46	76	24
2003	542	10	1.7	9.9	8.9	11	52	48	76	24
2004	481	10.4	1.6	10.3	9.4	11.4	42	58	67	33
2005	375	10.8	1.6	10.8	9.9	11.8	28	72	60	40
2006	385	10.9	1.6	10.9	10	11.8	25	75	54	46

Table 6.3.3: Distribution of Haemoglobin Concentration on Erythropoietin, HD patients 1997 – 2006

Year	No. of subjects	Mean	Std Dev	Median	LQ	UQ	% Patients ≤ 10 g/dL	% Patients >10 g/dL	% Patients ≤ 11 g/dL	% Patients >11 g/dL
1997	773	8.9	1.6	8.9	7.8	9.9	76	24	90	10
1998	971	9.1	1.6	9.1	7.9	10.2	71	29	88	12
1999	1503	9.2	1.5	9.1	8.1	10.2	71	29	89	11
2000	2332	9.4	1.7	9.4	8.3	10.5	65	35	85	15
2001	3049	9.4	1.6	9.4	8.3	10.5	65	35	85	15
2002	3859	9.5	1.7	9.5	8.4	10.7	62	38	81	19
2003	4782	9.6	1.6	9.6	8.5	10.7	61	39	81	19
2004	5804	9.8	1.6	9.9	8.8	10.9	54	46	77	23
2005	7213	10	1.6	10	8.9	11.1	50	50	73	27
2006	9311	10.1	1.6	10	9	11.1	50	50	73	27

Figure 6.3.3: Cumulative distribution of Haemoglobin Concentration on Erythropoietin, HD patients 1997-2006

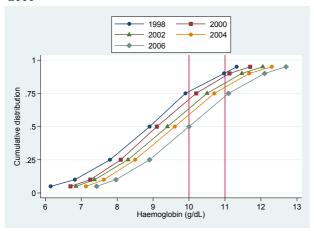


Figure 6.3.4: Cumulative distribution of Haemoglobin Concentration on Erythropoietin, CAPD patients 1997-2006

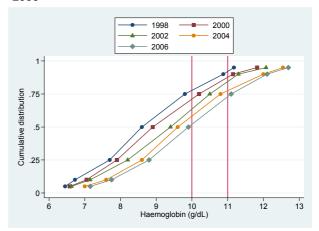


Table 6.3.4: Distribution of Haemoglobin Concentration on Erythropoietin, CAPD patients 1997 - 2006

Year	No. of subjects	Mean	Std Dev	Median	LQ	UQ	% Patients ≤ 10 g/dL	% Patients >10 g/dL	% Patients ≤ 11 g/dL	% Patients >11 g/dL
1997	175	8.8	1.5	8.6	7.7	9.8	79	21	94	6
1998	238	9	1.6	8.8	8	10.1	74	26	88	12
1999	262	9	1.6	8.9	7.9	10.2	73	27	89	11
2000	299	9.4	1.7	9.2	8.1	10.6	65	35	82	18
2001	345	9.3	1.6	9.4	8.2	10.5	65	35	86	14
2002	432	9.4	1.6	9.3	8.4	10.4	69	31	83	17
2003	639	9.7	1.7	9.6	8.6	10.8	59	41	78	22
2004	798	9.8	1.7	9.8	8.6	11	54	46	76	24
2005	970	9.9	1.7	9.9	8.8	11.1	53	47	73	27
2006	1119	10	1.6	10.1	9	11.1	50	50	74	26

In 2006, for HD patients on EPO, the median Hb in HD centres ranged 7.3 to 12.9 g/dl with the median at 10 g/dl. Similar trend was noted in the CAPD centres with lesser variation.

In 2006 for HD patients on EPO, the proportion of patients with Hb >10 g /dl varied between 0 to 89%, with median at 50%. Similarly for patients with Hb > 11g/dl, the range was from 0 to 80% with the median at 24%. This wide variation was not seen in the CAPD patients.

Table 6.3.5: Variation in Haemoglobin outcomes among HD centres 2006 (a) Median haemoglobin level among patients on erythropoietin

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	30	7.8	7.9	8.5	8.9	9.3	10.4	10.6
1998	34	7.6	7.6	8.5	9	9.4	10.2	10.5
1999	51	7.8	8.1	8.6	9	9.6	10.2	10.4
2000	76	7.8	8.2	8.8	9.2	9.8	10.5	14.6
2001	92	8.1	8.4	8.9	9.5	9.9	10.6	12.2
2002	114	8.2	8.5	9	9.4	10.1	10.8	11.4
2003	142	7.8	8.5	9.1	9.6	10	10.7	11.6
2004	179	7.8	8.6	9.2	9.7	10.3	11	11.2
2005	209	8.4	8.8	9.5	10	10.5	11.2	12.2
2006	263	7.3	8.8	9.6	10	10.5	11.2	12.9

Figure 6.3.5(a): Variation in median haemoglobin level among patients on Erythropoietin, HD centres 2006

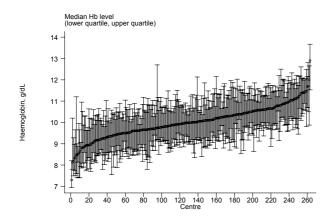
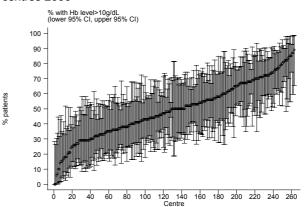


Figure 6.3.5(b): Variation in proportion of patients on erythropoietin with haemoglobin level > 10 g/dL, HD centres 2006



(b) Proportion of patients on erythropoietin with haemoglobin level > 10 g/dL

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	30	0	0	13	21	29	60	82
1998	34	0	0	13	27.5	38	57	71
1999	51	0	5	15	26	38	58	61
2000	76	0	5	20	30.5	43.5	64	97
2001	92	5	12	23.5	33	46	67	100
2002	114	8	15	27	36	50	69	86
2003	142	0	12	27	36	50	68	100
2004	179	8	17	31	41	58	73	86
2005	209	0	20	35	50	63	81	100
2006	263	0	18	36	50	65	80	89

(c) Proportion of patients on erythropoietin with haemoglobin level > 11 g/dL

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	30	0	0	0	7	14	21	33
1998	34	0	0	0	7	17	27	38
1999	51	0	0	0	8	16	29	39
2000	76	0	0	6.5	12.5	20	31	92
2001	92	0	0	8	13	24.5	38	60
2002	114	0	5	12	18	28	46	71
2003	142	0	0	8	15	27	44	61
2004	179	0	0	11	19	29	48	56
2005	209	0	4	14	24	36	54	75
2006	263	0	4	16	24	36	57	80

Figure 6.3.5(c): Variation in proportion of patients on erythropoietin with haemoglobin level > 11 g/dL, HD centres 2006

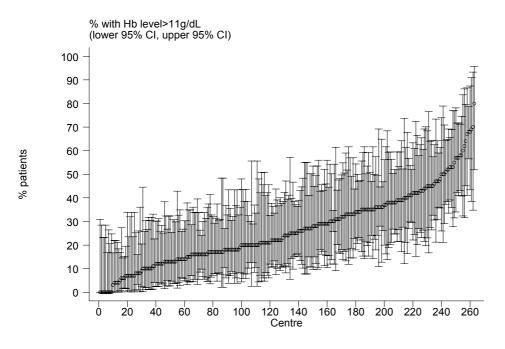


Table 6.3.6: Variation in Haemoglobin outcomes among CAPD centres 2006 (a) Median haemoglobin level among patients on erythropoietin

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	6	7.8	7.8	7.8	8.7	9	9.5	9.5
1998	6	7.9	7.9	8.4	8.9	9.3	9.5	9.5
1999	7	8.1	8.1	8.4	8.7	9.3	9.5	9.5
2000	9	8.2	8.2	8.9	9	9.3	10.3	10.3
2001	11	8.9	8.9	9.2	9.4	9.6	9.7	9.7
2002	12	8.8	8.8	9.1	9.3	9.5	9.9	9.9
2003	16	8.4	8.4	9.3	9.5	9.9	11.2	11.2
2004	17	8.4	8.4	9.2	9.6	10.1	11.2	11.2
2005	20	9	9.1	9.4	9.8	10.4	11	11.1
2006	23	8.8	9.1	9.7	10.1	10.4	10.6	11

Figure 6.3.6(a): Variation in median haemoglobin level among patients on Erythropoietin, CAPD centres 2006

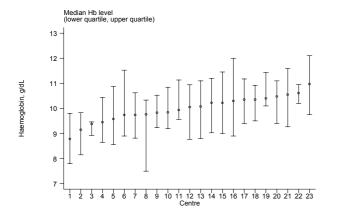
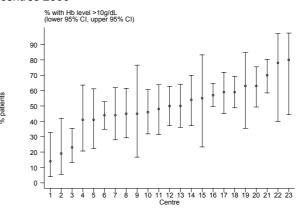


Figure 6.3.6(b): Variation in proportion of patients on erythropoietin with haemoglobin level > 10 g/dL, CAPD centres 2006



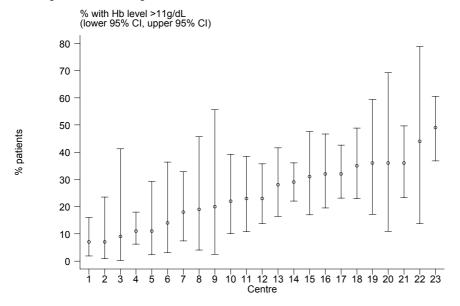
(b) Proportion of patients on erythropoietin with haemoglobin level > 10 g/dL

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	6	0	0	10	19	31	38	38
1998	6	19	19	19	25.5	29	40	40
1999	7	13	13	20	25	36	40	40
2000	9	19	19	30	36	38	55	55
2001	11	24	24	31	38	42	47	47
2002	12	15	15	25.5	31.5	36.5	48	48
2003	16	0	0	27	35	48.5	75	75
2004	17	13	13	39	44	52	72	72
2005	20	21	24.5	35.5	47	57.5	70	76
2006	23	14	19	44	50	59	78	80

(c) Proportion of patients on erythropoietin with haemoglobin level > 11 g/dL

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	6	0	0	0	5.5	8	10	10
1998	6	8	8	8	11	13	16	16
1999	7	0	0	8	9	13	16	16
2000	9	13	13	17	18	20	24	24
2001	11	7	7	10	16	20	23	23
2002	12	11	11	13	16.5	22	27	27
2003	16	0	0	10.5	16.5	22.5	52	52
2004	17	0	0	13	19	29	54	54
2005	20	7	8.5	18.5	28.5	36	55.5	60
2006	23	7	7	14	23	35	44	49

Figure 6.3.6(c): Variation in proportion of patients on erythropoietin with haemoglobin level > 11 g/dL, CAPD centres 2006



CHAPTER 7

Nutritional Status on Dialysis

Ahmad Fauzi Abdul Rahman Tilakavati Karupaiah Winnie Chee Siew Swee

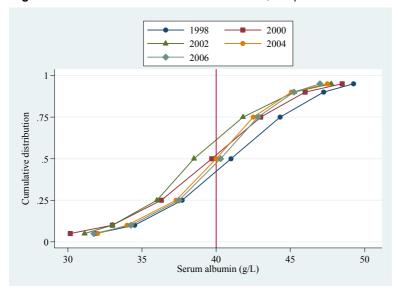
SECTION 7.1: SERUM ALBUMIN LEVELS ON DIALYSIS

Patient numbers increased by 2057 for HD in 2006. Mean serum albumin levels in 2006 was 39.9 g/L, which is just at the borderline for mortality risk (>40 g/L). However, the overall trend for percentage distribution of patients for serum albumin has remained unchanged since 2003. The percentage of well-nourished patients (serum albumin >40g/L) remained above 50% with the rest of the patients mainly in the 35-40g/L range. Percentage of patients with serum albumin levels <35g/L was only 11-18%. Improving trends are also shown from the cumulative distribution graph of serum albumin in HD patients for 2006 (Figure 7.1.1).

Table 7.1.1: Distribution of serum Albumin, HD patients 1997-2006

Year	No. of subjects	Mean	SD	Median	LQ	UQ	% patients <30g/L	% patients 30-<35g/L	% patients 35-<40g/L	% patients ≥40g/L
1997	1644	40.9	6.2	41	37.7	44.3	3	8	30	59
1998	2075	41.2	6.5	41	37.5	44.7	3	9	28	59
1999	2755	39.7	6.1	39.7	36.3	43	4	13	35	49
2000	3733	38.6	7	39	36	42	5	11	41	43
2001	4666	39	5.6	38.5	36	41.8	3	15	44	38
2002	5568	39.2	5.6	39	36.5	42	3	12	42	43
2003	6524	39.9	5.4	40	37.3	42.5	3	9	35	52
2004	7581	39.9	5.3	40	37	42.8	3	10	34	53
2005	8706	40	5.2	40.3	37.5	42.8	3	9	33	56
2006	10763	39.9	5.3	40.3	37.5	42.8	3	9	33	54

Figure 7.1.1: Cumulative distribution of Albumin, HD patients 1997-2006



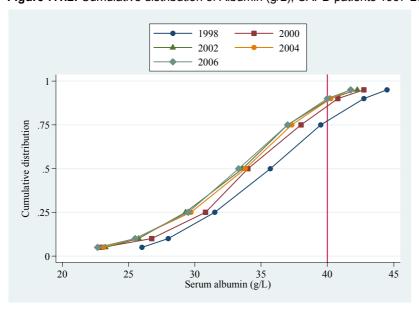
The trend in mean serum albumin levels for patients on CAPD has remained static since 2001 despite initial downward trend noted from 1997-2001. Percentage of patients at increased mortality risk (<35 g/L) also increased initially from 1997 until 2001 (44% in 1997 to 60% by 2001). However the proportion of patients with serum albumin <35g/L has remained static since 2001(56-60%). For the year 2006, mean serum albumin values was 33.5 ± 6.1 g/L. Overall 58% of this population has values less than 35 g/L compared to only 12% at \ge 40g/L. There was a 2% improvement in the number of patients with serum albumin >40g/L in 2006 compared to 2005. (Table 7.1.2)

The cumulative distribution graph in 2006, reflects the trend that percentage of CAPD patients with serum albumin <35 g/L had remained static since 2002 (Figure 7.1.2). The cumulative distribution of serum Albumin values for 2006 confirms that >50% of the population (of 1497) had level below 35g/L and >75% had level below 40g/L.

Table 7.1.2: Distribution of serum Albumin (g/L), CAPD patients 1997-2006

Year	No. of subjects	Mean	SD	Median	LQ	UQ	% patients <30g/L	% patients 30-<35g/L	% patients 35-<40g/L	% patients <u>></u> 40g/L
1997	471	35.7	6.8	35.7	31.5	39.5	16	28	34	22
1998	536	35.8	6.7	36	32	39.7	16	25	35	24
1999	597	34.1	6.6	34	30.8	38	21	33	32	14
2000	640	34.3	6.1	35	31	38.3	20	28	37	14
2001	750	33.3	6.2	33.6	29.3	37	27	33	28	12
2002	862	33.9	5.9	34.3	30.8	37.5	21	35	33	12
2003	1180	33.3	5.8	33.8	29.7	37.3	26	33	30	11
2004	1284	33	6	33.8	29.5	37.3	27	32	30	11
2005	1346	33.2	6.4	33.3	29.5	37	27	33	30	10
2006	1497	33.5	6.1	33.8	30	37	25	33	30	12

Figure 7.1.2: Cumulative distribution of Albumin (g/L), CAPD patients 1997-2006



There is a wide variation in the median for HD centres in the proportion of patients with serum albumin ≥40 g/L from 1997-2006. The trend had been encouraging for the past 4 years whereby the median percentages of serum albumin more than 40g/L is above 50%. For the year 2006, the median was 55%.

The best centre had all (100%) patients achieving serum albumin \geq 40g/L(target albumin). For all HD centres, a greater than 8-fold variation in meeting the albumin target was observed. (Table 7.1.3)

Figure 7.1.3 indicates the wide variation amongst the 272 HD centers reporting the proportion of patients able to achieve the target serum albumin (\geq 40g/L) for the year 2006.

24 CAPD centers participated in 2006. The median percentage of centers with serum albumin 40g/L shows a decreasing trend (28% in 1997 to 16% in 2001 and the latest in 2006 was only 11.5%). The median was 11.5 for the year 2006. The maximum proportion of patients achieving the target serum albumin >40g/L was 74% whilst majority of centers reported less than 40% of patients achieving this target. For all CAPD centres, greater than 36-fold variation in meeting the albumin target was observed. (Table 7.1.4)

Figure 7.1.4 indicates the wide variation amongst 24 CAPD centres reporting the proportion of patients achieving the target serum albumin \geq 40g/L for the year 2006.

Table 7.1.3: Variation in Proportion of patients with serum albumin ≥40g/L among HD centres 2006

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	44	0	10	32.5	61	76.5	95	97
1998	49	7	15	31	58	80	95	96
1999	70	2	8	23	52	65	91	100
2000	94	0	8	22	42.5	62	82	93
2001	116	0	3	17	41	57	82	100
2002	142	0	8	26	44	63	86	100
2003	166	0	17	39	54.5	70	92	100
2004	198	0	10	34	57	73	89	100
2005	226	4	9	43	56	72	88	100
2006	272	0	12	37	55	73	89	100

Figure 7.1.3: Variation in Proportion of patients with serum albumin ≥40g/L, HD centres 2006

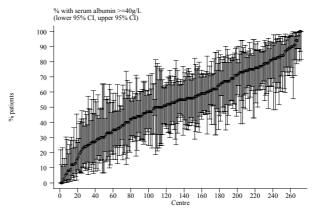
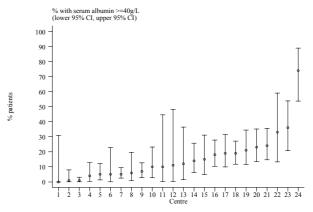


Table 7.1.4: Variation in Proportion of patients with serum albumin ≥40g/L among CAPD centres 2006

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	7	5	5	10	28	29	59	59
1998	9	3	3	18	27	34	45	45
1999	10	2	2	8	14.5	18	29	29
2000	11	0	0	5	13	28	42	42
2001	12	2	2	4.5	16	27	36	36
2002	15	4	4	7	12	21	36	36
2003	19	1	1	9	14	23	55	55
2004	19	1	1	8	13	22	34	34
2005	21	0	1	6	13	22	27	29
2006	24	0	1	5	11.5	20	36	74

Figure 7.1.4: Variation in Proportion of patients with serum albumin ≥40g/L, CAPD centres 2006



SECTION 7.2: BODY MASS INDEX (BMI) ON DIALYSIS

Table 7.2.1 indicates that overall mean BMI for HD patients from 1997 to 2006 was stabilizing at 23. An improving trend in mean BMI [22.9 in 2000 to 23.3 in 2006] was detected despite a 3-fold increase in patient numbers. The percentage of HD patients with BMI \geq 25 increased from 20% in 1997 to 29% in 2006. This may perhaps reflect an increasing number of overweight diabetic patients coming into dialysis or perhaps an improved dietary intake amongst patients.

For the year 2006 and a HD population of 9495, mean BMI value was 23.3 ± 7.7 . Encouragingly, 14% of this group had values less than 18.5 compared to 29% at \geq 25.

Figure 7.2.1 reflects the increasing BMI trends as the curve for 2006 is moving right. About 70% of the population (of 9495) were at BMI of 25

Table 7.2.1: Distribution of BMI, HD patients 1997-2006

Year	No of subjects	Mean	SD	Median	LQ	UQ	% patients <18.5	% patients 18.5-25	% patients >=25
1997	1545	23.6	16.3	21.5	19.1	24.2	19	61	20
1998	1980	24.1	18.3	21.6	19.1	24.3	19	60	21
1999	2708	23.5	15.9	21.4	19.2	24.4	18	61	21
2000	3854	22.9	11.7	21.7	19.3	24.5	18	60	22
2001	4544	23	11	21.9	19.3	24.7	18	59	23
2002	5090	23.2	10.6	22.1	19.5	24.9	16	59	24
2003	5971	23.1	9.7	22.1	19.5	25.1	16	58	26
2004	6749	23.3	9	22.4	19.8	25.4	14	58	28
2005	7799	23.4	9	22.5	19.8	25.6	14	57	29
2006	9526	23.3	7.7	22.5	19.9	25.7	14	56	29

Figure 7.2.1: Cumulative distribution of BMI, HD patients 1997-2006

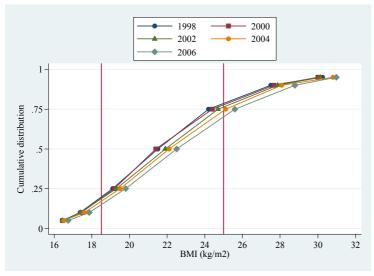


Table 7.2.2. indicates that mean BMI for CAPD patients from 1997 to 2006 was increasing from 22.6 to 23.4 despite a 3-fold increase in patient numbers. The percentage of CAPD patients with BMI \geq 25 increased from 23% in 1997 to 33% in 2006. Patients with BMI of <18.5 had dropped from 21% in year 1997 to 16% in year 2006. This may perhaps reflect an increased number of overweight diabetic patients coming into dialysis.

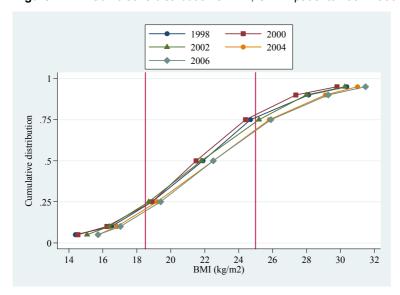
For the year 2006 and a CAPD population of 1413 patients, the mean BMI was 23.4 ± 8.2 and only 16% had BMI <18.5.

Figure 7.2.2 reflects increasing BMI trends as the curve for 2006 is moving right. For the year 2006, only about 65% of the population (of 1413) were at a BMI of <25. This figure is lower than 1998 [population =491] and 2000 [population =602] for which >75% of their populations had BMI below 25.

Table 7.2.2: Distribution of BMI, CAPD patients 1997-2006

Year	No of subjects	Mean	SD	Median	LQ	UQ	% patients <18.5	% patients 18.5-25	% patients >=25
1997	420	22.6	12.5	21.9	18.9	24.7	21	56	23
1998	491	22	11.1	21.3	18.7	24	23	57	20
1999	552	21.7	4.5	21.5	18.9	24.4	22	56	22
2000	602	21.7	4.5	21.5	18.6	24.6	25	53	22
2001	663	22.1	5	21.8	18.7	25.2	23	50	27
2002	750	22.3	4.9	22.1	18.7	25.5	23	47	30
2003	1068	22.9	6.8	22.5	19.2	25.8	20	50	30
2004	1172	23.1	7.2	22.6	19.5	26	18	51	31
2005	1221	23	7.1	22.5	19.4	25.9	19	51	30
2006	1413	23.4	8.2	22.6	19.7	26.1	16	51	33

Figure 7.2.2: Cumulative distribution of BMI, CAPD patients 1997-2006



Less variation was observed for BMI measurements amongst 259 HD centers for 2006.

The median of the participating centers achieving BMI of ≥ 18.5 was 87%. The best centre had all (100%) patients achieving BMI ≥ 18.5 (target), while the worst centre had 55% of patients achieving this target. For all HD centres, there was a 1.4-fold variation in meeting the target BMI (≥ 18.5). (Table 7.2.3)

Figure 7.2.3 indicates the variation amongst 258 HD centers reporting the proportion of patients achieving the target BMI (\geq 18.5) for the year 2006. The center with the least number of patients of BMI >18.5 recorded a percentage of 55%

There was a stable trend in the proportion of CAPD patients with BMI \geq 18.5 from 1997 to 2006. For 23 CAPD centers in 2006, the maximum proportion of patients achieving the target BMI(\geq 18.5) was 93% whilst the worst centres reported 30% of the patients achieving this target. This represented a 2.4-fold difference in variation. (Table 7.2.4)

Figure 7.2.4 indicates that only one center reported the lowest proportion of patients achieving the target BMI \geq 18.5 whilst 18 centers reported higher proportions (>75%).

Table 7.2.3: Variation in Proportion of patients with BMI ≥ 18.5 among HD centres 2006

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	44	62	64	74.5	81	88	100	100
1998	48	58	68	75.5	81.5	86	95	100
1999	72	59	67	78	82	90	95	100
2000	96	55	67	75	82	89	96	100
2001	110	60	68	77	83	89	94	100
2002	131	55	67	78	85	90	100	100
2003	156	55	69	79	84	91	100	100
2004	188	57	68	80	86	91	100	100
2005	208	64	70	81	88	93	100	100
2006	259	55	71	81	87	92	100	100

Figure 7.2.3: Variation in Proportion of patients with BMI ≥18.5, HD centres 2006

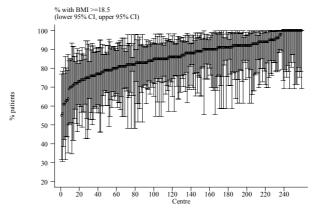
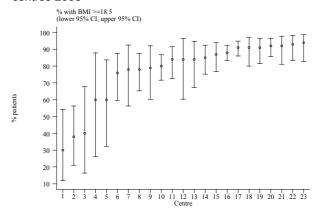


Table 7.2.4: Variation in Proportion of patients with BMI ≥ 18.5 among CAPD centres 2006

	•							
Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	7	50	50	74	81	88	93	93
1998	9	0	0	71	80	89	91	91
1999	9	0	0	71	75	83	93	93
2000	11	11	11	60	76	88	90	90
2001	11	15	15	72	77	88	92	92
2002	15	19	19	65	81	85	87	87
2003	19	17	17	63	81	88	96	96
2004	19	25	25	71	81	89	95	95
2005	19	17	17	67	84	88	91	91
2006	23	30	38	76	84	91	93	94

Figure 7.2.4: Variation in Proportion of patients with BMI ≥18.5, CAPD centres 2006



UQ

0.5

95^{tr}

Centile

5.5

Max

Table 7.2.5 shows that 20 out of 246 HD centres (8.1%) reported patients below BMI of 18.5 and serum albumin of < 30g/L with 7 of these centres (2.8%) at the 95th centile.

Figure 7.2.5 indicates one centre reports 20% of patients falling in this category of severe malnutrition, whilst for 17 of the 20 centres less than 10% of patients were reported in this category

Figure 7.2.5: Variation in Proportion of patients BMI ≤18.5 and serum albumin ≤30 , HD centres 2006

% with BMI <= 18.5 & ALB <= 30 (lower 95% CI, upper 95% CI)

serum albumin ≤30 g/dL among HD centres 2006

Min

No. of

centres

Year

5^{tr}

Centile

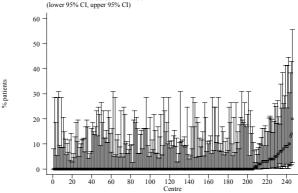


Table 7.2.6 shows that 21 out of 23 CAPD centres (91.3%) reported patients with BMI below 18.5 and serum albumin < 30g/L with 12 of these centres (52.2%) at the 95th centile.

Figure 7.2.6 indicates one centre reported 20% of patients falling in this category of severe of malnutrition, whilst 16 centres reported less than 10% of patients in this category.

Table 7.2.6: Variation in Proportion of patients with BMI <u>≤</u> 18.5 and serum albumin <u>≤</u> 30 g/dL among CAPD centres 2006

Table 7.2.5: Variation in Proportion of patients with BMI ≤ 18.5 and

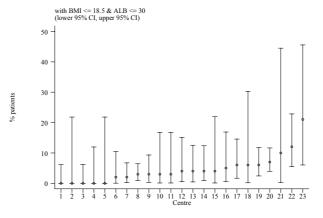
LQ

Median

n

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	7	0	0	3	6	7	13	13
1998	9	0	0	0	0	3	5	5
1999	9	0	0	0	3	6	29	29
2000	11	0	0	2	5	10	10	10
2001	11	2	2	4	6	9	15	15
2002	15	0	0	0	3	7	20	20
2003	19	0	0	0	4	8	25	25
2004	19	0	0	0	3	6	10	10
2005	19	0	0	3	5	8	12	12
2006	23	0	0	2	4	6	12	21

Figure 7.2.6: Variation in Proportion of patients BMI \leq 18.5 and serum albumin \leq 30 g/dL, CAPD centres 2006



CHAPTER 8

Blood Pressure Control And Dyslipidaemia

Prasad Menon Lee Wan Tin

SECTION 8.1: BLOOD PRESSURE CONTROL ON DIALYSIS

In 2006, the predialysis systolic blood pressure remains high and there appears a trend towards increasingly poor blood pressure control predialysis. The mean and median predialysis systolic blood pressure was 151.3 and 151 mmHg respectively.

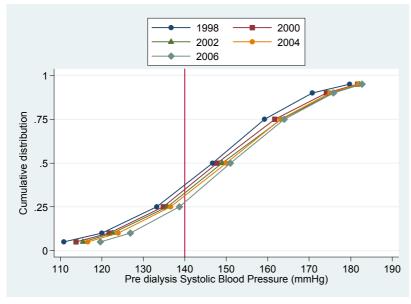
The trend towards an increase in uncontrolled predialysis systolic blood pressure continued in 2006 with the proportion of haemodialysis patients with predialysis systolic BP < 140 mmHg tending lower at 27% compared with 41% a decade ago in 1997. (Table 8.1.1)

This may be a reflection of the increasing numbers of elderly patients coming onto dialysis.

Table 8.1.1: Distribution of Pre dialysis Systolic Blood Pressure (mmHg), HD patients 1997-2006

Year	No. of subjects	Mean	SD	Median	LQ	UQ	% patients <120 mmHg	% patients 120-<140 mmHg	% patients 140-<160 mmHg	% patients 160-<180 mmHg	% patients ≥180 mmHg
1997	1659	144.5	20.8	144.2	130.8	158.1	11	30	35	19	4
1998	2108	146	20.5	146.7	133.2	159.2	10	27	39	19	5
1999	2965	148.7	20.8	148.5	135.3	162.2	8	25	38	23	6
2000	4310	148	20.6	147.8	134.8	161.7	9	25	38	23	6
2001	5147	148.8	20.9	148.8	134.9	162.6	8	25	37	23	7
2002	5911	149.2	20.6	149	135.8	163.3	8	24	38	24	6
2003	6834	149.7	20.2	149.8	136.4	162.9	7	24	39	23	7
2004	7937	149.7	20	150	136.6	163.1	7	23	39	25	6
2005	9221	149.9	19.4	149.6	137	162.8	6	24	40	24	6
2006	11346	151.3	19.3	151	138.7	164	5	22	41	25	7

Figure 8.1.1: Cumulative distribution of Pre dialysis Systolic Blood Pressure (mmHg), HD patients 1997-2006



In contrast, predialysis systolic blood pressure was better controlled in CAPD patients in 2006, with 54% of CAPD patients having a predialysis systolic BP <140 mmHg (Table 8.1.2). This is better than 47% in 2004 and 51% in 2005.

Table 8.1.2: Distribution of Pre dialysis Systolic Blood Pressure (mmHg), CAPD patients 1997-2006

Year	No. of subjects	Mean	SD	Median	LQ	UQ	% patients <120 mmHg	% patients 120-<140 mmHg	% patients 140-<160 mmHg	% patients 160-<180 mmHg	% patients >180 mmHg
1997	468	142.7	20.3	142.9	128.3	156.3	13	31	37	17	3
1998	519	141	21.2	140	126.4	157.5	16	34	29	18	3
1999	576	141	19.8	140	127.2	156	14	35	34	15	2
2000	638	137.2	20.4	136.1	123.3	150	18	39	29	13	2
2001	739	139	20.2	137.5	125.8	151.7	16	38	30	13	3
2002	843	139.8	20.5	140	127.1	151.8	14	36	34	12	4
2003	1154	140.5	20.1	140	126.7	154.1	15	35	32	15	3
2004	1259	141	19.8	140.9	127.4	154.5	13	34	36	14	3
2005	1351	140.4	20.2	139.3	127.3	153.2	13	38	32	14	3
2006	1522	139.4	19.3	138.4	126.7	151.6	14	40	32	11	2

Figure 8.1.2: Cumulative distribution of Pre dialysis Systolic Blood Pressure (mmHg), CAPD patients 1997-2006

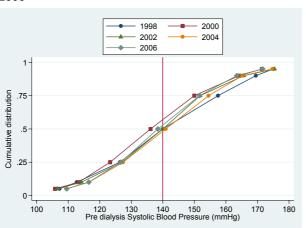
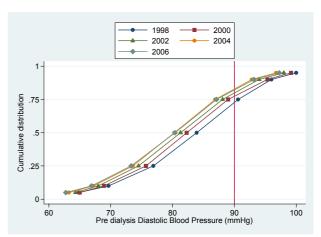


Figure 8.1.3: Cumulative distribution of Pre dialysis Diastolic Blood Pressure, HD patients 1997-2005



In comparison with predialysis systolic blood pressure, the predialysis diastolic blood pressure remained well controlled in haemodialysis patients in 2006, with 83% of haemodialysis patients having predialysis diastolic BP < 90 mmHg (Table 8.1.3). This has been stable over the last few years.

Table 8.1.3: Distribution of Pre dialysis Diastolic Blood Pressure, HD patients 1997-2006

Year	No. of subjects	Mean	SD	Median	LQ	UQ	% patients <70 mmHg	% patients 70-<80 mmHg	% patients 80-<90 mmHg	% patients 90-<100 mmHg	% patients >100 mmHg
1997	1660	83.7	10.9	84.2	77	90.7	10	23	38	22	6
1998	2108	83.5	10.7	83.9	76.9	90.6	10	24	38	23	5
1999	2965	83.5	10.5	83.5	77.1	90	10	24	40	21	6
2000	4309	82.2	10.4	82.3	75.7	89	11	28	39	18	4
2001	5146	81.6	10.4	81.7	75	88.3	12	30	37	17	4
2002	5907	81.2	10.4	81.3	74.5	88.1	13	30	37	16	3
2003	6832	80.6	10.2	80.8	73.9	87.2	14	32	37	14	3
2004	7935	80.3	10.2	80.3	73.6	86.9	15	33	36	14	3
2005	9221	80.3	10.6	80.4	73.5	87	15	32	36	14	3
2006	11345	80.4	11	80.4	73.3	87.1	16	32	35	14	3

Similarly, predialysis diastolic blood pressure was also well controlled in CAPD in 2006 with mean and median predialysis diastolic blood pressure at 81.4 mmHg and 81.5 mmHg respectively. 97% of CAPD patients have predialysis diastolic BP < 90 mmHg (Table 8.1.4).

Table 8.1.4: Distribution of Pre dialysis Diastolic Blood Pressure (mmHg), CAPD patients 1997-2006

Year	No. of subjects	Mean	SD	Median	LQ	UQ	% patients <70 mmHg	% patients 70-<80 mmHg	% patients 80-<90 mmHg	% patients 90-<100 mmHg	% patients ≥100 mmHg
1997	467	85.3	10.6	85.8	79.8	91.4	6	19	41	26	8
1998	519	84.3	11.3	85	77.1	90.1	8	24	36	24	8
1999	576	84	10.9	84.2	77.9	90	9	20	44	20	7
2000	638	82.9	11	83.3	76.6	89.6	10	24	41	20	5
2001	739	83.1	10.9	82.7	76.4	89.6	9	29	38	18	6
2002	843	82.8	10.8	83.4	76.1	90	11	24	41	21	5
2003	1156	82.2	10.9	82.3	75.6	89.4	12	26	38	19	4
2004	1258	82.2	10.5	83	75.4	89.2	11	28	38	18	4
2005	1351	81.6	10.9	82.2	75	88.3	12	29	40	15	5
2006	1521	81.4	10.6	81.5	74.8	88	13	28	40	15	3

Figure 8.1.4: Cumulative distribution of Pre dialysis Diastolic Blood Pressure (mmHg), CAPD patients 1997-2006

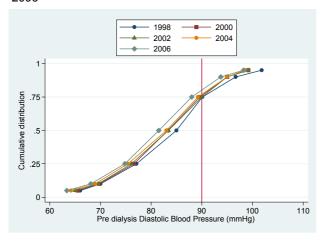
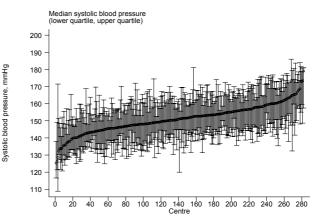


Figure 8.1.5(a): Variation in median systolic blood pressure (mmHg) among HD patients, HD centres 2006



The mild variation in median systolic blood pressure among haemodialysis centers was similar to previous year (Table 8.1.5 (a)).

Table 8.1.5: Variation in BP control among HD centres 2006 a) Median Systolic blood pressure (mmHg) among HD patients

Year	No. of centres	Min	5th Centile	LQ	Median	UQ	95th Centile	Max
1997	44	119.3	133.6	140.2	145.2	152.6	158	163.7
1998	48	131.8	135	141	146.3	150.9	158.7	159.2
1999	75	134.2	137.3	142.2	148.6	154.2	163.6	167.3
2000	107	130.6	136.6	142.7	147.6	155.6	163.2	180
2001	124	127.5	135.7	143	148.6	154.7	161.9	173.4
2002	152	125	137.1	143.5	149.2	154.6	163.4	170.9
2003	174	126.3	135.8	144.8	150.1	155.5	161.6	173.7
2004	204	120	138.1	145	150	155.2	162.5	171
2005	235	127.5	137	143.8	150.5	155.5	161	169.1
2006	281	125.3	137.5	146	151.1	156.3	163.9	179

The mild variation in median diastolic blood pressure among haemodialysis centers was also similar to previous years (Table 8.1.5 (b)).

Table 8.1.5(b) Median Diastolic blood pressure (mmHg) among HD patients

Year	No. of centres	Min	5th Centile	LQ	Median	UQ	95th Centile	Max
1997	44	76.4	80	82	84.2	85.6	88	93.9
1998	48	77.1	78.6	82	83.7	85.9	88.5	90
1999	75	70.9	77.3	81.7	83.7	86	89.4	91.8
2000	107	75.3	76.7	80	82.3	84.7	89.3	94.4
2001	124	73.9	76	79.8	81.9	84	87.5	91.3
2002	152	72.6	75.5	79.2	81.3	83.6	87.3	101.4
2003	174	72.4	74.9	78.5	80.9	83.5	86.7	97.5
2004	204	71.6	74	78.3	80.8	83	86.8	93.4
2005	235	67	73.1	78.1	80.8	82.8	87.1	91.8
2006	281	67	74.4	77.6	80.8	83	87.3	113.5

Figure 8.1.5(b): Variation in median diastolic blood pressure (mmHg) among HD patients, HD centres 2006

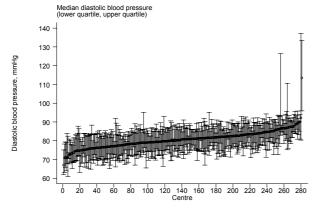
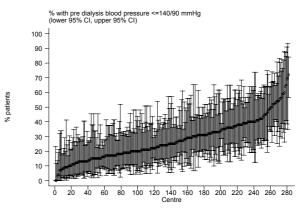


Figure 8.1.5(c): Variation in proportion of HD patients with pre dialysis blood pressure ≤ 140/90 mmHg, HD centres 2006



The variation in proportion of haemodialysis patients with predialysis BP < 140/90 mmHg was also similar to previous years (Table 8.1.5 (c)).

Table 8.1.5(c) Proportion of HD patients with Pre dialysis Blood Pressure < 140/90 mmHq

Year	No. of centres	Min	5th Centile	LQ	Median	UQ	95th Centile	Max
1997	44	15	17	28.5	37.5	44.5	64	87
1998	48	9	20	27	35	42	58	72
1999	75	4	11	23	32	41	56	67
2000	107	0	13	23	32	43	63	83
2001	124	0	11	21	31	43	60	70
2002	152	0	10	22	30	41	56	73
2003	174	4	11	21	29	38	58	80
2004	204	0	10	20	29	37	56	90
2005	235	4	10	20	27	40	57	88
2006	281	0	9	17	25	36	54	72

Similar to haemodialysis centers, the variation in median predialysis systolic blood pressure was mild and similar to previous years in CAPD centers (Table 8.1.6 (a))

Table 8.1.6: Variation in BP control among CAPD centres 2006

(a) Median Systolic blood pressure (mmHg) among CAPD patients

Year	No. of centres	Min	5th Centile	LQ	Median	UQ	95th Centile	Max
1997	7	124	124	139.4	142.5	150	151.6	151.6
1998	9	110.3	110.3	135	138.6	141.6	147.5	147.5
1999	9	116.7	116.7	132.5	137.8	140	152.8	152.8
2000	11	115.5	115.5	131.1	134.9	137.8	149.1	149.1
2001	11	119.6	119.6	130.7	137.5	138.8	149	149
2002	15	123.2	123.2	134.8	140	144.5	148.2	148.2
2003	19	120.7	120.7	131.8	140.8	146.6	151.8	151.8
2004	19	115.9	115.9	133.3	139.5	142.5	149.7	149.7
2005	21	109.9	116.9	134.2	136.6	138.1	147.5	158
2006	24	110.8	116.5	130.6	136.4	141.4	146.2	155.3

Figure 8.1.6(a): Variation in median systolic blood pressure (mmHg) among CAPD patients, CAPD centres 2006

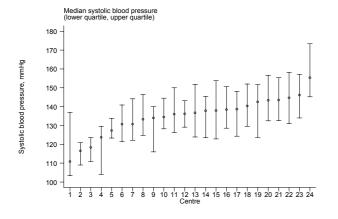
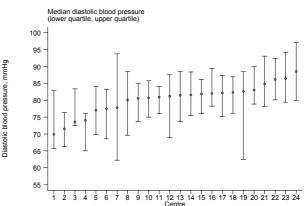


Figure 8.1.6(b): Variation in median diastolic blood pressure (mmHg) among CAPD patients, CAPD centres 2006



The variation in median predialysis diastolic blood pressure was also mild and very similar to previous years (Table 8.1.6 (b)).

Table 8.1.6(b) Median Diastolic blood pressure (mmHg) among CAPD patients

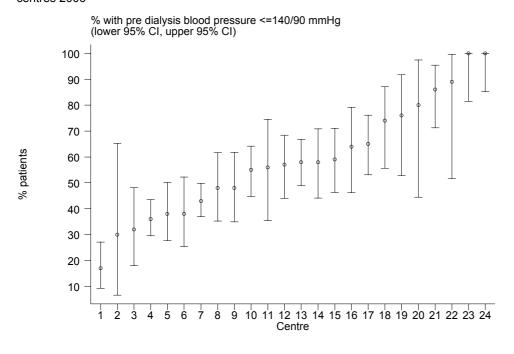
Year	No. of centres	Min	5th Centile	LQ	Median	UQ	95th Centile	Max
1997	7	82.5	82.5	85.3	86	86	88.7	88.7
1998	9	75	75	85.2	85.5	86	87	87
1999	9	77.5	77.5	82.3	84.5	85	86.7	86.7
2000	11	73.3	73.3	80	83	84.4	88	88
2001	11	79	79	80.9	83	84.7	88	88
2002	15	76.8	76.8	81.8	83.3	85.5	89.5	89.5
2003	19	73.6	73.6	81	82.9	84.4	88	88
2004	19	76	76	80.6	83.1	84.4	87.5	87.5
2005	21	64.8	75.5	80	82	83.8	86	86.3
2006	24	69.9	71.5	77.6	81.3	82.4	86.1	88.5

As in previous years, the proportion of CAPD patients with BP < 140/90 mmHg was high suggesting good BP control in CAPD patients. BP control in CAPD was much better than in haemodialysis patients (Table 8.1.6 (c)).

(c) Proportion of CAPD patients with Pre dialysis Blood Pressure ≤140/90 mmHg

Year	No. of centres	Min	5th Centile	LQ	Median	UQ	95th Centile	Max
1997	7	26	26	35	41	46	59	59
1998	9	36	36	44	47	49	100	100
1999	9	30	30	41	52	60	100	100
2000	11	24	24	53	58	65	91	91
2001	11	36	36	46	52	65	84	84
2002	15	19	19	33	46	56	92	92
2003	19	28	28	38	46	66	91	91
2004	19	29	29	40	49	59	82	82
2005	21	23	29	45	55	67	94	100
2006	24	17	30	40.5	57.5	75	100	100

Figure 8.1.6(c): Variation in proportion of CAPD patients with pre dialysis blood pressure ≤ 140/90 mmHg, CAPD centres 2006



SECTION 8.2: DYSLIPIDAEMIA IN DIALYSIS PATIENTS

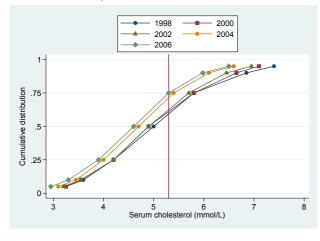
The previous year's trend of improving total cholesterol levels in HD patients continued in 2006, with 76% of HD patients achieving total cholesterol < 5.3 mmol/l (Table 8.2.1).

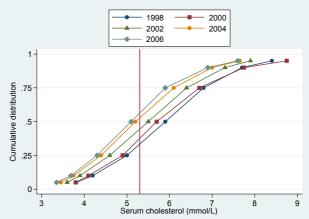
Table 8.2.1: Distribution of serum Cholesterol, HD patients 1997-2006

Year	No. of subjects	Mean	SD	Median	LQ	UQ	% patients <3.5 mmol/L	% patients 3.5-<5.3 mmol/L	% patients 5.3-<6.2 mmol/L	% patients >6.2 mmol/L
1997	1158	5.1	1.4	5.1	4.2	5.9	8	49	24	19
1998	1166	5.1	1.3	5	4.2	5.8	7	53	22	17
1999	1871	5	1.3	4.9	4.1	5.7	10	54	20	15
2000	2956	5	1.2	4.9	4.2	5.8	8	53	23	16
2001	3898	5.1	1.3	4.9	4.2	5.8	8	52	24	16
2002	4751	5	1.2	4.9	4.2	5.7	9	55	24	13
2003	5806	4.8	1.1	4.8	4.1	5.5	9	59	21	11
2004	6710	4.7	1.1	4.7	4	5.4	11	60	21	8
2005	7906	4.7	1.1	4.6	4	5.3	12	61	19	8
2006	9977	4.6	1.1	4.6	3.9	5.3	14	62	17	7

Figure 8.2.1: Cumulative distribution of serum Cholesterol, HD patients 1997-2006







In 2006, as in previous years, total cholesterol levels in CAPD patients was less optimally controlled in comparison with HD patients, with 57% of CAPD patients achieving total cholesterol < 5.3 mmol/l (Table 8.2.2).

Table 8.2.2: Distribution of serum Cholesterol, CAPD patients 1997-2006

Year	No. of subjects	Mean	SD	Median	LQ	UQ	% patients <3.5 mmol/L	% patients 3.5-<5.3 mmol/L	% patients 5.3-<6.2 mmol/L	% patients >6.2 mmol/L
1997	420	6.1	1.4	6	5.1	6.9	2	27	28	43
1998	348	6	1.4	5.9	5	6.8	3	29	28	41
1999	434	5.7	1.4	5.6	4.9	6.4	3	37	30	31
2000	526	5.9	1.6	5.7	4.9	6.7	3	31	30	36
2001	581	5.8	1.4	5.7	4.8	6.6	2	36	27	35
2002	766	5.6	1.4	5.5	4.6	6.4	4	38	28	29
2003	1104	5.4	1.4	5.3	4.4	6.1	5	45	27	23
2004	1230	5.3	1.4	5.2	4.4	6.1	5	48	26	21
2005	1242	5.2	1.3	5	4.3	5.9	5	55	22	18
2006	1394	5.2	1.4	5.1	4.3	5.9	6	51	25	18

In 2006, serum triglyceride control was better in HD patients than CAPD patients, with 75% of HD patients achieving serum triglyceride levels < 2.3 mmol/l (Table 8.2.3) compared to 69% of CAPD patients achieving serum triglyceride level <2.3 mmol/l (Table 8.2.4). This trend of better control of triglyceride levels in HD patients has been demonstrated in previous years.

Table 8.2.3: Distribution of serum Triglyceride, HD patients 1997-2006

Year	No. of subjects	Mean	SD	Median	LQ	UQ	% patients <1.7 mmol/L	% patients 1.7-<2.3 mmol/L	% patients 2.3-<3.5 mmol/L	% patients >3.5 mmol/L
1997	1074	2.1	1.4	1.8	1.3	2.5	45	24	18	12
1998	1089	2.2	1.5	1.8	1.3	2.6	42	26	20	12
1999	1633	2.1	1.3	1.7	1.2	2.5	49	21	18	11
2000	2393	2.1	1.4	1.7	1.3	2.6	48	22	19	12
2001	3162	2.1	1.4	1.7	1.2	2.5	48	22	17	13
2002	3861	2.1	1.4	1.8	1.2	2.5	47	22	18	12
2003	4710	2	1.3	1.7	1.2	2.5	48	23	18	11
2004	5607	2	1.2	1.7	1.2	2.4	51	23	17	10
2005	6950	2	1.3	1.7	1.2	2.4	50	22	18	10
2006	9377	2	1.3	1.6	1.2	2.3	54	21	16	9

Figure 8.2.3: Cumulative distribution of serum Triglyceride (mmol/L), HD patients 1997-2006

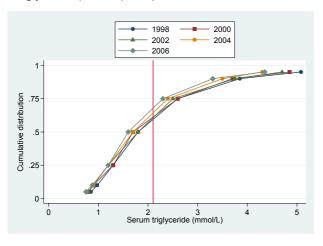


Figure 8.2.4: Cumulative distribution of serum Triglyceride, CAPD patients 1997-2006

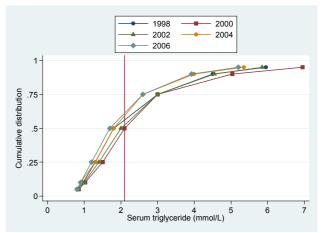


Table 8.2.4: Distribution of serum Triglyceride, CAPD patients 1997-2006

Year	No. of subjects	Mean	SD	Median	LQ	UQ	% patients <1.7 mmol/L	% patients 1.7-<2.3 mmol/L	% patients 2.3-<3.5 mmol/L	% patients ≥3.5 mmol/L
1997	413	2.6	1.9	2.2	1.4	3	36	22	25	18
1998	344	2.4	1.8	1.8	1.3	3	42	22	17	19
1999	421	2.4	1.6	2	1.4	3	38	25	18	19
2000	520	2.7	2.2	2.1	1.5	3	33	24	23	21
2001	576	2.6	1.8	2	1.4	3	36	22	22	20
2002	767	2.5	1.7	2	1.4	3	39	21	22	18
2003	1100	2.3	1.6	1.8	1.2	2.8	45	20	21	14
2004	1223	2.2	1.6	1.8	1.3	2.6	47	23	17	13
2005	1241	2.2	1.5	1.8	1.3	2.7	43	24	18	14
2006	1390	2.2	1.6	1.7	1.2	2.6	47	22	18	13

The mild variation in median serum cholesterol levels and median serum triglyceride levels among HD centers were similar to previous years (Table 8.2.5 (a) and (c)). It is noted that the median of the proportion of patients with serum cholesterol level < 5.3 mmol/l in HD centers has gradually increased from 56% in 1997 to 75% in 2006 (Table 8.2.5 (b)). The median of the proportion of patients with serum triglyceride level <2.3 mmol/l in HD centers has also gradually increased from 65% in 1997 to 70% in 2006 (Table 8.2.5 (d)).

Table 8.2.5: Variation in dyslipidaemia among HD centres 2005 (a) Median serum cholesterol level among HD patients

Year	No. of centres	Min	5th Centile	LQ	Median	UQ	95th Centile	Max
1997	34	4.1	4.3	4.6	5	5.3	5.8	5.9
1998	30	4.2	4.4	4.7	5	5.3	5.5	5.5
1999	47	3.5	4.2	4.6	4.8	5	5.6	5.7
2000	77	4.2	4.3	4.7	4.9	5.2	5.5	5.9
2001	95	4.1	4.4	4.7	5	5.2	5.6	6.3
2002	124	4.4	4.5	4.7	4.9	5.1	5.4	6
2003	150	4.2	4.3	4.6	4.8	5	5.3	5.8
2004	179	3.7	4.1	4.5	4.7	4.9	5.3	6.2
2005	209	3.8	4.2	4.4	4.6	4.8	5.2	5.6
2006	253	3.5	4	4.3	4.6	4.8	5.2	5.6

Figure 8.2.5(a): Variation in median serum cholesterol level among HD patients, HD centres 2006

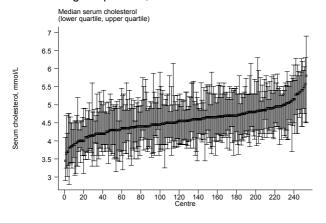
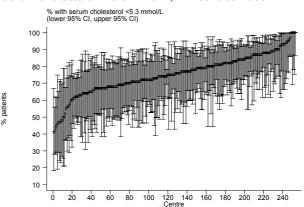


Figure 8.2.5(b): Variation in proportion of patients with serum cholesterol < 5.3 mmol/L, HD centres 2006



(b) Proportion of patients with serum cholesterol < 5.3 mmol/L

Year	No. of centres	Min	5th Centile	LQ	Median	UQ	95th Centile	Max
1997	34	32	35	49	56	65	81	92
1998	30	30	30	51	62	68	86	90
1999	47	38	38	57	64	77	85	89
2000	77	27	36	52	61	68	86	94
2001	95	14	36	53	60	68	79	83
2002	124	32	47	56	64	71	77	94
2003	150	36	46	60	67.5	76	84	92
2004	179	36	47	63	71	78	90	94
2005	209	29	53	65	74	81	89	100
2006	253	41	52	68	75	83	93	100

Table 8.2.5(c) Median serum triglyceride level among HD patients

Year	No. of centres	Min	5th Centile	LQ	Median	UQ	95th Centile	Max
1997	33	1.3	1.3	1.6	1.8	1.9	2.5	2.9
1998	28	1.4	1.5	1.7	1.8	2	2.1	2.3
1999	41	1.2	1.3	1.5	1.7	1.9	2.2	2.5
2000	60	1.2	1.4	1.5	1.8	2	2.6	3
2001	80	1.1	1.4	1.5	1.7	2	2.3	2.5
2002	99	1.1	1.4	1.6	1.8	2	2.5	3.2
2003	128	1.2	1.3	1.5	1.7	1.9	2.2	2.5
2004	157	1	1.3	1.5	1.7	1.8	2.2	2.8
2005	187	0.9	1.3	1.5	1.7	1.9	2.2	2.6
2006	243	0.9	1.2	1.5	1.6	1.8	2.2	2.9

Figure 8.2.5(c): Variation in median serum triglyceride level among HD patients, HD centres 2006

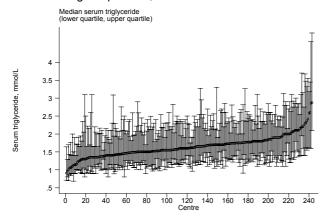


Figure 8.2.5(d): Variation in proportion of patients with serum triglyceride < 2.1 mmol/L, HD centres 2006

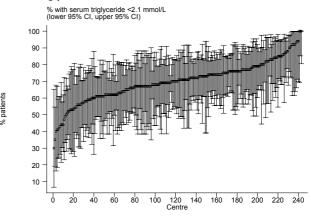


Table 8.2.5(d) Proportion of patients with serum triglyceride < 2.1 mmol/L

Year	No. of centres	Min	5th Centile	LQ	Median	UQ	95th Centile	Max
1997	33	23	36	59	65	77	84	87
1998	28	44	50	56.5	62.5	69.5	82	82
1999	41	36	47	60	67	73	81	91
2000	60	23	36.5	57	65.5	73.5	83	85
2001	80	38	44.5	55.5	65	75	83	90
2002	99	0	42	54	65	72	81	94
2003	128	27	44	56.5	67	75.5	89	100
2004	157	21	47	60	69	78	86	94
2005	187	33	45	60	67	73	83	94
2006	243	30	50	62	70	76	89	100

In 2006 the variation in median serum cholesterol levels and median serum triglyceride levels among CAPD centers were similar to previous years (Table 8.2.6 (a) and (c)). It is noted that the median of the proportion of patients with cholesterol level < 5.3 mmol/l in CAPD centers has gradually increased markedly from 29% in 1997 to 59% in 2006 (Table 8.2.6 (b)). The median of the proportion of patients with serum triglyceride level 2.3 mmol/l in CAPD centers has also gradually increased from 52% in 1997 to 61.5% in 2006 (Table 8.2.6 (d)).

Table 8.2.6: Variation in dyslipidaemia among CAPD centres 2006 (a) Median serum cholesterol level among CAPD patients

Year	No. of centres	Min	5th Centile	LQ	Median	UQ	95th Centile	Max
1997	6	5.8	5.8	5.9	5.9	6.1	6.1	6.1
1998	6	4.8	4.8	5.6	5.8	6.1	6.2	6.2
1999	8	5.1	5.1	5.4	5.7	5.8	6	6
2000	10	5.2	5.2	5.4	5.6	5.9	6.4	6.4
2001	10	5	5	5.6	5.9	6.1	6.2	6.2
2002	15	4.9	4.9	5.4	5.5	5.8	6.3	6.3
2003	19	4.5	4.5	4.9	5.3	5.8	6.1	6.1
2004	19	4.6	4.6	4.9	5.3	5.5	6.2	6.2
2005	19	4.4	4.4	4.7	5	5.3	6	6
2006	23	4.2	4.5	4.8	5.1	5.4	6.1	6.2

Figure 8.2.6(a): Variation in median serum cholesterol level among CAPD patients, CAPD centres 2006

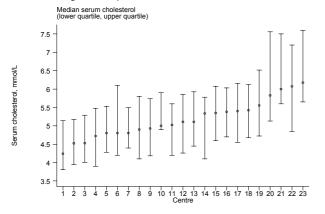


Figure 8.2.6(b): Variation in proportion of patients with serum cholesterol < 5.3 mmol/L, CAPD centres 2006

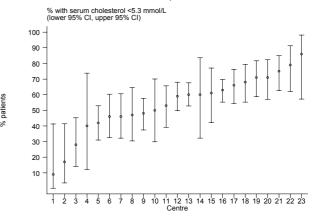


Table 8.2.6(b) Proportion of patients with serum cholesterol < 5.3 mmol/L

Year	No. of centres	Min	5th Centile	LQ	Median	UQ	95th Centile	Max
1997	6	27	27	27	29	31	33	33
1998	6	24	24	27	32	37	53	53
1999	8	10	10	36.5	39.5	45	56	56
2000	10	11	11	18	31	46	54	54
2001	10	22	22	30	34.5	45	63	63
2002	15	16	16	33	42	46	80	80
2003	19	8	8	35	47	58	82	82
2004	19	9	9	40	50	60	72	72
2005	19	28	28	50	60	70	75	75
2006	23	9	17	46	59	68	79	86

Table 8.2.6(c) Median serum triglyceride level among CAPD patients

Year	No. of centres	Min	5th Centile	LQ	Median	UQ	95th Centile	Max
1997	6	1.7	1.7	1.9	2.1	2.2	2.4	2.4
1998	6	1.2	1.2	1.5	1.7	1.9	2.1	2.1
1999	8	1.6	1.6	1.9	2	2.1	2.6	2.6
2000	10	1.8	1.8	2	2.3	2.5	2.6	2.6
2001	10	1.5	1.5	1.9	2	2.1	3	3
2002	15	1.5	1.5	1.8	1.9	2	2.4	2.4
2003	19	1.4	1.4	1.7	1.8	2	2.3	2.3
2004	19	1.3	1.3	1.6	1.8	1.8	2.1	2.1
2005	19	1.3	1.3	1.6	1.9	2	2.2	2.2
2006	22	1.2	1.4	1.6	1.8	1.9	2.4	2.5

Figure 8.2.6(c): Variation in median serum triglyceride level among CAPD patients, CAPD centres 2006

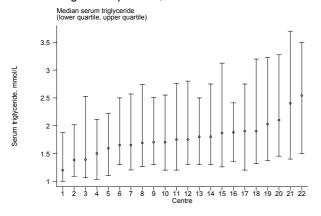


Figure 8.2.6(d): Variation in proportion of patients with serum triglyceride < 2.1 mmol/L, CAPD centres 2006

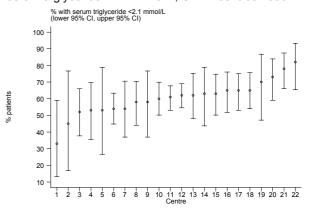


Table 8.2.5(d) Proportion of patients with serum triglyceride < 2.1 mmol/L

	` ' '	•	•	•				
Year	No. of centres	Min	5th Centile	LQ	Median	UQ	95th Centile	Max
1997	6	40	40	46	52	56	61	61
1998	6	51	51	55	61	70	85	85
1999	8	37	37	53.5	56	60.5	64	64
2000	10	18	18	42	49	54	62	62
2001	10	27	27	50	53	58	68	68
2002	15	37	37	52	53	57	78	78
2003	19	46	46	54	58	62	91	91
2004	19	50	50	60	62	69	89	89
2005	19	38	38	53	60	69	89	89
2006	22	33	45	54	61.5	65	78	82

CHAPTER 9

Management of Renal Bone Disease in Dialysis Patients

Fan Kin Sing Rozina Ghazalli Ching Chen Hua Liew Yew Fong

SECTION 9.1: TREATMENT OF RENAL BONE DISEASE

In 2006, no major changes were found in the treatment of renal bone disease. Majority of HD patients (92%) and CAPD patients (84%) received calcium carbonate as phosphate binder. From 1997 to 2006, calcium carbonate remained as the main phosphate binder used in both HD and CAPD patients. The percentage of patients on aluminium phosphate binders remained low in both groups of patients. The percentage of patients on Vitamin D therapy was increasing in HD patients since 2001. The number of parathroidectomies done in 2006 was 151 compared to 43 in 2005. The data regarding parathyroidectomies prior to 2005 is not available. (tables 9.1.1 & 9.1.2)

Table 9.1.1: Treatment for Renal Bone Disease, HD patients 1997-2006

Year	No. of subjects	No. of subjects on CaCO ₃	% on CaCO ₃	No. of subjects on Al(OH) ₃	% on Al(OH) ₃	No. of subjects on Vitamin D	% on Vitamin D	No. of subjects with parathyroid ectomy	% with parathyroid ectomy
1997	1695	1543	91	417	25	694	41	NA	NA
1998	2141	1956	91	343	16	652	30	NA	NA
1999	2996	2693	90	244	8	770	26	NA	NA
2000	4392	3977	91	239	5	1084	25	NA	NA
2001	5194	4810	93	145	3	1145	22	NA	NA
2002	6108	5536	91	171	3	1375	23	NA	NA
2003	7018	6425	92	118	2	1690	24	NA	NA
2004	8164	7408	91	106	1	2029	25	NA	NA
2005	9351	8568	92	98	1	2555	27	43	0
2006	11497	10600	92	69	1	3712	32	151	1

NA = not available

Table 9.1.2: Treatment for Renal Bone Disease, CAPD patients 1997-2006

Year	No. of subjects	No. of subjects on CaCO ₃	% on CaCO₃	No. of subjects on Al(OH) ₃	% on Al(OH) ₃	No. of subjects on Vitamin D	% on Vitamin D	No. of subjects with parathyroid ectomy done	% with parathyroid ectomy done
1997	476	393	83	57	12	114	24	NA	NA
1998	541	425	79	46	9	110	20	NA	NA
1999	610	450	74	36	6	75	12	NA	NA
2000	662	522	79	15	2	96	15	NA	NA
2001	781	588	75	5	1	84	11	NA	NA
2002	891	713	80	6	1	130	15	NA	NA
2003	1231	1039	84	10	1	238	19	NA	NA
2004	1327	1124	85	18	1	304	23	NA	NA
2005	1398	1186	85	13	1	314	22	4	0
2006	1550	1321	85	7	0	375	24	15	1

NA = not available

SECTION 9.2: SERUM CALCIUM AND PHOSPHATE CONTROL

The median corrected serum calcium level remained at 2.3 mmol/l in HD patients (table 9.2.1 & fig. 9.2.1) and 2.4 mmol/l in CAPD patients (table 9.2.2 & fig 9.2.2). In 2006, 50% of patients on HD and 38% of CAPD patients had achieved the target serum calcium level of 2.1 to 2.37 mmol/l as advocated by K/DOQI guidelines.

Table 9.2.1: Distribution of corrected Serum Calcium, HD patients 1997-2006

Year	No. of Subjects	Mean	SD	Median	LQ	UQ	% patients ≥2.1 & ≤2.37 mmol/L
1997	1633	2.3	0.3	2.3	2.2	2.5	40
1998	2060	2.3	0.3	2.3	2.2	2.5	44
1999	2732	2.3	0.3	2.3	2.2	2.5	39
2000	3703	2.4	0.3	2.3	2.2	2.5	42
2001	4618	2.4	0.2	2.4	2.2	2.5	40
2002	5485	2.3	0.3	2.3	2.2	2.5	43
2003	6466	2.3	0.2	2.3	2.2	2.4	46
2004	7536	2.3	0.2	2.3	2.2	2.4	47
2005	8630	2.3	0.2	2.3	2.2	2.4	49
2006	10717	2.3	0.2	2.3	2.1	2.4	50

Figure 9.2.1: Cumulative distribution of corrected Serum Calcium, HD patients 1997-2006

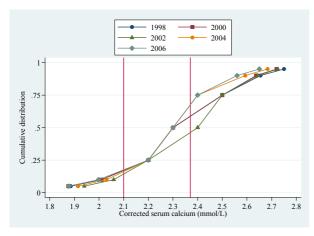


Figure 9.2.2: Cumulative distribution of corrected Serum Calcium, CAPD patients 1997-2006

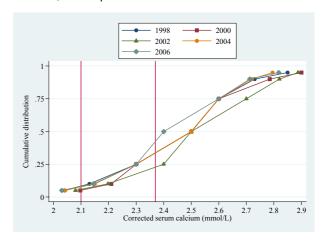


Table 9.2.2: Distribution of corrected Serum Calcium, CAPD patients 1997-2006

Year	No. of Subjects	Mean	SD	Median	LQ	UQ	% patients ≥2.1 & ≤2.37 mmol/L
1997	469	2.5	0.3	2.5	2.3	2.6	25
1998	535	2.4	0.3	2.4	2.3	2.6	30
1999	593	2.5	0.2	2.5	2.3	2.6	25
2000	635	2.5	0.2	2.5	2.3	2.6	25
2001	744	2.5	0.3	2.5	2.4	2.7	22
2002	859	2.5	0.2	2.5	2.3	2.6	24
2003	1167	2.4	0.2	2.5	2.3	2.6	27
2004	1276	2.5	0.2	2.5	2.3	2.6	23
2005	1338	2.4	0.2	2.4	2.3	2.6	30
2006	1494	2.4	0.2	2.4	2.3	2.5	38

The median serum phosphate levels were higher among HD patients (1.7 mmol/l) compared to CAPD patients (1.6 mmol/l). (tables and figs 9.2.3 & 9.2.4). In 2006, 46 % of patients on HD and 54% of CAPD patients had achieved the target serum phosphate of 1.13 to 1.78 mmol/l as advocated by K/DOQI guidelines.

Table 9.2.3: Distribution of Serum Phosphate, HD patients 1997-2006

Year	No of Subjects	Mean	SD	Median	LQ	UQ	% patients <1.13 mmol/L	% patients ≥1.13 & <1.78 mmol/L	% patients ≥1.78 & ≤2.6 mmol/L	% patients > 2.6 mmol/L
1997	1649	1.9	0.5	1.9	1.6	2.3	5	37	48	11
1998	2051	1.9	0.5	1.9	1.6	2.2	4	35	52	10
1999	2861	1.9	0.5	1.9	1.5	2.2	7	37	47	9
2000	4080	1.9	0.6	1.8	1.5	2.2	8	37	46	9
2001	4765	1.9	0.5	1.8	1.5	2.2	7	40	45	8
2002	5679	1.9	0.5	1.8	1.5	2.2	7	38	45	10
2003	6588	1.8	0.5	1.8	1.5	2.2	7	41	43	9
2004	7620	1.8	0.5	1.8	1.5	2.2	8	42	42	7
2005	8834	1.8	0.5	1.7	1.4	2.1	9	45	40	6
2006	10963	1.8	0.5	1.7	1.4	2.1	9	46	39	6

Figure 9.2.3: Cumulative distribution of Serum Phosphate, HD patients 1997-2006

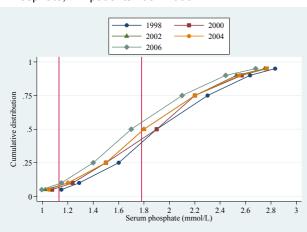


Figure 9.2.4: Cumulative distribution of Serum Phosphate, CAPD patients 1997-2006

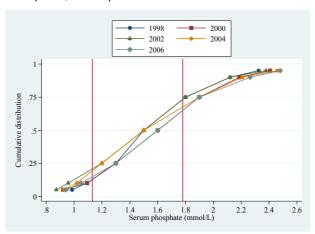


Table 9.2.4: Distribution of Serum Phosphate, CAPD patients 1997-2006

Year	No of Subjects	Mean	SD	Median	LQ	UQ	% patients <1.13 mmol/L	% patients ≥1.13 & <1.78 mmol/L	% patients ≥1.78 & ≤2.6 mmol/L	% patients > 2.6 mmol/L
1997	470	1.6	0.4	1.5	1.3	1.8	13	58	27	2
1998	537	1.6	0.5	1.6	1.3	1.9	12	55	30	3
1999	583	1.6	0.5	1.6	1.3	1.9	11	56	30	3
2000	633	1.5	0.5	1.5	1.3	1.8	17	55	26	2
2001	732	1.5	0.5	1.5	1.2	1.8	21	53	24	2
2002	862	1.5	0.5	1.5	1.2	1.8	21	52	25	2
2003	1173	1.6	0.5	1.5	1.2	1.9	16	53	28	3
2004	1278	1.6	0.5	1.6	1.3	1.9	15	52	29	3
2005	1343	1.6	0.5	1.6	1.3	1.9	15	52	29	3
2006	1510	1.6	0.5	1.6	1.3	1.9	13	54	29	4

The median corrected calcium phosphate product had declined from 3.9 mmol^2/L^2 in 2005 to 3.8 mmol^2/L^2 in HD patients and remained at 3.7 mmol^2/L^2 in CAPD patients. (tables and figs 9.2.5 & 9.2.6).

Table 9.2.5: Distribution of corrected calcium x phosphate product, HD patients 1997-2006

Year	No of Subjects	Mean	SD	Median	LQ	UQ	% patients <3.5 mmol ² /L ²	% patients ≥3.5 & <4 mmol²/L²	% patients ≥4 & <4.5 mmol²/L²	% patients ≥4.5 & <5 mmol²/L²	% patients ≥5 & <5.5 mmol ² /L ²	% patients ≥5.5 mmol²/L²
1997	1615	4.5	1.3	4.5	3.6	5.3	23	14	15	17	12	20
1998	2020	4.5	1.2	4.4	3.7	5.2	21	15	18	15	13	19
1999	2698	4.4	1.3	4.3	3.4	5.2	27	14	15	14	11	18
2000	3650	4.4	1.3	4.3	3.5	5.2	25	15	16	15	10	19
2001	4555	4.3	1.3	4.2	3.4	5.2	27	16	16	13	11	18
2002	5403	4.4	1.3	4.3	3.4	5.2	27	16	15	13	10	19
2003	6383	4.2	1.3	4.1	3.3	5.1	30	16	15	13	10	16
2004	7414	4.2	1.3	4.1	3.3	5	32	16	15	12	10	15
2005	8496	4	1.3	3.9	3.2	4.8	36	17	14	11	9	12
2006	10595	4	1.2	3.8	3.1	4.7	38	17	15	11	8	11

Figure 9.2.5: Cumulative distribution of corrected Calcium x Phosphate product, HD patients 1997-2006

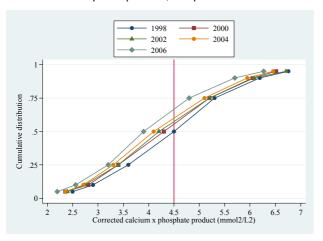


Figure 9.2.6: Cumulative distribution of corrected Calcium x Phosphate product, CAPD patients 1997-2006

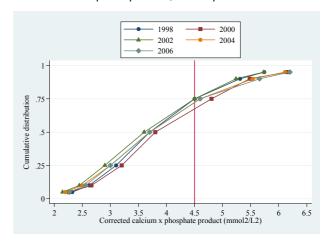


Table 9.2.6: Distribution of corrected calcium x phosphate product, CAPD patients 1997-2006

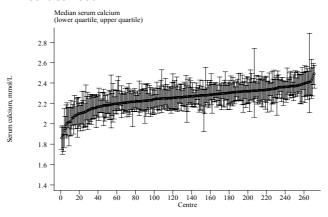
Year	No of Subjects	Mean	SD	Median	LQ	UQ	% patients <3.5 mmol ² /L ²	% patients ≥3.5 & <4 mmol²/L²	% patients ≥4 & <4.5 mmol²/L²	% patients ≥4.5 & <5 mmol ² /L ²	% patients ≥5 & <5.5 mmol²/L²	% patients ≥5.5 mmol²/L²
1997	468	3.9	1.1	3.7	3.1	4.5	40	20	15	10	6	7
1998	533	4	1.1	3.8	3.2	4.6	38	18	16	10	6	11
1999	580	4	1.2	3.8	3.2	4.8	36	20	13	12	9	10
2000	621	3.8	1.1	3.7	3.1	4.5	44	19	12	10	7	8
2001	723	3.8	1.1	3.6	2.9	4.5	46	18	12	10	8	7
2002	856	3.8	1.2	3.6	2.9	4.5	45	17	12	11	7	8
2003	1162	3.9	1.2	3.7	3	4.6	43	17	13	10	8	10
2004	1274	4	1.2	3.8	3	4.7	41	15	14	10	8	12
2005	1333	3.9	1.3	3.7	3	4.6	43	15	14	11	6	11
2006	1493	3.9	1.2	3.7	3.1	4.6	43	17	14	10	7	9

In 2006, the median corrected serum calcium level among HD patients from 271 centres ranged from 1.9 to 2.5 mmol/l. The median corrected serum calcium level among CAPD patients from 24 centres ranged from 2.2 to 2.6 mmol/l.(tables 9.2.7a & 9.2.8a).

Table 9.2.7: Variation in corrected serum calcium levels among HD centres, 2006 (a) Median serum calcium level among HD patients

` '			•	•				
Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	44	2.1	2.2	2.3	2.3	2.4	2.5	2.5
1998	49	2	2.1	2.3	2.3	2.4	2.5	2.5
1999	69	1.6	2	2.3	2.3	2.4	2.5	2.6
2000	93	2	2.1	2.3	2.3	2.4	2.6	3.2
2001	116	2	2.1	2.3	2.4	2.4	2.5	2.6
2002	139	1.9	2.1	2.2	2.3	2.4	2.5	2.6
2003	164	2	2.1	2.2	2.3	2.4	2.5	2.5
2004	198	1.9	2.1	2.2	2.3	2.4	2.4	2.5
2005	222	1.8	2	2.2	2.3	2.4	2.4	2.6
2006	271	1.9	2.1	2.2	2.3	2.3	2.4	2.5

Figure 9.2.7(a): Variation in median serum calcium level among HD patients, HD centres 2006



We reviewed the proportion of patients with serum calcium range 2.1 to 2.37 mmol/l. The median was 50% for HD centres (table 9.2.7b) and 47% for CAPD centres (table 9.2.8b) for year 2006. There is great variation between HD centres in proportion patients with serum calcium 2.1 to 2.37 mmol/l, ranging from 8% to 88%. proportion of patients within a centre with serum calcium 2.1 to 2.37 mmol/l among CAPD centres ranges from 0% to 76%.

Table 9.2.7(b) Proportion of patients with serum calcium 2.1 to 2.37 mmol/L, HD centres

	(-) -1						- ,	
Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	44	6	15	29.5	42	46.5	63	70
1998	49	12	21	36	45	53	70	76
1999	69	0	10	25	39	46	63	80
2000	93	0	13	32	42	48	65	96
2001	116	0	12	29	40	50	64	85
2002	139	5	17	33	44	54	68	80
2003	164	13	24	36.5	46.5	55	68	83
2004	198	7	20	38	47	58	72	82
2005	222	0	18	37	49.5	57	70	80
2006	271	8	30	42	50	60	71	88

Figure 9.2.7(b): Variation in proportion of patients with serum calcium 2.1 to 2.37 mmol/L, HD centres 2006

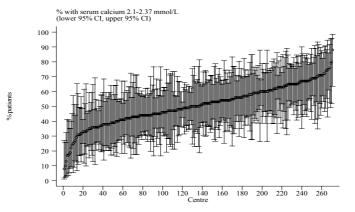
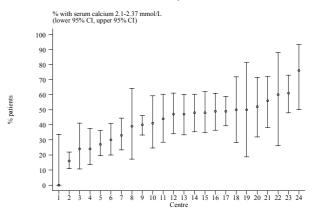


Table 9.2.8: Variation in corrected serum calcium levels among CAPD centres, 2006 (a) Median serum calcium level among CAPD patients

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	7	2.1	2.1	2.4	2.4	2.5	2.6	2.6
1998	9	2.2	2.2	2.3	2.4	2.4	2.6	2.6
1999	10	2.4	2.4	2.4	2.5	2.5	2.6	2.6
2000	11	2.4	2.4	2.4	2.5	2.5	2.6	2.6
2001	12	2.3	2.3	2.4	2.5	2.5	2.6	2.6
2002	15	2.4	2.4	2.4	2.4	2.5	2.6	2.6
2003	19	2.3	2.3	2.4	2.4	2.5	2.6	2.6
2004	19	2.3	2.3	2.4	2.4	2.5	2.5	2.5
2005	21	2.2	2.3	2.3	2.4	2.4	2.5	2.6
2006	24	2.2	2.2	2.3	2.4	2.4	2.5	2.6

Figure 9.2.8(a): Variation in median serum calcium level among CAPD patients, CAPD centres 2006

Figure 9.2.8(b): Variation in proportion of patients with serum calcium 2.1-2.37 mmol/L, CAPD centres 2006



(b) Proportion of patients with serum calcium 2.1 - 2.37 mmol/L, CAPD centres

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	7	10	10	18	26	29	53	53
1998	9	11	11	27	38	40	64	64
1999	10	5	5	21	28	31	42	42
2000	11	13	13	18	25	33	48	48
2001	12	11	11	18	23.5	36	38	38
2002	15	12	12	20	25	33	46	46
2003	19	10	10	19	35	40	64	64
2004	19	10	10	18	26	31	50	50
2005	21	16	18	24	33	40	45	51
2006	24	0	16	31.5	47	50	61	76

With regards to the proportion of patients with serum phosphate level 1.13-1.78 mmol/L, the CAPD centres have a higher median proportion of patients achieving this target (51.5%) compared with HD centres (46%) (tables 9.2.9a & 9.2.9b).

Table 9.2.9: Variation in serum phosphate levels among HD centres, 2006 (a) Median serum phosphate level among HD patients

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	44	1.3	1.5	1.7	1.9	2.1	2.3	2.8
1998	49	1.5	1.5	1.8	1.9	2.1	2.2	2.6
1999	70	1.1	1.6	1.8	1.9	2	2.1	2.1
2000	100	1.4	1.6	1.7	1.8	2	2.2	3.7
2001	116	1.3	1.5	1.7	1.8	1.9	2.1	2.3
2002	146	1.3	1.5	1.8	1.9	2	2.2	2.4
2003	172	0.9	1.5	1.7	1.8	1.9	2.2	2.4
2004	197	1.3	1.5	1.7	1.8	1.9	2.1	2.2
2005	227	8.0	1.4	1.6	1.7	1.9	2.1	2.3
2006	274	0.9	1.4	1.6	1.7	1.8	2.1	2.5

Figure 9.2.9 (a): Variation in median serum phosphate level among HD patients, HD centres 2006

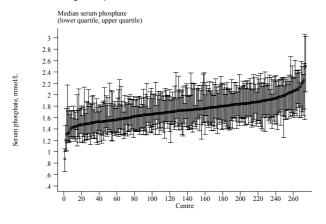
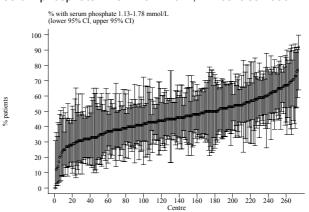


Figure 9.2.9(b): Variation in proportion of patients with serum phosphate 1.13-1.78 mmol/L, HD centres 2006



(b) Proportion of patients with serum phosphate 1.13-1.78 mmol/L, HD centres 2006

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	44	7	18	27.5	37	44.5	59	68
1998	49	9	17	29	34	44	63	70
1999	70	8	17	27	35	44	57	63
2000	100	9	18.5	29.5	37	44	57.5	73
2001	116	0	21	33	38	47	61	70
2002	146	5	14	29	36	46	61	91
2003	172	8	19	31	40	48	64	93
2004	197	0	19	31	41	53	68	92
2005	227	10	24	35	44	53	67	77
2006	274	0	27	38	46	54	67	92

Table 9.2.10: Variation in serum phosphate levels among CAPD centres, 2006 (a) Median serum phosphate level among CAPD patients

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	7	1.4	1.4	1.5	1.5	1.6	1.7	1.7
1998	9	1.5	1.5	1.5	1.6	1.6	1.8	1.8
1999	9	1.5	1.5	1.6	1.6	1.6	1.7	1.7
2000	11	1.3	1.3	1.4	1.5	1.6	1.7	1.7
2001	12	1.3	1.3	1.4	1.5	1.7	1.9	1.9
2002	15	1.4	1.4	1.4	1.5	1.6	2.1	2.1
2003	19	1.1	1.1	1.5	1.5	1.6	1.7	1.7
2004	19	1.4	1.4	1.5	1.5	1.7	1.8	1.8
2005	21	1.4	1.4	1.5	1.6	1.7	1.8	1.9
2006	24	1.3	1.4	1.5	1.6	1.7	1.9	2.1

Figure 9.2.10(a): Variation in median serum phosphate level among CAPD patients, CAPD centres 2006

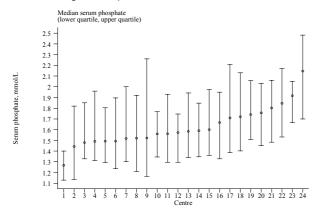
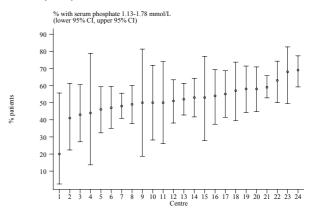


Figure 9.2.10(b): Variation in proportion of patients with serum phosphate 1.13-1.78 mmol/L, CAPD centres 2006



(b) Proportion of patients with serum phosphate 1.13-1.78 mmol/L, CAPD centres 2006

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	7	53	53	54	58	66	76	76
1998	9	43	43	54	60	61	82	82
1999	9	45	45	51	58	65	68	68
2000	11	46	46	48	53	61	65	65
2001	12	40	40	48.5	54	58	79	79
2002	15	43	43	46	54	60	82	82
2003	19	36	36	47	54	60	81	81
2004	19	34	34	49	52	64	76	76
2005	21	40	43	46	52	58	64	76
2006	24	20	41	47.5	51.5	57.5	68	69

A higher number of CAPD centres have median calcium phosphate product of less than 4.5 mmol $^2/L^2$ as compared to HD centres (71.5% versus 70%). There is an increasing trend among HD centres achieving a calcium phosphate product of less than 4.5 mmol $^2/L^2$ and the difference between CAPD centres and HD centres is decreasing. (tables and figs 9.2.11 & 9.2.12)

Table 9.2.11: Variation in corrected calcium x phosphate product among HD centres, 2006 (a) Median corrected calcium x phosphate product among HD patients

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	44	2.9	3.7	4.1	4.4	4.8	5.3	6.2
1998	49	3.2	3.3	4.1	4.5	4.7	5.3	5.3
1999	68	2.3	3.2	4.1	4.4	4.7	5.2	5.3
2000	91	3.1	3.7	4	4.3	4.6	5.1	6.2
2001	112	2.9	3.6	3.9	4.3	4.6	5	5.7
2002	139	2.9	3.6	3.9	4.2	4.6	5.2	6.2
2003	164	2.1	3.4	3.8	4.1	4.5	4.9	5.7
2004	195	2.9	3.3	3.8	4.1	4.4	5	5.5
2005	217	2.1	3.2	3.6	3.9	4.3	4.7	5.6
2006	268	1.8	3.2	3.5	3.9	4.2	4.7	5.2

Figure 9.2.11(a): Variation in median corrected calcium x phosphate product among HD patients, HD centres 2006

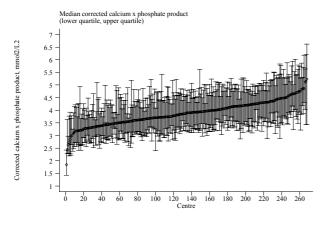
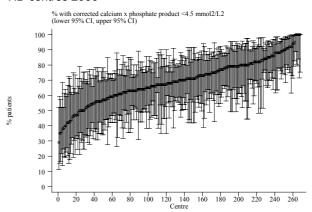


Figure 9.2.11(b): Variation in proportion of patients with corrected calcium x phosphate product $< 4.5 \text{ mmol}^2/L^2$, HD centres 2006



(b) Proportion of patients with corrected calcium x phosphate product < 4.5 mmol²/L²

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	44	15	26	38.5	51.5	66	77	100
1998	49	20	30	40	52	65	83	91
1999	68	20	30	47	55	64	90	95
2000	91	12	33	47	57	67	80	88
2001	112	18	38	47	57	70.5	82	91
2002	139	14	33	48	57	69	89	100
2003	164	25	33	51.5	61.5	73	88	100
2004	195	17	36	53	64	74	91	100
2005	217	23	43	58	69	77	90	100
2006	268	29	43	61	70	80	91	100

Table 9.2.12: Variation in corrected calcium x phosphate product among CAPD centres, 2006 (a) Median corrected calcium x phosphate product among CAPD patients

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	7	3.5	3.5	3.6	3.7	3.8	3.9	3.9
1998	9	3.4	3.4	3.6	3.7	3.9	4	4
1999	9	3.6	3.6	3.7	3.9	4.1	4.2	4.2
2000	11	3.4	3.4	3.5	3.7	4	4.3	4.3
2001	12	3.1	3.1	3.4	3.7	3.8	4.3	4.3
2002	15	3.3	3.3	3.4	3.7	4	4.9	4.9
2003	19	2.7	2.7	3.4	3.7	4	4.1	4.1
2004	19	3.2	3.2	3.5	3.8	4	4.4	4.4
2005	21	3.3	3.4	3.6	3.7	4.1	4.3	4.4
2006	24	3	3.3	3.5	3.7	4	4.4	4.8

Figure 9.2.12(a): Variation in median corrected calcium x phosphate product among CAPD patients, CAPD centres 2006

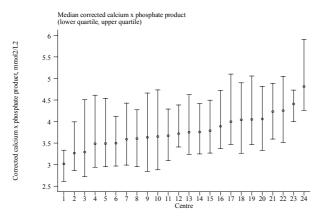


Figure 9.2.12(b): Variation in proportion of patients with corrected calcium x phosphate product $< 4.5 \text{ mmol}^2/L^2$, CAPD centres 2006

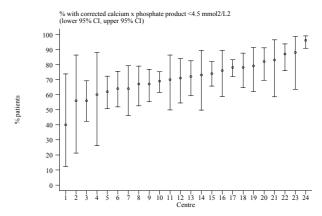


Table 9.2.12(b) Proportion of patients with corrected calcium x phosphate product $< 4.5 \text{ mmol}^2/L^2$

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	7	70	70	74	78	82	94	94
1998	9	66	66	71	73	79	91	91
1999	9	59	59	65	72	74	80	80
2000	11	65	65	70	73	81	85	85
2001	12	50	50	71.5	75	80.5	84	84
2002	15	43	43	65	78	83	88	88
2003	19	61	61	64	75	81	100	100
2004	19	57	57	67	72	79	91	91
2005	21	54	56	63	74	77	85	85
2006	24	40	56	64	71.5	78.5	88	96

SECTION 9.3: SERUM iPTH

Among patients on HD, the mean iPTH was 220.3 ng/ml while the median was 94.5 ng/ml. Among patients on CAPD, the mean iPTH was 224.57 ng/ml while the median iPTH was 188.9 ng/ml.(table 9.3.1, 9.3.2, 9.3.3, 9.3.4). The majority of patients (61% of HD patients, 54% of CAPD patients) have iPTH level < 150 ng/ml.

Tables and figures 9.3.1 and 9.3.2 (b) and (c) show that diabetics have consistently lower iPTH values than non diabetics.

Table 9.3.1(a): Distribution of iPTH, HD patients 1997 - 2006

Year	No. of Subjects	Mean	SD	Median	LQ	UQ	% patients <150 ng/ml	% patients ≥150 & ≤300 ng/ml	% patients >300 & ≤500 ng/ml	% patients >500 ng/ml
1997	1088	195.1	282.9	76.8	26	240.3	66	13	9	11
1998	938	126.1	202	44	15	141	76	12	6	6
1999	1533	185.6	260.7	78.9	23.5	240	64	16	10	10
2000	2244	149.3	230	58	17.6	178.3	72	13	8	7
2001	2760	141.2	219.5	57	18	164.8	73	15	6	7
2002	3391	161.6	248	64	19	191	70	14	8	8
2003	4068	219.1	328.8	79	24.3	263.3	64	14	9	14
2004	4748	212.1	325.6	74.3	22.6	257.3	65	13	9	13
2005	5826	221.6	312.5	83.8	26.5	297	61	14	11	14
2006	7645	220.3	307.8	89	29.3	293.6	61	14	11	14

Figure 9.3.1(a): Cumulative distribution of iPTH, HD patients 1997-2006

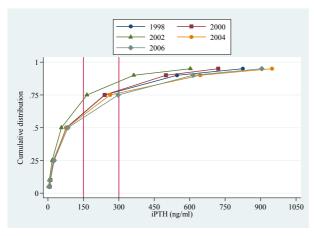


Figure 9.3.1(b): Cumulative distribution of iPTH, diabetic HD patients 1997-2006

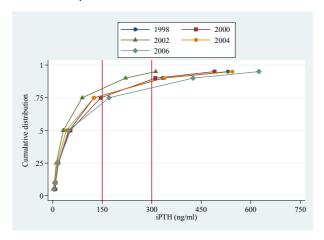


Table 9.3.1(b) Distribution of iPTH, diabetic HD patients 1997 - 2006

Year	No. of Subjects	Mean	SD	Median	LQ	UQ	% patients <150 ng/ml	% patients ≥150 & ≤300 ng/ml	% patients >300 & ≤500 ng/ml	% patients >500 ng/ml
1997	200	127.8	216.6	49.5	17.5	125	78	11	7	5
1998	185	82.6	137.2	24	11	83	84	9	5	2
1999	336	121.5	181.8	53.5	16	145.8	75	14	6	5
2000	530	87.5	137.1	35.8	11	101	83	9	6	2
2001	720	82.5	139.6	32	10.9	89.5	83	11	3	2
2002	966	92.6	161.6	35	11	99	83	10	4	3
2003	1244	122.5	211.2	40.6	13.4	124.8	78	10	6	6
2004	1574	113.7	196.6	38	14	118	80	10	5	5
2005	2151	151.1	248.6	47.6	16.4	171	72	12	8	8
2006	3059	155.2	252.1	55	20.8	174.5	72	12	8	7

Table 9.3.1(c) Distribution of iPTH, non diabetic HD patients 1997 - 2006

Year	No. of Subjects	Mean	SD	Median	LQ	UQ	% patients <150 ng/ml	% patients ≥150 & ≤300 ng/ml	% patients >300 & ≤500 ng/ml	% patients >500 ng/ml
1997	888	210.2	293.8	83.8	28.3	272.5	63	14	10	13
1998	753	136.8	213.7	50	17	154	74	12	7	7
1999	1197	203.6	276.3	93.2	26.5	267.2	61	17	11	11
2000	1714	168.4	248.8	65.6	21.8	204	69	14	9	9
2001	2040	162	238.1	71	23.5	198	69	16	7	8
2002	2425	189.1	270.2	85	26	236	65	15	10	10
2003	2824	261.6	360.8	108	33.5	331	57	16	10	17
2004	3174	260.9	363.6	102.5	31	340.3	58	14	12	17
2005	3675	262.9	337.7	114.6	36	364.5	55	15	13	17
2006	4586	263.8	333	125.3	40	366	54	16	13	18

Figure 9.3.1(c): Cumulative distribution of iPTH, non diabetic HD patients 1997-2006

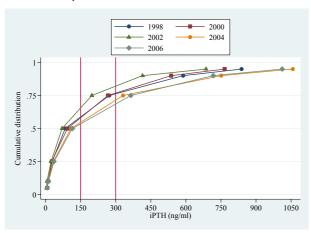


Figure 9.3.2(a): Cumulative distribution of iPTH, CAPD patients 1997-2006

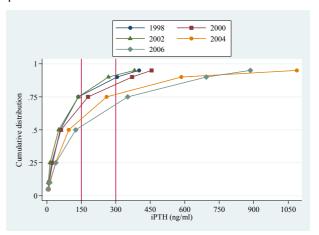


Table 9.3.2(a): Distribution of iPTH, CAPD patients 1997 – 2006

Year	No. of Subjects	Mean	SD	Median	LQ	UQ	% patients <150 ng/ml	% patients ≥150 & ≤300 ng/ ml	% patients >300 & ≤500 ng/ml	% patients >500 ng/ml
1997	293	112.3	151	58	25	137	78	12	7	3
1998	280	93.7	117.4	47.5	18.5	126	81	13	5	1
1999	365	132.8	176.4	61.5	21	179.3	71	15	10	4
2000	406	109.8	192.4	46.8	15.5	118	80	12	5	4
2001	531	108	155.8	51.5	13.5	137.6	76	15	6	3
2002	681	160.6	219.1	82	26	196	67	17	8	7
2003	938	230.3	340.3	95	37.4	260	61	18	9	12
2004	1115	216.4	302.9	105	39.5	260	60	19	10	11
2005	1071	247.1	306.4	125.3	39	352	54	18	13	15
2006	1264	224.7	271.9	128	41.5	318.5	54	20	14	12

Table 9.3.2(b) Distribution of iPTH, diabetic CAPD patients 1997 - 2006

Year	No. of Subjects	Mean	SD	Median	LQ	UQ	% patients <150 ng/ml	% patients ≥150 &≤300 ng/ ml	% patients >300 &≤500 ng/ ml	% patients >500 ng/ml
1997	90	61.3	66.1	39	15	90.5	91	7	2	0
1998	84	59.2	68.4	34.3	10.3	88.5	90	7	2	0
1999	100	95.8	145.2	41	17	111.6	81	11	5	3
2000	114	66.2	174.5	27.7	6	69	89	9	2	1
2001	165	65.7	87.6	33.5	7.5	82.5	87	10	2	1
2002	205	101.1	155.5	60	16	132	80	14	3	2
2003	326	122.9	175	68	29	154.3	74	16	6	4
2004	380	131	191.4	65.5	24.4	148.3	75	15	4	6
2005	367	159.4	235.6	69.2	23.9	190.5	70	16	7	7
2006	462	151.7	198	91.5	33	187.8	67	19	8	5

Figure 9.3.2(b): Cumulative distribution of iPTH, diabetic CAPD patients 1997-2006

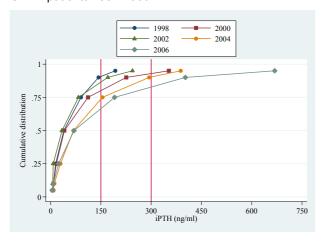


Figure 9.3.2(c): Cumulative distribution of iPTH, non diabetic CAPD patients 1997-2006

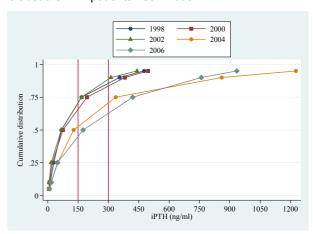


Table 9.3.2(c) Distribution of iPTH, non diabetic CAPD patients 1997 - 2006

Year	No. of Sub- jects	Mean	SD	Median	LQ	UQ	% patients <150 ng/ml	% patients ≥150 &≤300 ng/ml	% patients >300 &≤500 ng/ml	% patients >500 ng/ml
1997	203	134.9	171.3	68	29.5	167	72	14	9	4
1998	196	108.5	130.3	57.5	22.3	139.3	77	16	6	2
1999	265	146.8	185.2	75	22.5	194	67	16	12	5
2000	292	126.7	196.6	57.3	22.7	139	76	13	6	5
2001	366	127.1	175	67	16.7	168	72	17	7	4
2002	476	186.2	237	98.5	32.3	242	62	19	10	10
2003	612	287.6	389.6	128	50	336.5	54	18	10	17
2004	735	260.6	338.6	140	50	329	53	21	12	14
2005	704	292.8	328.4	175	48.2	420	46	19	16	19
2006	802	266.8	298.7	167.3	50	390	47	21	16	16

A higher number of CAPD centres had median iPTH 150-300 ng/ml as compared to HD centres (21.5% versus 14%). The proportion of patients with iPTH 150-300 ng/ml among HD centres ranged from 0%-33% while the proportion of patients with iPTH 150-300 ng/ml among CAPD centres ranged from 5% to 31% (Fig 9.3.3, 9.3.4)

Table 9.3.3: Variation in iPTH among HD centres, 2006 (a) Median iPTH among HD patients

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	36	8	26	45.5	71.8	112.4	252.5	402.5
1998	30	8	13	24	45.5	105	196.6	221
1999	39	10	17	40.3	80.3	151	304.5	380
2000	59	5.6	15	31.5	48	86.7	352.5	481.8
2001	70	7.3	12.3	27.6	55	101.8	240	550
2002	93	2.9	13	28.4	54	145.5	319.3	466
2003	113	4	10.8	37	92.4	188.5	375.2	624.5
2004	135	3.6	13	29.8	76	223.5	412	854
2005	167	5.8	13.6	38	96	215.3	396.8	612.3
2006	213	9.3	16.8	40	94.5	206.2	410.5	704.5

Figure 9.3.3(a): Variation in iPTH among HD patients, HD centres 2006

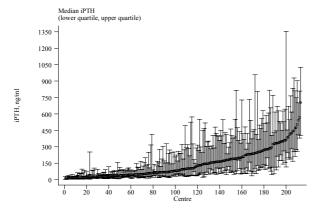


Figure 9.3.3(b): Variation in proportion of patients with iPTH 150-300 ng/ml, HD centres 2006

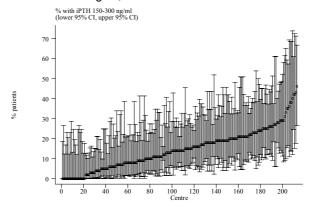


Table 9.3.3(b) Proportion of patients with iPTH 150-300 ng/ml

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	36	0	0	8	11.5	17.5	29	29
1998	30	0	0	5	9.5	16	26	29
1999	39	0	0	9	16	23	36	38
2000	59	0	0	4	10	15	33	42
2001	70	0	0	6	10	21	35	40
2002	93	0	0	2	10	21	31	43
2003	113	0	0	7	14	21	38	45
2004	135	0	0	5	11	20	38	50
2005	167	0	0	6	13	20	33	50
2006	213	0	0	7	14	21	33	46

Table 9.3.4: Variation in iPTH among CAPD centres, 2006 (a)Median iPTH among CAPD patients

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	5	36.5	36.5	44.8	47	81	120	120
1998	5	16	16	57.5	66.3	68	73	73
1999	8	16.5	16.5	52.8	74.5	87.5	263.9	263.9
2000	9	16	16	33	43.3	60.8	122	122
2001	11	15.4	15.4	42.5	56.3	91	274	274
2002	14	27.3	27.3	50	81.1	107	319	319
2003	17	22.3	22.3	70	131	170.5	393	393
2004	18	41	41	76.3	138.8	169.3	329.1	329.1
2005	19	25	25	83.8	181	321.5	496.9	496.9
2006	22	34.5	38.3	96	188.9	230	382	494

Figure 9.3.4(a): Variation in median iPTH among CAPD patients, CAPD centres 2006

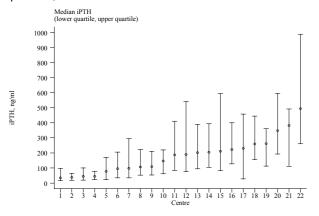


Figure 9.3.4(b): Variation in proportion of patients with iPTH 150-300 ng/ml, CAPD centres 2006

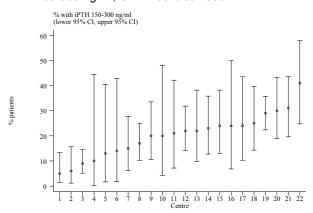


Table 9.3.4(b) Proportion of patients with iPTH 150-300 ng/ml

	•	•		•				
Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	5	7	7	10	13	15	27	27
1998	5	0	0	13	15	17	24	24
1999	8	6	6	7	14	20.5	26	26
2000	9	0	0	5	12	17	18	18
2001	11	0	0	9	14	17	30	30
2002	14	0	0	10	15.5	21	24	24
2003	17	2	2	16	19	23	33	33
2004	18	7	7	15	21	26	29	29
2005	19	0	0	9	16	23	31	31
2006	22	5	6	14	21.5	24	31	41

Conclusion

There is no major change in the types of phosphate binders used. Calcium carbonate still remains the major phosphate binder for both HD and CAPD patients in year 2006. The number of patients on aluminium hydroxide is on a decreasing trend. The number of patients on lanthanum and sevelamer is still very small and is not included in this report.

The use of vitamin D has increased. The number of patients on paricalcitol is also very small and is not included in this report. The number of patients who underwent parathyroidectomy has increased by three fold compared to 2005. This may reflect active participation of our endocrine surgeon and is also due to the extension of parathyroidectomy service in other hospitals outside Kuala Lumpur Hospital and Putrajaya Hospital.

Mean corrected serum calcium remains unchanged for both HD (2.3mmol/l) and CAPD (2.4mmol/l) patients since 2002. The availability of low calcium dialysate in hemodialysis helps to keep the mean calcium level lower in HD patients. More CAPD patients achieved target serum calcium (2.1-2.37 mmol/l) compared to year 2005 (from 30 to 38%). Phosphate control continues to be better in CAPD patients. Median corrected serum calcium phosphate product for HD patients shows a decreasing trend from 1997 to 2005 but remained static at 3.9 mmol²/l² in 2006. Median corrected serum calcium phosphate product for CAPD patients also remains static but is still lower compared to HD (3.7 mmol²/l² vs 3.9 mmol²/l²). Furthermore, more CAPD patients achieved corrected serum calcium phosphate product of less than 4.5 mmol²/l². Differences in dialysis management have resulted in wide variation of outcome results in serum calcium, phosphate and calcium phosphate product.

Mean serum iPTH level remained static in HD patients (220.3ng/ml); however it has decreased from 247.1ng/ml to 224.7ng/ml in CAPD patients between 2005 and 2006. More HD patients have serum iPTH level of <150ng/ml compared to CAPD patients (61% vs 54%). The percentage of patients achieving the target iPTH level of 150-300ng/ml is low. Only 20% of CAPD patients have serum iPTH between 150-300ng/ml while only 14% in the HD population achieved this target. Again a wide variation of serum iPTH was observed due to differences in dialysis management among dialysis centres. As expected, diabetics had lower iPTH values.

Management of renal bone disease seems to have improved among HD patients over last few years but is still not as good as CAPD patients. Unfortunately, the relationship of these factors in relation to cardiovascular disease outcome has not been determined as yet but hopefully will be done in the future.

CHAPTER 10

Hepatitis on Dialysis

Teo Sue Mei Claire Tan Hui Hong Foo Sui Mei

Table 10.1: Prevalence of positive HBsAg and positive Anti-HCV at annual survey, HD patients 1997-2006

Table 10.2: Prevalence of positive HBsAg and positive Anti-HCV at annual survey, CAPD patients 1997-2006

					J, - 1	
No. of subjects	Prevalence of HBsAg+ (%)	Prevalence of Anti-HCV+ (%)	Year	No. of subjects	Prevalence of HBsAg+ (%)	Prevalence of Anti- HCV+ (%)
1694	6	23	1997	476	3	5
2139	6	22	1998	541	3	6
2991	6	23	1999	610	2	5
4386	6	25	2000	662	2	5
5187	6	23	2001	781	2	3
6106	5	20	2002	891	3	4
6977	5	19	2003	1223	3	4
7618	5	17	2004	1200	4	5
8957	4	14	2005	1318	4	5
11116	5	12	2006	1489	5	4
	subjects 1694 2139 2991 4386 5187 6106 6977 7618 8957	No. of subjects of HBsAg+ (%) 1694 6 2139 6 2991 6 4386 6 5187 6 6106 5 6977 5 7618 5 8957 4	No. of subjects of HBsAg+ (%) Prevalence of Anti-HCV+ (%) 1694 6 23 2139 6 22 2991 6 23 4386 6 25 5187 6 23 6106 5 20 6977 5 19 7618 5 17 8957 4 14	No. of subjects of HBsAg+ (%) Prevalence of Anti-HCV+ (%) Year 1694 6 23 1997 2139 6 22 1998 2991 6 23 1999 4386 6 25 2000 5187 6 23 2001 6106 5 20 2002 6977 5 19 2003 7618 5 17 2004 8957 4 14 2005	No. of subjects Prevalence of HBsAg+ (%) Prevalence of Anti-HCV+ (%) Year No. of subjects 1694 6 23 1997 476 2139 6 22 1998 541 2991 6 23 1999 610 4386 6 25 2000 662 5187 6 23 2001 781 6106 5 20 2002 891 6977 5 19 2003 1223 7618 5 17 2004 1200 8957 4 14 2005 1318	No. of subjects Prevalence of HBsAg+ (%) Prevalence of Anti-HCV+ (%) Year No. of subjects Prevalence of HBsAg+ (%) 1694 6 23 1997 476 3 2139 6 22 1998 541 3 2991 6 23 1999 610 2 4386 6 25 2000 662 2 5187 6 23 2001 781 2 6106 5 20 2002 891 3 6977 5 19 2003 1223 3 7618 5 17 2004 1200 4 8957 4 14 2005 1318 4

The prevalence of Hepatitis B infection has remained low and was similar when comparing HD and CAPD patients whereas prevalence of HCV was still higher in HD than CAPD patients as a result of the higher risk of nosocomial transmission with HD. However with effective and more stringent implementation of infection control measures, HCV prevalence has continued to decline by 2-3% annually from 2001 onwards.

There was only small center variation in the proportion of hepatitis B positive patients in both HD and CAPD.

Table 10.3: Variation in Proportion of patients with positive HBsAg at annual survey among HD centres, 2006

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	44	0	0	2.5	5	9	17	19
1998	50	0	0	0	5	9	18	23
1999	76	0	0	0	4	9.5	19	30
2000	109	0	0	0	4	8	15	80
2001	125	0	0	0	5	9	15	90
2002	156	0	0	0	3	8	14	27
2003	178	0	0	0	3.5	8	15	64
2004	198	0	0	0	3	8	15	100
2005	231	0	0	0	1	6	15	100
2006	283	0	0	0	0	6	16	94

Figure 10.3: Variation in Proportion of patients with positive HBsAg among HD centres, 2006

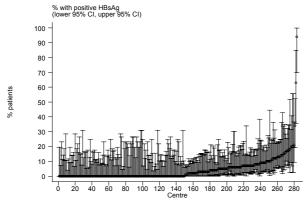


Table 10.4: Variation in Proportion of	natients with positive H	RsAg at annual survey am	ong CAPD centres 2006
rable lu.t. valiation in laportion of	patients with positive in	Dong at ailiual sulvey all	ong OAI D Contics, 2000

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	7	0	0	0	2	3	8	8
1998	9	0	0	0	1	3	6	6
1999	10	0	0	0	2	2	4	4
2000	11	0	0	0	1	4	5	5
2001	12	0	0	0	2	3	9	9
2002	15	0	0	1	3	6	18	18
2003	19	0	0	1	3	6	8	8
2004	19	0	0	1	3	6	11	11
2005	21	0	0	0	3	5	5	11
2006	23	0	0	0	4	6	9	13

Figure 10.4: Variation in Proportion of patients with positive HBsAg among CAPD centres, 2006

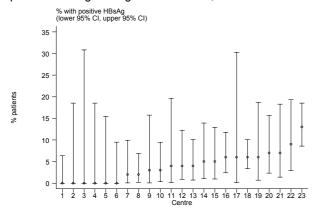
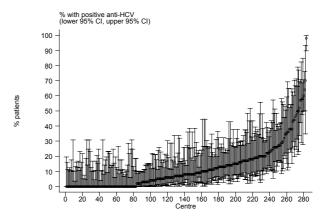


Figure 10.5: Variation in Proportion of patients with positive anti-HCV among HD centres, 2006



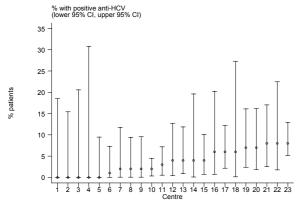
Even though the median proportion of HCV infected HD patients decline continued to annually from 21% in 1997 to 8% in 2006, there was still wide center variation (table and figure 10.5). This probably reflected the differences infection control practices and criteria for intake of patients among centers.

Table 10.5: Variation in Proportion of patients with positive anti-HCV at annual survey among HD centres, 2006

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	44	0	0	14.5	21	29	47	63
1998	50	0	0	9	19.5	30	61	79
1999	76	0	0	7	19.5	32.5	60	79
2000	109	0	0	9	18	30	70	91
2001	125	0	0	7	17	30	64	91
2002	156	0	0	5	14	24.5	51	96
2003	178	0	0	5	13	24	50	96
2004	201	0	0	4	11	24	50	100
2005	232	0	0	0	9.5	19	44	98
2006	283	0	0	0	8	17	44	98

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	7	0	0	0	6	7	9	9
1998	9	0	0	3	3	8	11	11
1999	10	0	0	3	4	7	14	14
2000	11	0	0	2	3	8	10	10
2001	12	0	0	0	3	4.5	7	7
2002	15	0	0	0	3	8	12	12
2003	19	0	0	1	4	7	9	9
2004	19	0	0	0	4	7	10	10
2005	21	0	0	1	4	8	10	11
2006	23	0	0	1	4	6	8	8

Figure 10.6: Variation in Proportion of patients with positive anti-HCV among CAPD centres, 2006



Similar to Hepatitis B infection, the prevalence of HCV infection was low in CAPD patients and did not vary greatly between centers.

Table 10.7a: Risk factors in relation to HD practices for seroconversion to anti-HCV positive among sero-negative patients

Risk factor	Number of patients	Risk Ratio	95% (CI	p-value
Assistance to Perform HD					
(1) Self care ^{ref}	151	1.000			
(2) Partial self care	108	0.701	(0.542	0.908)	0.007
(3) Completely assisted	239	0.469	(0.377	0.583)	0.000
Dialyzer Reuse					
(1) less than 10 ^{ref}	161	1.000			
(2) more than 10	362	0.733	(0.607	0.885)	0.001
Dialyzer					
Reprocessing System					
(1) Fully Auto ^{ref}	275	1.000			
(2) Semi Auto	29	0.702	(0.478	1.030)	0.070
(3) Manual	29	0.710	(0.484	1.042)	0.081
(4) No Reuse	0	0.000	•		
Age					
(1) <=20 ^{ref}	29	1.000			
(2) 21-40	189	0.914	(0.619	1.432)	0.778
(3) 41-60	242	0.538	(0.356	0.813)	0.003
(4) >60	63	0.243	(0.154	0.385)	0.000
Gender					
(1)Female ^{ref}	213	1.000			
(2)Male	310	1.168	(0.977	1.395)	0.088
Diabetes					
(1)No ^{ref}	420	1.000			
(2)Yes	103	0.330	(0.267	0.408)	0.000
Previous			`	,	
Renal Transplant					
(1)No ^{ref}	474	1.000			
(2)Yes	49	3.357	(2.426	4.645)	0.000
History of					
Blood Transfusion					
(1)No ^{ref}	319	1.000			
(2)Yes	204	1.287	(1.075	1.540)	0.006

Risk factors associated with occurrence of HCV seroconversion can be divided into patient-level factors and center-level factors. Patient-level factors such as previous renal transplant and history of blood transfusion were associated with a significantly higher risk of HCV seroconversion. There was also a trend of increasing risk with men and younger patients and lower risk with diabetes.

Completely assisted HD patients had a significantly lower risk of acquiring HCV infection. This was not surprising as these patients were fully assisted by trained HD staffs who were more stringent with infection control measures. However a rather interesting finding was that center-level factors such as increased frequency of dialyser reuse, the use of semi-automated and manual reprocessing systems seemed to be associated with lower risk of acquiring HCV infection. As the number of study patients was small, it is possible that these factors did not actually influence HCV seroconversion, but rather reflected differences in practices among centers. Total segregation and good practice of universal precautions are probably more important contributing factors.

Table 10.7b: Risk factors for seroconversion to anti-HCV positive among sero-negative patients in CAPD

Risk factor	Number of patients	Risk Ratio	95% CI	p-value
Age				
(1) <=20 ^{ref}	4			
(2) 21-40	8	1.553	(0.474 5.095)	0.467
(3) 41-60	8	0.713	(0.213 2.387)	0.583
(4) >60	2	0.338	(0.062 1.859)	0.213
Gender				
(1)Female ^{ref}	9			
(2)Male	13	1.384	(0.604 3.171)	0.323
Diabetes				
(1)No ^{ref}	18			
(2)Yes	4	0.334	(0.113 0.986)	0.047
Switched from HD to CAPD				
(1)No ^{ref}	18			
(2)Yes	4	2.215	(0.744 6.591)	0.153
Previous Renal Transplant				
(1)No ^{ref}	22			
(2)Yes	1	1.144	(0.152 8.610)	0.896
History of Blood Transfusion				
(1)No ^{ref}	11			
(2)Yes	12	2.191	(0.961 4.994)	0.062

CAPD patients who were switched from HD, had previous renal transplant and blood transfusion had a tendency for increased risk of seroconversion, but these findings did not reach statistical significance as the number of study patients was small.

Conclusion:

Nosocomial transmission in HD has been implicated for the higher HCV prevalence in HD compared to CAPD. Our efforts to curb the epidemic of hepatitis in HD has been shown to be effective with the decreasing prevalence of HCV annually. However there should be further investigations into the mechanism of transmission and aspects of our current HD practices which may put patients at risk of HCV infection.

CHAPTER 11

Haemodialysis Practices

Tan Chwee Choon Shahnaz Shah Firdaus Khan Rafidah Abdullah Norleen Bt Zulkarnain Sim

SECTION 11.1: VASCULAR ACCESS AND ITS COMPLICATIONS

There was a progressive decline in the percentage of patients having native vascular access from 98% in 1997 to 93% in 2006. The ratio of brachiocephalic fistula (BCF) to arteriovascular fistula (AVF) has increased. In 2006, 25% of native vascular access was BCF. The proportion of patients with artificial graft and permanent catheter remained at 2% respectively. (Table 11.1.1)

Table 11.1.1: Vascular Access on Haemodialysis, 1997-2006

Access types	199	7	199	98	199	99	200	00	200	1
	No.	%	No.	%	No.	%	No.	%	No.	%
Wrist AVF	1427	85	1763	84	2406	81	3561	82	4049	79
BCF*	213	13	273	13	431	14	655	15	897	17
Venous graft	4	0	6	0	8	0	11	0	19	0
Artificial graft	13	1	20	1	34	1	31	1	64	1
Permanent CVC	4	0	8	0	17	1	19	0	25	0
Temporary CVC*	20	1	37	2	77	3	77	2	90	2
Temporary FVC*	0	0	0	0	0	0	0	0	0	0
TOTAL	1681	100	2107	100	2973	100	4354	100	5144	100
Access types	200)2	200	03	200)4	200)5	200	6
	No.	%	No.	%	No.	%	No.	%	No.	%
Wrist AVF	4680	78	5249	75	5891	73	6405	69	7704	68
BCF*	1068	18	1359	19	1693	21	2169	23	2793	25
Venous graft	14	0	23	0	41	1	30	0	22	0
Artificial graft	78	1	114	2	150	2	221	2	275	2
Permanent CVC	43	1	62	1	99	1	180	2	229	2
Temporary CVC*	138	2	180	3	233	3	269	3	291	3
Temporary FVC*	0	0	0	0	0	0	7	0	21	0
TOTAL	6021	100	6987	100	8107	100	9281	100	11335	100

^{*} BCF=Brachiocephalic fistula

Table 11.1.2: Difficulties reported with Vascular Access, 1997-2006

Access difficulty	199	97	199	98	199	9	200	00	200)1
	No.	%	No.	%	No.	%	No.	%	No.	%
Difficulty with needle placement	55	47	82	4	133	5	146	4	217	5
Difficulty in obtaining desired blood flow rate	48	41	60	3	112	5	136	4	239	5
Other difficulties	12	10	30	2	55	2	32	1	39	1
No difficulties	1	1	1778	91	2155	88	3402	92	4276	90
TOTAL	116	100	1950	100	2455	100	3716	100	4771	100
Access difficulty	200)2	200)3	200)4	200	5	200	6
	No.	%	No.	%	No.	%	No.	%	No.	%
Difficulty with needle placement	215	4	217	3	255	3	319	4	391	3
Difficulty in obtaining desired blood flow rate	235	4	243	4	301	4	354	4	354	3
Other difficulties	57	1	60	1	67	1	58	1	45	0
No difficulties	5073	91	5970	92	6957	92	8339	92	10416	93
TOTAL	5580	100	6490	100	7580	100	9070	100	11206	100

4000

^{*} CVC= Central venous catheter

^{*} FVC= Femoral venous catheter

Complication rates have remained similar despite an increase in intake of elderly and diabetic patients on dialysis in recent years. (Table 11.1.3)

Table 11.1.3: Complications reported with Vascular Access, 1997-2006

Complication	199	97	199	98	199	9	200	00	200)1
Complication	No.	%								
Thrombosis	71	19	69	3	129	5	148	4	209	4
Bleed	23	6	37	2	23	1	30	1	62	1
Aneurysmal dilatation	121	33	134	6	159	6	208	5	212	4
Swollen limb	35	9	36	2	51	2	44	1	67	1
Access related infection, local/systemic	29	8	21	1	34	1	52	1	49	1
Distal limb ischaemia	4	1	12	1	9	0	26	1	22	0
Venous outflow obstruction	45	12	50	2	71	3	78	2	123	2
Carpal tunnel	23	6	19	1	35	1	42	1	41	1
Others	18	5	48	2	64	2	37	1	74	1
No complications	0	0	1636	79	2119	79	3237	83	4204	83
TOTAL	369	100	2062	100	2694	100	3902	100	5063	100
O and Parties	200)2	200)3	200)4	200)5	200	16
Complication	No.	%								
Thrombosis	202	3	220	3	284	4	289	3	313	3
Bleed	66	1	54	1	67	1	73	1	68	1
Aneurysmal dilatation	211	4	199	3	193	2	179	2	244	2
Swollen limb	56	1	55	1	77	1	84	1	88	1
Access related infection, local/systemic	52	1	43	1	70	1	63	1	76	1
Distal limb ischaemia	17	0	13	0	37	0	35	0	30	0
Venous outflow obstruction	101	2	119	2	151	2	170	2	197	2
Carpal tunnel	44	1	63	1	49	1	55	1	48	0
Others	118	2	118	2	133	2	109	1	113	1
No complications	4988	85	5963	87	6896	87	8113	88	9987	89

SECTION 11.2: HD PRESCRIPTION

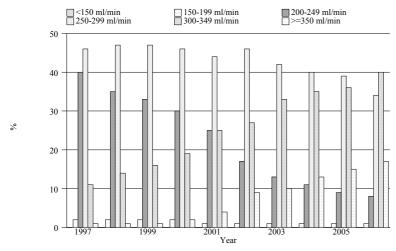
There was an increasing trend towards the use of higher blood flow rates from 1997 to 2006. The proportion of patients with blood flow of 300-349mls/min has increased from 11% to 40% and those with blood flow rates \geq 350mls/min from 1% to 17%. Fifty seven percent of patients had blood flow rates of \geq 300mls/min in 2006. (Table 11.2.1 and Fig 11.2.1)

Table 11.2.1: Blood Flow Rates in HD centres, 1997-2006

Table 11.2.1. Blood	I Flow Rat	65 111 110	Centres, 1	991 – 200	U					
Blood flow rates	199	97	199	98	199	9	200	00	200	1
Blood flow rates	No.	%	No.	%	No.	%	No.	%	No.	%
<150 ml/min	2	0	4	0	6	0	9	0	7	0
150-199 ml/min	34	2	36	2	65	2	85	2	69	1
200-249 ml/min	649	40	735	35	962	33	1282	30	1233	25
250-299 ml/min	734	46	968	47	1367	47	1938	46	2229	44
300-349 ml/min	176	11	298	14	455	16	814	19	1276	25
>=350 ml/min	18	1	30	1	31	1	94	2	216	4
TOTAL	1613	100	2071	100	2886	100	4222	100	5030	100
Blood flow rates	200)2	200)3	200	14	200	5	200	6
blood flow rates	No.	%	No.	%	No.	%	No.	%	No.	%
<150 ml/min	9	0	4	0	11	0	7	0	5	0

Blood flow rates	200	2	200)3	200	4	200)5	200	6
Blood flow fales	No.	%	No.	%	No.	%	No.	%	No.	%
<150 ml/min	9	0	4	0	11	0	7	0	5	0
150-199 ml/min	69	1	84	1	86	1	94	1	100	1
200-249 ml/min	973	17	882	13	879	11	814	9	904	8
250-299 ml/min	2692	46	2865	42	3112	40	3523	39	3738	34
300-349 ml/min	1590	27	2241	33	2711	35	3226	36	4471	40
>=350 ml/min	505	9	690	10	1020	13	1328	15	1900	17
TOTAL	5838	100	6766	100	7819	100	8992	100	11118	100

Figure 11.2.1: Blood Flow Rates in HD centres, 1997–2006



Ninety seven percent of patients were on 3 haemodialysis (HD) sessions / week. The small percentage of patients on 2 HD sessions / week is likely to be patients who were dialyzing in private centres and unable to afford 3 HD sessions / week. (Table 11.2.2)

Table 11.2.2: Number of HD Sessions per week, 1997 - 2006

HD sessions	199	7	199	98	199	9	200	00	200	1
per week	No.	%	No.	%	No.	%	No.	%	No.	%
1	1	0	1	0	4	0	8	0	8	0
2	6	0	5	0	153	5	341	8	337	7
3	1664	99	2110	100	2811	95	3982	92	4761	92
4	9	1	2	0	3	0	10	0	50	1
TOTAL	1680	100	2118	100	2971	100	4341	100	5156	100
HD sessions	200)2	200)3	200)4	200	5	200	6
per week	No.	%	No.	%	No.	%	No.	%	No.	%
1	10	0	15	0	11	0	7	0	25	0
2	369	6	343	5	281	3	265	3	267	2
3	5603	93	6558	95	7709	96	9010	97	11148	97
4	18	0	9	0	30	0	31	0	33	0

100

8031

100

9313

100

11473

100

Majority of patients (99%) were on 4 hours HD session. (Table 11.2.3)

6925

100

Table 11.2.3: Duration of HD, 1997 – 2006

TOTAL

6000

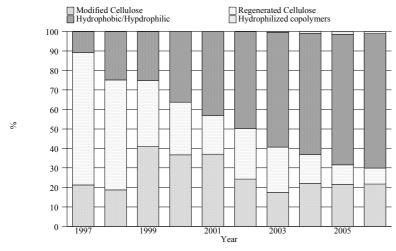
Duration of	199	7	199	98	199	9	200	0	200	1
HD per session	No.	%	No.	%	No.	%	No.	%	No.	%
<=3 hours	7	0	3	0	4	0	8	0	6	0
-3.5 hours	3	0	18	1	9	0	12	0	33	1
-4 hours	1594	95	1993	94	2735	92	4053	93	4956	96
-4.5 hours	69	4	91	4	160	5	189	4	106	2
-5 hours	8	0	8	0	61	2	77	2	59	1
>5 hours	1	0	3	0	0	0	13	0	0	0
TOTAL	1682	100	2116	100	2969	100	4352	100	5160	100
Duration of HD	200)2	200)3	200)4	200	5	200	6
Duration of HD per session	200 No.)2 %	200 No.)3 %	200 No.)4 %	200 No.	5 %	200 No.	6 %
per session	No.	%	No.	%	No.	%	No.	%	No.	%
per session <=3 hours	No. 18	0	No.	0	No. 23	% 0	No.	% 0	No. 28	0
per session <=3 hours -3.5 hours	No. 18 15	% 0 0	No. 11 7	% 0 0	No. 23 17	% 0 0	No. 22 17	% 0 0	No. 28 7	% 0 0
per session <=3 hours -3.5 hours -4 hours	No. 18 15 5845	% 0 0 98	No. 11 7 6761	% 0 0 98	No. 23 17 7830	% 0 0 97	No. 22 17 9152	% 0 0 98	No. 28 7 11321	% 0 0 99
per session <=3 hours -3.5 hours -4 hours -4.5 hours	No. 18 15 5845 68	% 0 0 98 1	No. 11 7 6761 76	% 0 0 98 1	No. 23 17 7830 119	% 0 0 97 1	No. 22 17 9152 67	% 0 0 98 1	No. 28 7 11321 68	% 0 0 99 1

The use of synthetic membrane (hydrophobic/ hydrophilic and hydrophilised copolymer) has increased from 11% in 1997 to 70% in 2006. Regenerated cellulose membrane usage has progressively declined from 67% in 1997 to 8% in 2006. The use of modified cellulose membrane has remained relatively the same. (Table 11.2.4 and Fig. 11.2.4)

Table 11.2.4: Dialyser membrane types in HD centres, 1997 – 2006

Dialyser	19	97	19	98	19	99	20	00	20	01
membrane	No.	%	No.	%	No.	%	No.	%	No.	%
Modified Cellulose	360	21	395	19	1224	41	1605	37	1666	37
Regenerated Cellulose	1148	68	1195	56	1012	34	1183	27	890	20
Hydrophobic/Hypdrophilic	184	11	524	25	754	25	1589	36	1944	43
Hydrophilized copolymers	1	0	2	0	1	0	0	0	0	0
TOTAL	1693	100	2116	100	2991	100	4377	100	4500	100
Dialyser	2002		2003		2004		2005		2006	
membrane	No.	%	No.	%	No.	%	No.	%	No.	%
Modified Cellulose	1376	24	1115	17	1718	22	1919	21	2333	22
Regenerated Cellulose	1473	26	1502	23	1150	15	901	10	882	8
Hydrophobic/Hypdrophilic	2828	50	3782	59	4843	62	5976	67	7432	69
Hydrophilized copolymers	1	0	35	1	74	1	132	1	121	1
TOTAL	5678	100	6434	100	7785	100	8928	100	1076 8	100

Figure 11.2.4: Dialyser membrane types in HD centres, 1997 – 2006



Reuse if dialysers is a common practice in Malaysia whereby 95% reuse the dialyser. The frequency of reuse depends on the type of dialyser membrane. Five percent of patients did not reuse dialysers. In 2006, 75% of patients reuse their dialysers 10 times or more. (Table 11.2.5)

Table 11.2.5: Dialyser Reuse Frequency in HD centres, 1997-2006

Dialyser reuse	199	97	199	98	199	9	200	00	200	1
frequency	No.	%	No.	%	No.	%	No.	%	No.	%
1*	21	1	16	1	65	2	116	3	152	3
2	9	1	5	0	13	0	17	0	15	0
3	996	63	215	11	191	7	205	5	232	5
4	174	11	113	6	250	9	477	12	416	9
5	194	12	137	7	264	10	312	8	357	7
6	154	10	1072	55	1414	51	1730	43	1413	29
7	2	0	37	2	46	2	69	2	85	2
8	4	0	66	3	122	4	357	9	793	16
9	30	2	109	6	179	6	101	2	132	3
10	0	0	84	4	96	3	246	6	400	8
11	0	0	23	1	6	0	4	0	43	1
12	0	0	64	3	118	4	333	8	470	10
>=13	0	0	0	0	0	0	91	2	331	7
TOTAL	1584	100	1941	100	2764	100	4058	100	4839	100
Dialyser reuse	200)2	200)3	200)4	200)5	200	16
frequency	No.	%	No.	%	No.	%	No.	%	No.	%
1*	197	4	251	4	319	4	196	4	392	5
2	41	1	19	0	42	1	1	0	5	0
3	316	6	349	5	194	3	81	2	36	0
4	337	6	339	5	192	3	85	2	75	1
5	318	6	267	4	192	3	137	3	187	3
				•		U				
6	1216	22	915	14	806	11	555	10	589	8
6 7	1216 124							10 1	589 63	8 1
		22	915	14	806	11	555			
7	124	22 2	915 71	14 1	806 89	11 1	555 44	1	63	1
7 8	124 866	22 2 16	915 71 852	14 1 13	806 89 809	11 1 11	555 44 477	1 9	63 410	1 6
7 8 9	124 866 59	22 2 16 1	915 71 852 87	14 1 13 1	806 89 809 50	11 1 11 1	555 44 477 46	1 9 1	63 410 115	1 6 2
7 8 9 10	124 866 59 538	22 2 16 1	915 71 852 87 880	14 1 13 1	806 89 809 50 1160	11 1 11 1 1	555 44 477 46 770	1 9 1 15	63 410 115 911	1 6 2 13
7 8 9 10 11	124 866 59 538 36	22 2 16 1 10 1	915 71 852 87 880 25	14 1 13 1 14 0	806 89 809 50 1160 42	11 1 11 1 16 1	555 44 477 46 770 12	1 9 1 15 0	63 410 115 911 100	1 6 2 13 1

Ninety nine percent of patients were on bicarbonate dialysate buffer in 2006 compared to 67% in 1997. In 2006, 147 patients were still using acetate as a buffer. (Table 11.2.6)

Table 11.2.6: Dialysate Buffer used in HD centres,	1997 – 2006
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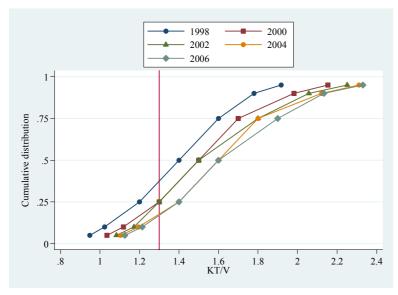
Dialysate buffer	1997		1998		1999		2000		2001	
	No.	%	No.	%	No.	%	No.	%	No.	%
Acetate	551	33	627	30	552	19	393	9	240	5
Bicarbonate	1123	67	1492	70	2429	81	3969	91	4920	95
TOTAL	1674	100	2119	100	2981	100	4362	100	5160	100
Dialysate buffer	2002		2003		2004		2005		2006	
	No.	%	No.	%	No.	%	No.	%	No.	%
Acetate	138	2	76	1	33	0	58	1	147	1
Bicarbonate	5880	98	6815	99	7957	100	9268	99	11457	99
TOTAL	6018	100	6891	100	7990	100	9326	100	11604	100

The median prescribed KT/V was 1.6. The percentage of patients with KT/V \geq 1.3 has increased from 60% in 1997 to 84% in 2006. Since 2002, the median KT/V has remained the same. (Table 11.2.7a and Fig. 11.2.7a)

Table 11.2.7a: Distribution of prescribed KT/V, HD patients 1997-2006

Year	No. of subjects	Mean	SD	Median	LQ	UQ	% patients ≤1.3
1997	1558	1.4	0.3	1.4	1.2	1.6	59
1998	2022	1.4	0.3	1.4	1.2	1.6	67
1999	2831	1.5	0.3	1.5	1.3	1.7	73
2000	4087	1.5	0.4	1.5	1.3	1.8	75
2001	4908	1.6	0.4	1.5	1.3	1.8	78
2002	5496	1.6	0.4	1.6	1.4	1.8	81
2003	6516	1.6	0.4	1.6	1.4	1.8	82
2004	7453	1.6	0.4	1.6	1.4	1.8	82
2005	8749	1.6	0.4	1.6	1.4	1.9	83
2006	10919	1.7	0.4	1.6	1.4	1.9	84

Figure 11.2.7a: Cumulative distribution of prescribed KT/V, HD patients 1997-2006



In 2006, 142 (37.4%) centres returned data on delivered KT/V. Overall delivered KT/V data were reported in 37.9% of HD patients. The methods of measuring KT/V were urea kinetic modeling (64.1%), online modeling (23.5%) and others (12.4%).

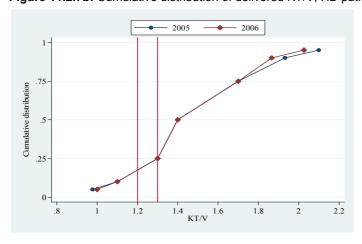
Table 11.2.7b and Fig. 11.2.7b shows that although the prescribed median KT/V was 1.6, the delivered median KT/V was only 1.4. In 2006, the percentage of patients with a delivered KT/V \geq 1.2 and KT/V \geq 1.3 was 82% and 66% respectively.

Table 11.2.7b: Distribution of delivered KT/V, HD patients 2005-2006

Year	No. of subjects	Mean	SD	Median	LQ	UQ	% patients ≤1.2	% patients ≤1.3	Variance*
2005	1760	1.6	2.7	1.4	1.3	1.7	82	65	0
2006	5553	1.5	1.3	1.4	1.3	1.7	82	66	0

^{* (}prescribed KT/V - delivered KT/V)/ prescribed KT/V

Figure 11.2.7b: Cumulative distribution of delivered KT/V, HD patients 2005-2006



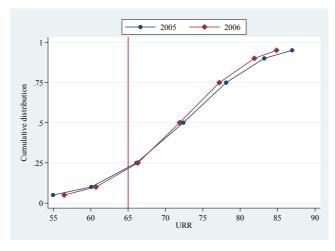
Data returns for Urea Reduction Ratio (URR) were better than that of delivered KT/V. Overall 201 centres (52.9%) representing 55.1% of HD population submitted data on URR. Post dialysis blood sampling methodologies were not requested for and hence not specified.

In 2006, the median URR was 71.9% and the percentage of patients with URR \geq 65% was 79%. (Table 11.2.7c and Fig. 11.2.7c)

Table 11.2.7c Distribution of URR, HD patients 2005 – 2006

Year	No. of subjects	Mean	SD	Median	LQ	UQ	% patients ≤65
2005	2514	71.8	10.3	72.4	66.1	78.1	79
2006	8065	71.4	9.2	71.9	66.3	77.2	79

Figure 11.2.7c Cumulative distribution of URR, HD patients 2005 - 2006



The median blood flow rates among centres had increased from 250 mls/min in 1997 to 300mls/min in 2006. There is still a wide variation in practices among centres. The median blood flow rates among centres ranges from 220mls/min to 400mls/min. (Table 11.2.8(a) and Fig. 11.2.8 (a))

Table 11.2.8: Variation in HD prescription among HD centres 2006

Table 11.2.8(a) Median	blood	flow rates	in HD	patients
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Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	45	200	200	220	250	250	280	300
1998	46	200	200	230	250	250	300	300
1999	67	200	200	230	250	250	300	300
2000	100	200	200	240	250	275	300	300
2001	116	200	220	250	252.5	300	300	350
2002	137	200	230	250	280	300	300	350
2003	155	200	240	250	280	300	325	350
2004	184	220	250	257.5	287.5	300	350	400
2005	228	200	250	260	300	300	350	400
2006	279	210	250	270	300	300	350	400

Figure 11.2.8(a): Variation in median blood flow rates in HD patients among HD centres 2006

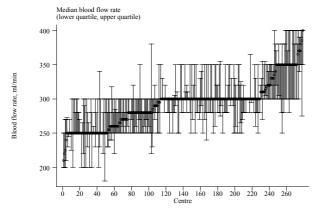
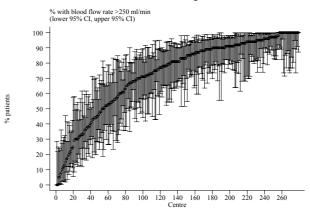


Figure 11.2.8(b): Variation in Proportion of patients with blood flow rates > 250 ml/min among HD centres 2006



There was an increase in the proportion of patients with blood flow rates from > 250mls/min. In 2006, 50% of centres had 81% of their patients with blood flow rates of > 250mls/min compared to only 13% in 1997. (Table 11.2.8(b))

There was still a wide variation in the proportion of patients with blood flow rate > 250mls/min among centres. Two centres that had no patients with blood flow rates of > 250mls/min. (Fig. 11.2.8 (b))

Table 11.2.8(b) Proportion of patients with blood flow rates > 250 ml/min

	· / ·							
Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	45	0	0	4	13	27	60	64
1998	46	0	2	9	20.5	38	79	100
1999	67	0	2	8	28	49	85	100
2000	100	0	0	10.5	31.5	59.5	85.5	91
2001	116	0	0	22.5	49.5	73.5	92	100
2002	137	0	2	36	61	82	95	100
2003	155	0	4	42	70	85	98	100
2004	184	0	17	50	73	86	96	100
2005	228	0	17	54.5	77	90.5	99	100
2006	279	0	19	57	81	92	100	100

The majority of centres had 100% of their patients with 3 HD sessions/ week. There are some centres with significant proportion of their patients with their patients with less than 3 HD session/ week. (Table 11.2.8(c) and Fig. 11.2.8 (c))

Table 12.2.8(c) Proportion of patients with 3 HD sessions per week

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	47	80	92	99	100	100	100	100
1998	46	80	98	100	100	100	100	100
1999	69	17	45	97	100	100	100	100
2000	100	25	44.5	90.5	100	100	100	100
2001	118	23	50	92	100	100	100	100
2002	137	28	48	94	99	100	100	100
2003	160	36	55	97	100	100	100	100
2004	188	37	70	98	100	100	100	100
2005	231	40	75	99	100	100	100	100
2006	283	52	83	98	100	100	100	100

Figure 11.2.8(c): Variation in proportion of patients with 3 HD sessions per week among HD centres 2006

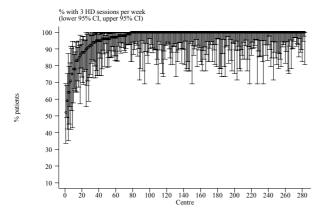
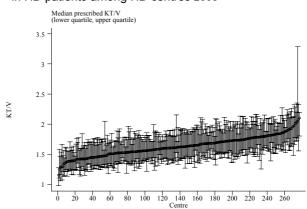


Figure 11.2.8(d): Variation in median prescribed KT/V in HD patients among HD centres 2006



The median prescribed KT/V in HD patients was 1.6 in 2006. The minimum prescribed KT/V was 1.2 and maximum prescribed KT/V was 2.1. (Table 11.2.8(d)) The variation of prescribed KT/V among centres (fig. 11.2.8 (d)) was less marked than the variation in proportion of patients with blood flow rates of > 250 mls/min. (fig. 11.2.8(b)).

Table 11.2.8(d): Median prescribed KT/V in HD patients

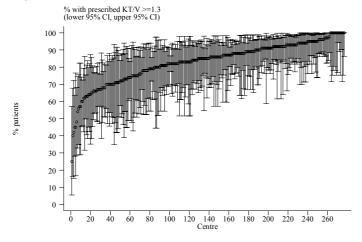
Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	44	1.2	1.2	1.3	1.4	1.4	1.5	1.8
1998	45	1.1	1.3	1.4	1.4	1.5	1.6	1.7
1999	67	1.2	1.3	1.4	1.5	1.6	1.7	1.8
2000	99	1.2	1.3	1.4	1.5	1.6	1.8	2.8
2001	114	1.2	1.3	1.5	1.5	1.7	1.8	1.9
2002	132	1.2	1.4	1.5	1.6	1.7	1.8	2.1
2003	150	1.2	1.4	1.5	1.6	1.7	1.9	2.1
2004	181	1.2	1.4	1.5	1.6	1.7	1.9	2.2
2005	224	1.3	1.4	1.5	1.6	1.7	1.9	2
2006	277	1.2	1.4	1.5	1.6	1.7	1.9	2.1

In 2006, half the centres had 85% of their patients with a prescribed KT/V \geq 1.3. However there was variation in proportion of patients with prescribed KT/V \geq 1.3 among the HD centres ranging from below 30% in one centre to 100% in a small number of centres. (Fig. 11.2.8 (e))

Table 11.2.8(e): Proportion of patients with prescribed KT/V \geq 1.3	Table 11.2.8	:): Proportion	of patients with	prescribed KT/V ≥ 1.3
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Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
1997	44	32	44	51.5	59.5	70	90	100
1998	45	0	46	60	67	74	85	96
1999	67	36	50	67	73	83	94	100
2000	99	26	47	66	79	86	94	100
2001	114	42	52	71	81.5	89	96	100
2002	132	35	58	74.5	82	90	97	100
2003	150	30	57	77	84	91	96	100
2004	181	28	61	74	83	91	100	100
2005	224	47	63	75	85	92	100	100
2006	277	25	62	76	85	92	100	100

Figure 11.2.8(e): Variation in proportion of patients with prescribed KT/V ≥ 1.3 among HD centres 2006

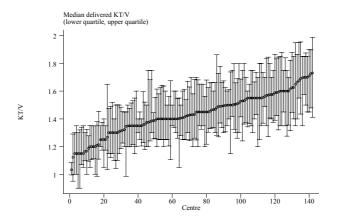


The median delivered KT/V in HD patients was 1.4 in 2006. The minimum delivered KT/V was 1.0 and maximum delivered KT/V was 1.7. There was marked variation of median delivered KT/V among HD centres. (Table 11.2.8 f and fig 11.2.8 f)

Table 11.2.8(f): Median delivered KT/V in HD patients

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
2005	52	1.2	1.2	1.3	1.4	1.5	1.7	1.7
2006	142	1	1.2	1.3	1.4	1.5	1.7	1.7

Figure 11.2.8(f): Variation in median delivered KT/V in HD patients among HD centres 2006



In 2006, 50% of centres had 82.5% of their patients with a delivered KT/V \geq 1.2. There was however a wide variation in the proportion of patients with a delivered KT/V \geq 1.2 among centres. Eleven centres had < 50% of their patients with a delivered KT/V \geq 1.2. Two centres had 100% of their patients with delivered KT/V \geq 1.2. (Table 11.2.8 g and Fig. 11.8.2 g)

The median URR in HD centres was 71.6% in 2006. The minimum URR was 55.4% and maximum URR was 94.4%. (Table 11.2.8 h) There was variation of median URR among HD centres but the variation was less wide than the variation of delivered KT/V among HD centres as shown by the less steep curve. (fig 11.2.8 h and fig 11.2.8 h)

In 2006, 50% of centres had 80% of their patients with URR \geq 65%. There was however a wide variation in the proportion of patients with URR \geq 65% among centres. Six centres had < 50% of their patients with a URR \geq 65%. One centre had no patients with URR \geq 65%. Four centres had 100% of their patients with a URR \geq 65%. (Table 11.2.8 j and Fig. 11.8.2 j)

Table 11.2.8(g) Proportion of patients with delivered KT/V ≥1.2

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
2005	52	40	53	74	84	90.5	100	100
2006	142	21	47	75	82.5	90	96	100

Figure 11.2.8(g): Variation in proportion of patients with delivered KT/V \geq 1.2 among HD centres 2006

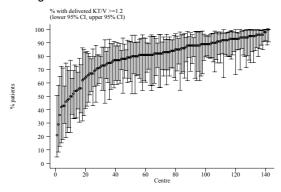


Table 11.2.8(h) Median URR among HD patients

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
2005	73	61.3	66.6	69.8	71.9	74.4	85.9	96.2
2006	210	55.4	64.8	69.1	71.6	74.4	78.2	94.4

Figure 11.2.8(h): Variation in median URR among HD patients, HD centres 2006

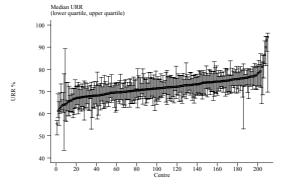
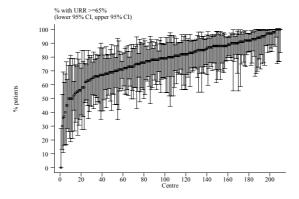


Table 11.2.8 (i) Proportion of HD patients with URR ≥ 65%

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
2005	73	40	53	70	81	88	100	100
2006	210	0	50	70	80	88	97	100

Figure 11.2.8(i): Variation in proportion of patients with URR \geq 65% among HD centres 2006



SECTION 11.3: TECHNIQUE SURVIVAL ON DIALYSIS

The unadjusted HD technique survival at 1 year, 5 years and 10 years was 89%, 56% and 33% respectively. The CAPD unadjusted technique survival was 81% at 1 year, 29% at 5 years and 7% at 10 years. (Table 11.3.2 and Fig. 11.3.1)

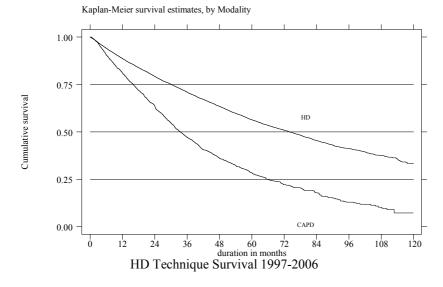
Table 11.3.1: Unadjusted technique survival by Dialysis modality, 1997-2006

Dialysis modality		CAPD			HD			All Dialysis	
Interval (months)	No.	% Survival	SE	No.	% Survival	SE	No.	% Survival	SE
6	2537	91	1	17560	94	0	20097	94	0
12	2087	81	1	14950	89	0	17037	88	0
24	1376	64	1	10955	79	0	12331	77	0
36	847	47	1	7895	71	0	8742	68	0
48	497	36	1	5592	63	0	6089	60	0
60	293	29	1	3809	56	0	4102	53	0
72	160	22	1	2536	51	1	2695	47	0
84	88	18	1	1578	45	1	1666	42	1
96	44	13	1	884	41	1	927	37	1
108	19	10	1	372	38	1	390	34	1
120	3	7	2	27	33	1	29	30	1

^{*} No. = Number at risk

SE=standard error

Figure 11.3.1: Unadjusted technique survival by Dialysis modality, 1997-2006



There was no apparent difference in the unadjusted HD technique survival by year of starting dialysis for the years 1997 to 2006. (Table 11.3.2 and fig 11.3.2)

Table 11.3.2: Unadjusted technique survival by year of entry, 1997-2006

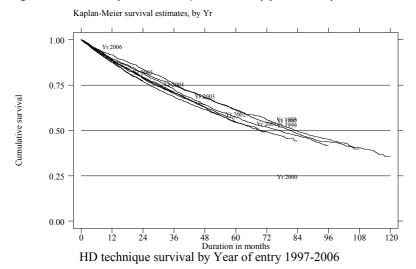
Year		1997			1998			1999	
Interval (months)	No.	% Survival	SE	No.	% Survival	SE	No.	% Survival	SE
6	946	93	1	1101	95	1	1321	95	1
12	891	89	1	1051	92	1	1236	90	1
24	810	82	1	942	84	1	1097	82	1
36	735	75	1	839	76	1	961	73	1
48	660	68	1	744	68	1	839	64	1
60	587	62	2	664	61	1	742	57	1
72	512	55	2	603	56	1	670	52	1
84	450	48	2	528	49	2	602	47	-
96	406	44	2	479	45	2	-	-	-
108	372	40	2	-	-	-	-	-	-
120	27	36	2	-	-	-	-	-	-

Year	2000			2001				2002		2003		
Interval (months)	No.	% Survival	SE									
6	1602	94	1	1766	93	1	2009	94	1	2157	94	0
12	1481	89	1	1620	87	1	1883	89	1	1998	88	1
24	1277	79	1	1404	77	1	1616	79	1	1764	79	1
36	1124	71	1	1235	69	1	1435	70	1	1565	71	1
48	982	63	1	1098	62	1	1274	63	1	-	-	-
60	853	55	1	966	55	1	-	-	-	-	-	-
72	756	49	1	-	-	-	_	-	-	-	-	-

Year		2004			2005			2006			
Interval (months)	No.	% Survival	SE	No.	% Survival	SE	No.	% Survival	SE		
6	2534	94	0	2648	93	0	1482	94	1		
12	2341	88	1	2453	88	1	-	-	-		
24	2050	79	1	-	-	-	-	-	-		

^{*} No. = Number at risk

Figure 11.3.2: Unadjusted technique survival by year of entry, 1997-2006



SE=standard error

The unadjusted HD technique survival was better in the younger age groups than the older age group, ten years unadjusted HD technique survival in the age group of 25-34, 35-44, 44-54, 55-64 and \geq 65 years old was 71%, 46%, 36%, 20% and 13% respectively. (Table 11.3.3 and fig 11.3.3)

Table 11.3.3: Unadjusted technique survival by age, 1997-2006

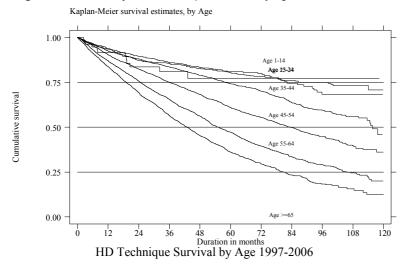
Age group (years)		<=14			15-24			25-34			35-44	
Interval (months)	No.	% Survival	SE	No.	% Survival	SE	No.	% Survival	SE	No.	% Survival	SE
6	66	95	0	678	96	1	1412	96	1	2478	96	0
12	55	92	0	584	94	1	1242	94	1	2169	92	1
24	41	84	1	430	87	1	961	89	1	1727	88	1
36	28	81	1	334	85	1	770	86	1	1359	84	1
48	19	77	1	251	83	2	610	83	1	1051	79	1
60	13	77	1	190	81	2	444	80	1	803	74	1
72	11	77	1	141	80	2	347	78	1	575	70	1
84	9	77	1	97	75	3	235	76	2	367	64	1
96	5	77	1	56	68	3	144	75	2	226	59	2
108	3	77	1	31	68	3	65	73	2	98	56	2
120	-	-	-	4	68	3	4	71	3	4	46	4

Age group (years)		45-54			55-64			>=65	
Interval (months)	No.	% Survival	SE	No.	% Survival	SE	No.	% Survival	SE
6	4468	95	0	4888	93	0	3571	91	0
12	3817	90	0	4154	87	0	2931	84	1
24	2881	82	1	2958	76	1	1961	69	1
36	2111	75	1	2068	66	1	1231	56	1
48	1540	68	1	1393	56	1	730	45	1
60	1053	61	1	889	47	1	423	36	1
72	710	55	1	535	39	1	226	30	1
84	442	50	1	322	34	1	113	23	1
96	234	45	1	169	29	1	56	18	1
108	93	40	2	69	24	1	19	16	2
120	9	36	2	6	20	2	3	13	2

^{*} No. = Number at risk

SE=standard error

Figure 11.3.3: Unadjusted technique survival by age, 1997-2006



Unadjusted HD technique survival in non diabetics at 1 year, 5 years and 10years was 91%, 69% and 47% respectively. Unadjusted HD technique survival for diabetics was worse than non diabetics with at 86%, 43% and 15% at one, five and ten years respectively. (Table 11.3.4 and Fig 11.3.4)

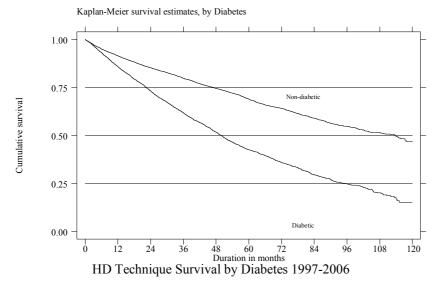
Table 11.3.4: Unadjusted technique survival by Diabetes status, 1997-2006

Diabetes status		Non-Diabetic			Diabetic	
Interval (months)	No.	% Survival	SE	No.	% Survival	SE
6	8790	95	0	8770	93	0
12	7690	91	0	7260	86	0
24	5980	85	0	4975	73	0
36	4637	80	0	3258	62	1
48	3519	75	1	2073	52	1
60	2569	69	1	1240	43	1
72	1807	64	1	731	36	1
84	1181	59	1	398	30	1
96	680	55	1	205	25	1
108	311	51	1	62	20	1
120	23	47	2	5	15	2

^{*} No. = Number at risk

SE=standard error

Figure 11.3.4: Unadjusted technique survival by Diabetes status, 1997-2006



CHAPTER 12

Chronic Peritoneal Dialysis Practices

Sunita Bavanandan Lily Mushahar

SECTION 1: PERITONEAL DIALYSIS (PD) PRACTICES

12.1: Mode of PD (Tables 12.1.1 to 12.1.4)

In 2006, CAPD still remained the commonest mode of PD therapy accounting for 90% of the total. Automated peritoneal dialysis (APD) penetration has been steadily increasing from 1% when it was first introduced in 2004 to 6% at present. As compared to PD practice in certain countries where Icodextrin is widely used in high transporter PD patients, a significant 4% of our patients were still practicing DAPD. This is most likely due to our local financial constraints. Most patients (92%) were on the Baxter disconnect system. The majority of patients (91%) were on 4 exchanges per day but there is a trend for an increased percentage of patients on 3 exchanges a day from 1% in 2004 to 4% presently. This may be a reflection of more aggressive management of advanced chronic kidney disease, with earlier initiation of dialysis allowing incremental dialysis. Conversely, up to 5% of patients were doing 5 exchanges per day. Most patients (82%) used a fill volume of 2L but up to 11% were using larger fill volumes while 7%, mainly paediatric, were on smaller exchange volumes than 2L.

Table 12.1.1: Chronic Peritoneal Dialysis Regimes, 1997-2006

DD ragima	199	97	199	98	199	9	200	00	200)1
PD regime	No.	%								
Standard CAPD	440	94	492	93	577	96	633	97	755	98
DAPD	26	6	32	6	16	3	16	2	17	2
Automated PD/ CCPD	4	1	6	1	6	1	5	1	2	0
TOTAL	470	100	530	100	599	100	654	100	774	100
-								_		_
DD regime	200)2	200)3	200)4	200)5	200)6
PD regime	No.)2 %	200 No.)3 %	200 No.)4 %	200 No.)5 %	200 No.)6 %
PD regime Standard CAPD		-				· ·				
	No.	%								
Standard CAPD	No. 837	97	No.	97	No.	% 96	No.	93	No.	90

Table 12.1.2: CAPD Connectology, 1997-2006

CAPD Connectology	199	97	199	98	199	99	200	00	200	01
CAPD Connectology	No.	%	No.	%	No.	%	No.	%	No.	%
UVXD	27	5	10	2	3	1	0	0	0	0
Baxter disconnect	461	93	511	95	347	58	235	39	436	57
B Braun disconnect	10	2	18	3	248	41	370	61	324	43
Fresenius disconnect	0	0	0	0	0	0	0	0	0	0
Others	0	0	0	0	0	0	0	0	0	0
TOTAL	498	100	539	100	598	100	605	100	760	100
CARD Compostology	20	02	200)3	200)4	200)5	200	06
CAPD Connectology	No.	%	No.	%	No.	%	No.	%	No.	%
UVXD	0	0	0	0	0	0	0	0	0	0
Baxter disconnect	719	87	1038	87	1142	88	1252	90	1423	92
B Braun disconnect	93	11	7	1	14	1	1	0	0	0
Fresenius disconnect	11	1	154	13	145	11	111	8	119	8
Others	0	0	1	0	0	0	28	2	6	0
TOTAL	823	100	1200	100	1301	100	1392	100	1548	100

Table 12.1.3: CAPD Number of Exchanges per day, 1997-2006

No. of	1997		1998		199	99	2000		2001	
Exchanges/ day	No.	%	No.	%	No.	%	No.	%	No.	%
2	0	0	2	0	0	0	2	0	1	0
3	3	1	4	1	4	1	1	0	5	1
4	454	97	508	96	579	97	624	96	735	95
5	12	3	16	3	13	2	23	4	31	4
TOTAL	469	100	530	100	596	100	650	100	772	100

No. of	200)2	200)3	200)4	200)5	200)6
Exchanges/ day	No.	%	No.	%	No.	%	No.	%	No.	%
2	0	0	4	0	6	0	3	0	4	0
3	11	1	14	1	12	1	25	2	55	4
4	834	96	1136	96	1225	95	1280	94	1357	91
5	28	3	32	3	52	4	48	4	74	5
TOTAL	873	100	1186	100	1295	100	1356	100	1490	100

Table 12.1.4: CAPD Volume per Exchange, 1997–2006

		-								
Volume per	199	97	199	98	199	99	200	00	200)1
Exchange (L)	No.	%								
<1.5	24	5	35	7	32	5	46	7	51	7
1.5-1.9	0	0	20	4	22	4	25	4	27	4
2.0	444	95	476	90	535	91	570	88	684	89
>2.0	0	0	0	0	2	0	7	1	9	1
TOTAL	468	100	531	100	591	100	648	100	771	100
Volume per	200	02	200	03	200)4	200)5	200	06
Exchange (L)	No.	%								
<1.5	64	7	64	5	73	6	85	6	67	4

<1.5	64	7	64	5	73	6	85	6	67	4
1.5-1.9	24	3	36	3	36	3	38	3	45	3
2.0	769	88	1051	85	1117	84	1153	83	1262	82
>2.0	14	2	80	6	101	8	113	8	168	11
TOTAL	871	100	1231	100	1327	100	1389	100	1542	100

SECTION 2: SOLUTE CLEARANCE ACHIEVEMENT AND PERITONEAL TRANSPORT

The median delivered weekly Kt/V has remained unchanged at 2.1 since 2003, with 59% of patients achieving K/DOQI recommended Kt/V of more than or equal to 2.0. Comparison between PD centres has shown a narrowing of the gap between the highest- and lowest- performing centers, with less than 3-fold variation in terms of the percentage of patients in each center achieving a Kt/V of > 2.0 per week. Half of the centers were able to have at least 58% of their patients achieving the K/DOQI target (Tables and figures 12.2.1 and 12.2.2).

Table 12.2.1: Distribution of delivered KT/V, CAPD patients 2003-2006

Year	No of Subjects	Mean	SD	Median	LQ	UQ	% patients ≥2.0 per week
2003	789	3.7	19.9	2.1	1.8	2.5	59
2004	1068	2.8	9.9	2.1	1.8	2.5	61
2005	1124	3.3	13.7	2.1	1.8	2.5	58
2006	1290	2.4	3.6	2.1	1.8	2.4	59

Figure 12.2.1: Cumulative distribution of delivered KT/V, CAPD patients 2003-2006

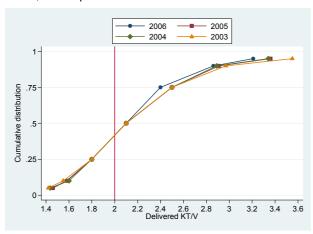


Figure 12.2.2: Variation in proportion of patients with $KT/V \ge 2.0$ per week among CAPD centres 2006

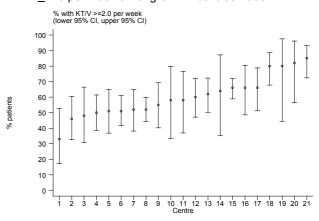


Table 12.2.2: Variation in proportion of patients with KT/V ≥ 2.0 per week among CAPD

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
2003	14	0	0	51	59	62	73	73
2004	17	43	43	53	56	67	85	85
2005	18	11	11	50	53.5	63	92	92
2006	21	33	46	51	58	66	82	85

Among incident PD patients low average transport status was commonest (42%) followed by high average transport status (37%). However, a proportion of patients developed changes in their membrane characteristics over time. There was high PET status in 15% of prevalent patients compared to 9% in new PD patients.

Table 12.2.3: Peritoneal transport status by PET D/P creatinine at 4 hours, new PD patients 2003-2006

PET	200)3	200	04	200	05	2006		
PEI	No.	%	No.	%	No.	%	No.	%	
Low	10	6	67	15	69	12	105	12	
Low average	85	51	187	41	246	41	360	42	
High average	62	37	176	38	223	37	314	37	
High	11	7	29	6	62	10	75	9	
TOTAL	168	100	459	100	600	100	854	100	

^{*} New PD patients=patients commencing dialysis since 2003

Table 12.2.4: Peritoneal transport status by PET D/P creatinine at 4 hours, prevalent PD patients 2003-2006

PET	200	03	200	04	200	05	2006		
PEI	No.	%	No.	%	No.	%	No.	%	
Low	10	3	39	9	44	13	23	8	
Low average	174	44	180	42	130	39	106	38	
High average	171	43	168	39	118	35	106	38	
High	39	10	41	10	42	13	41	15	
TOTAL	394	100	428	100	334	100	276	100	

^{*}Prevalent PD patients=patients commencing dialysis before 2003

SECTION 3: TECHNIQUE SURVIVAL ON PD

In terms of technique survival, CAPD fared worse compared to haemodialysis with Kaplan-Meir cumulative survival curves diverging as early as 6 months. One-, three- and five-year technique survival for CAPD was 81%, 47% and 29% as compared to haemodialysis at 89%, 71% and 56% respectively. Median technique survival time was less than 36 months. Overall these trends in technique survival remained unchanged by year of entry (Tables and figures 12.3.1 and 12.3.2).

The youngest age group between 1-14 years has the best technique survival rate while the oldest age group aged >65 years consistently had the worst technique survival (Table and figure 12.3.3). Diabetics have a poorer technique survival than the non-diabetics (Table and figure 12.3.4). However, there was no gender difference (Table and figure 12.3.5).

The commonest factor for PD drop-out between 1997-2006 was peritonitis (44%), followed by membrane failure (19%) and technical problems (19%) (Table12.3.9). Patient preference contributed to a further 13% of conversions to HD.

Table 12.3.1: Unadjusted technique survival by Dialysis modality, 1997-2006

Dialysis modality		CAPD			HD			All Dialysis	
Interval (months)	No.	% Survival	SE	No.	% Survival	SE	No.	% Survival	SE
6	2537	91	1	17560	94	0	20097	94	0
12	2087	81	1	14950	89	0	17037	88	0
24	1376	64	1	10955	79	0	12331	77	0
36	847	47	1	7895	71	0	8742	68	0
48	497	36	1	5592	63	0	6089	60	0
60	293	29	1	3809	56	0	4102	53	0
72	160	22	1	2536	51	1	2695	47	0
84	88	18	1	1578	45	1	1666	42	1
96	44	13	1	884	41	1	927	37	1
108	19	10	1	372	38	1	390	34	1
120	4	7	2	54	33	1	57	30	1

^{*} No. = Number at risk

Figure 12.3.1: Unadjusted technique survival by Dialysis modality, 1997-2006

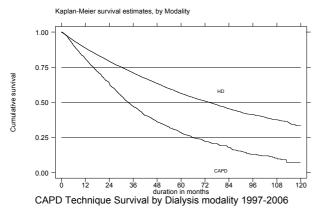


Figure 12.3.2: Unadjusted technique survival by year of entry, 1997-2006

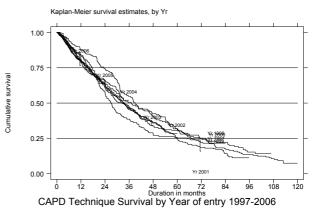


Table 12.3.2: Unadjusted technique survival by year of entry, 1997-2006

Year		1997			1998		1999				2000		
Interval (months)	No.	% Survival	SE	No.	% Survival	SE	No.	% Survival	SE	No.	% Survival	SE	
6	187	94	2	144	92	2	189	90	2	206	91	2	
12	170	88	2	128	83	3	175	84	3	185	81	3	
24	141	74	3	96	65	4	117	58	3	138	63	3	
36	101	55	4	75	51	4	78	39	3	101	46	3	
48	76	42	4	59	40	4	57	29	3	78	36	3	
60	57	32	3	45	32	4	50	25	3	67	31	3	
72	44	25	3	35	25	4	37	19	3	47	22	3	
84	32	18	3	31	22	3	27	15	3	1	-	-	
96	24	14	3	21	15	3	1	-	-	1	-	-	
108	19	11	2	1	-	-	1	-	-	1	-	-	
120	4	8	2	1	-	-	1	-	-	1	-	-	

Year		2001			2002		2003				2004		
Interval (months)	No.	% Survival	SE	No.	% Survival	SE	No.	% Survival	SE	No.	% Survival	SE	
6	303	90	2	341	92	1	370	89	2	301	89	2	
12	265	80	2	292	80	2	333	80	2	266	80	2	
24	198	61	3	227	63	3	254	63	2	213	66	3	
36	151	47	3	164	47	3	183	46	2	5	-	-	
48	107	34	3	125	36	3	1	-	-	5	-	-	
60	78	25	2	1	-	-	1	-	-	5	-	-	

Year		2005			2006	
Interval (months)	No.	% Survival	SE	No.	% Survival	SE
6	321	89	2	183	92	2
12	280	79	2	2	-	-

^{*} No. = Number at risk

SE=standard error

Table 12.3.3: Unadjusted technique survival by age, 1997-2006

Age group (years)		<=14		15-24			25-34			35-44		
Interval (months)	No.	% Survival	SE	No.	% Survival	SE	No.	% Survival	SE	No.	% Survival	SE
6	229	98	1	253	93	2	238	93	2	355	94	1
12	204	95	1	210	84	2	203	86	2	298	86	2
24	142	84	3	139	72	3	150	75	3	208	71	2
36	106	73	3	95	59	3	112	65	3	144	57	3
48	76	66	4	55	48	4	68	49	4	96	46	3
60	46	57	4	30	40	4	50	43	4	50	34	3
72	29	48	5	17	35	5	27	31	4	27	26	3
84	15	44	6	7	23	6	19	26	4	15	23	4
96	7	29	7	4	17	7	12	19	4	8	20	4
108	4	29	7	2	9	7	5	16	5	4	15	5
120	1	-	-	1	-	-	1	-	-	3	-	

Age group (years)		45-54			55-64		>=65			
Interval (months)	No.	% Survival	SE	No.	% Survival	SE	No.	% Survival	SE	
6	615	92	1	543	90	1	310	80	2	
12	513	82	1	428	78	2	236	65	2	
24	338	63	2	274	58	2	131	42	3	
36	192	45	2	155	39	2	49	21	2	
48	104	33	2	83	26	2	21	13	2	
60	73	28	2	40	18	2	9	8	2	
72	41	21	2	20	13	2	5	6	2	
84	23	17	2	11	9	2	4	4	2	
96	14	14	2	3	3	2	2	4	2	
108	7	10	3	2	3	2	1	-	-	
120	2	6	3	1	-	-	1	-	-	

^{*} No. = Number at risk

SE=standard error

Figure 12.3.3: Unadjusted technique survival by age, 1997-2006

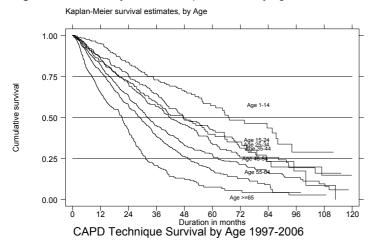


Table 12.3.4:	Unadjusted	technique survival b	v Gender	. 1997-2006
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Gender		Male			Female	
Interval (months)	No.	% Survival	SE	No.	% Survival	SE
6	1270	91	1	1267	90	1
12	1040	81	1	1048	81	1
24	682	63	1	695	64	1
36	415	45	2	433	49	1
48	230	34	2	268	39	2
60	131	26	2	163	31	2
72	73	20	2	88	24	2
84	40	16	2	50	20	2
96	22	13	2	23	14	2
108	10	11	2	10	10	2
120	2	5	3	3	9	2

^{*} No. = Number at risk

Figure 12.3.4: Unadjusted technique survival by Gender, 1997-2006

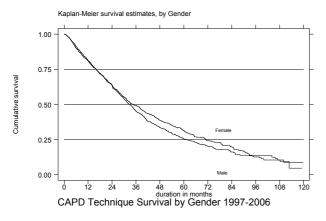


Figure 12.3.5: Unadjusted technique survival by Diabetes status, 1997-2006

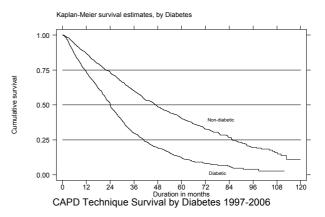


Table 12.3.5: Unadjusted technique survival by Diabetes status, 1997-2006

Diabetes status		Non-Diabetic			Diabetic	
Interval (months)	No.	% Survival	SE	No.	% Survival	SE
6	1497	94	1	1040	86	1
12	1277	87	1	810	74	1
24	909	73	1	469	51	2
36	621	60	1	227	30	2
48	389	49	2	109	20	1
60	236	40	2	58	13	1
72	132	33	2	29	8	1
84	76	27	2	13	6	1
96	39	20	2	6	4	1
108	18	16	2	3	3	1
120	4	11	3	1	-	-

^{*} No. = Number at risk

Table 12.3.6: Reasons for change of dialysis modality to HD, 1997-2006.

Cause	N	Percentage
Peritonitis	287	44
Catheter related infection	21	3
Membrane failure	123	19
Technical problem	55	8
Patient preference	88	13
Others	52	8
Unknown	27	4
Total	653	100

SE=standard error

SE=standard error

SECTION 4: PD PERITONITIS

The median peritonitis rate was 33 patient-months per episode (Table 12.4.1). There was a wide variation between centre with the highest and lowest peritonitis rates of 16 vs 103 patient-months per episode. Gram-positive organisms accounted for 32% of peritonitis episodes while 22% were due to gram positive organisms. Fungal organisms accounted for 4% of cases and mycobacterial infection rates remained low (1%). The culture–negative rate increased to 39% compared with 30% in 2005 (Table 12.4.2). There was a trend for increased peritonitis rates with increasing age and diabetic status but gender did not appear to have any influence (Table 12.4.3).

Catheter removal rate was highest in fungal infection (75%), followed by pseudomonas (36%) and staphylococcus aureus (21%) infections. Pseudomonas aeruginosa peritonitis has the highest mortality outcome of 23% compared to other organisms (Table 12.4.4).

Table 12.4.1: Variation in peritonitis rate (pt-month/epi) among CAPD centres 2006

Year	No. of centres	Min	5 th Centile	LQ	Median	UQ	95 th Centile	Max
2000	12	10.9	10.9	18	21.5	27.6	1019.7	1019.7
2001	11	13.4	13.4	18.5	22.7	30.9	55.5	55.5
2002	13	14.3	14.3	20.2	25.3	34.4	49	49
2003	13	16.5	16.5	22.8	30.1	41.6	253	253
2004	15	0	0	23.2	32.6	41.8	47.7	47.7
2005	15	23.6	23.6	28	35.7	41.8	66.3	66.3
2006	21	16.1	21	27.7	33	61.4	88.5	102.9

^{*} Criteria for combination of centre with less than 10 subjects not applied

Figure 12.4.1: Variation in peritonitis rate among CAPD centres 2006

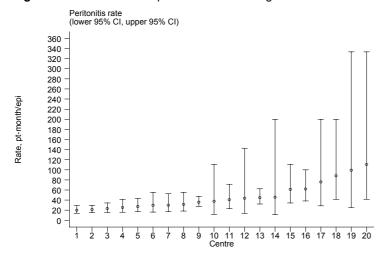


Table 12.4.2: Causative organism in PD peritonitis, 2000-2006

Microorganism	20	00	200)1	200)2	200)3	200)4	200)5	200	06
Wilchoorganism	No.	%												
(A) Gram Positives														
 Staph. Aureus 	35	11	41	13	62	17	45	12	51	14	40	12	50	14
 Staph Coagulase Neg. 	39	13	34	11	41	11	52	14	43	12	47	15	35	10
Strep	12	4	13	4	9	2	11	3	11	3	6	2	13	4
Others	4	1	6	2	7	2	15	4	4	1	8	2	13	4
(B) Gram Negatives														
 Pseudomonas 	19	6	14	4	23	6	20	5	28	8	27	8	23	6
Others	45	15	56	18	67	19	75	21	83	22	86	27	58	16
(C) Polymicrobial	9	3	10	3	8	2	3	1	2	1	0	0	1	0
(D) Others														
Fungal	19	6	21	7	12	3	12	3	15	4	7	2	16	4
 Mycobacterium 	6	2	4	1	1	0	3	1	4	1	2	1	4	1
Others	2	1	14	4	14	4	13	4	8	2	3	1	11	3
(E) No growth	119	39	99	32	118	33	115	32	123	33	96	30	142	39
TOTAL	309	100	312	100	362	100	364	100	372	100	322	100	366	100

Table 12.4.3: Factors influencing peritonitis rate, 2000-2006

Factors	N (No. at risk)	Annualised rate: Epi/ pt-year	(95)	% CI)
Age (years):				
<=14	69	0.405	(0.33	0.498)
15-24	38	0.465	(0.354	0.612)
25-34	82	0.416	(0.35	0.495)
35-44	94	0.444	(0.373	0.528)
45-54	143	0.523	(0.455	0.6)
55-64	121	0.571	(0.489	0.667)
>=65	51	0.663	(0.52	0.845)
Gender:				
Male	282	0.492	(0.444	0.545)
Female	316	0.482	(0.44	0.528)
Diabetes:				
No	414	0.456	(0.421	0.494)
Yes	184	0.591	(0.519	0.674)

Table 12.4.4: Outcome of peritonitis by Causative organism, 2006

			Outo	come				
Causative Organism	Reso	olved		solved, removed	De	ath	To	otal
	No.	%	No.	%	No.	%	No.	%
Gram Positives								
 Staph. Aureus 	33	69	10	21	5	10	48	100
 Staph Coagulase Neg. 	28	85	1	3	4	12	33	100
Gram Negatives								
 Pseudomonas 	9	41	8	36	5	23	22	100
Others	35	60	11	19	12	21	58	100
Others								
Fungal	1	6	12	75	3	19	16	100
 Mycobacterium 	0	0	2	50	2	50	4	100
No growth	94	68	21	15	23	17	138	100

CHAPTER 13

Renal Transplantation

Editor: Dr. Goh Bak Leong

Expert Panel:
Dato' Dr. Zaki Morad (Chair)
Dr. Goh Bak Leong (Co-chair)
Dr. Fan Kin Sing
Dr. Lily Mushahar
Mr. Rohan Malek
Dr. S Prasad Menon
Prof. Dr. Tan Si Yen

13.1 STOCK AND FLOW

New renal transplant patients showed a modest increase from 126 transplants per year in 1997 to 161 per year in 2005. By end of 2006, the number of functioning renal transplants reported to NRR has increased to 1725 (Table 13.1.1).

Table 13.1.1: Stock and Flow of Renal Transplantation, 1997-2006

Year	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006
New transplant patients	126	104	127	143	161	168	160	189	161	126
Died	31	26	25	30	37	32	37	41	41	49
Graft failure	38	49	36	32	40	38	41	44	20	30
Lost to follow up	0	1	1	9	2	3	5	16	7	5
Functioning graft at 31st December	1083	1111	1176	1248	1330	1425	1502	1590	1683	1725

Figure 13.1.1: Stock and Flow of Renal Transplantation, 1975-2006

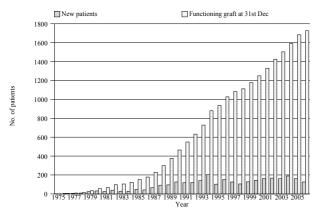
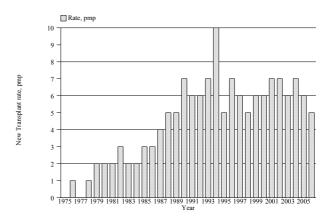


Figure 13.1.2: New transplant rate, 1975-2006



Incident rate for renal transplantation stabilised at a modest rate of 5-7 per million population for the last decade (Table 13.1.2), while the transplant prevalence rate maintained at 50-65 per million population for the last 10 years (Table 13.1.3).

Table 13.1.2: New transplant rate per million population (pmp), 1997-2006

Year	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006
New transplant patients	126	104	127	143	161	168	160	189	161	126
New transplant rate, pmp	6	5	6	6	7	7	6	7	6	5

Table 13.1.3: Transplant prevalence rate per million population (pmp), 1997-2006

Year	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006
Functioning graft at 31st December	1083	1111	1176	1248	1330	1425	1502	1590	1683	1725
Transplant prevalence rate, pmp	50	50	52	53	55	58	60	62	64	65

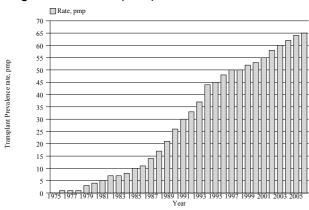


Figure 13.1.3: Transplant prevalence rate, 1975-2006

Table 13.1.4: Place of transplantation, 1997-2006

Year	19	97	19	98	19	99	20	00	20	01
	No.	%								
HKL	29	23	33	32	36	28	28	20	33	20
UMMC	6	5	7	7	16	13	19	13	23	14
Selayang Hospital	0	0	0	0	0	0	4	3	11	7
Other local	0	0	0	0	1	1	3	2	4	2
China	79	63	50	48	62	49	80	56	82	51
India	7	6	7	7	5	4	9	6	7	4
Other overseas	3	2	3	3	2	2	0	0	1	1
Unknown	2	2	4	4	5	4	0	0	0	0
TOTAL	126	100	104	100	127	100	143	100	161	100

Year	20	02	20	03	20	04	20	05	20	06	TOT	ĀL
	No.	%	No.	%	No.	No.	%	No.	%	No.	%	No.
HKL	28	17	26	16	20	11	32	20	34	27	331	20
UMMC	14	8	6	4	7	4	7	4	5	4	117	7
Selayang Hospital	11	7	11	7	11	6	5	3	7	6	60	4
Other local	1	1	1	1	2	1	5	3	2	2	19	1
China	102	61	111	69	136	72	107	66	71	56	985	61
India	12	7	4	3	11	6	5	3	6	5	79	5
Other overseas	0	0	1	1	2	1	0	0	0	0	13	1
Unknown	0	0	0	0	0	0	0	0	1	1	12	1
TOTAL	168	100	160	100	189	100	161	100	126	100	1616	100

13.2 RECIPIENTS' CHARACTERISTICS

Table 13.2.1: Renal Transplant Recipients' Characteristics, 1997-2006

Year	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006
New Transplant Patients	126	104	127	143	161	168	160	189	161	126
Age at transplant (years), Mean	36	37	37	40	41	41	42	41	38	37
Age at transplant (years), SD	12	11	13	13	13	13	13	13	14	16
% Male	63	58	61	64	63	57	66	62	70	68
% Diabetic (co-morbid / primary renal disease)	11	9	10	15	19	15	22	21	19	21
% HBsAg positive	6	6	4	5	5	7	8	5	4	7
% Anti-HCV positive	7	18	11	8	15	9	10	8	2	7

The mean age for new transplant recipients is between 36+6 years to 42+13 years over the last 10 years (Table 13.2.1). Men are still in the majority among renal transplant recipients and they made up 68% of all recipients in year 2006. Over the last 10 years, the proportion of diabetic transplant recipients has increased, from 11% in 1997 to about 21% in 2006.

In 2006, 7% were HbsAg positive and 7% had anti-HCV antibodies at the time of transplantation. The proportion of HbsAg positivity had reduced from 10-20% in the period 1985-1994 to 3-7% for the last 5 years while the number of recipients with anti-HCV antibodies at the time of transplantation had also reduced from 20-30% in the early 1990's to 2-10% for the last 5 years since the screening test was introduced in 1989. For those transplanted prior to the screening test, anti-HCV antibodies were found in 40-60%.

Chronic glomerulonephritis was the primary cause of ESRF in 25-37% for the last 5 years (Table 13.2.2). As expected, patients with diabetes mellitus had become increasingly frequent renal transplant recipients, from 7% in 1997 to 17% in 2006. Majority of renal transplant recipients still presented late with unknown primary renal disease, contributing to 30-44% of all the recipients for the last 5 years.

Table 13.2.2: Primary causes of end stage renal failure, 1997-2006

Year	19	97	19	98	19	99	20	00
	No.	%	No.	%	No.	%	No.	%
New transplant patients	126	100	104	100	127	100	143	100
Glomerulonephritis	30	24	28	27	41	32	49	34
Diabetes Mellitus	9	7	5	5	10	8	16	11
Hypertension	5	4	5	5	7	6	18	13
Obstructive uropathy	3	2	4	4	4	3	3	2
ADPKD	2	2	1	1	1	1	3	2
Drugs/toxic nephropathy	0	0	0	0	0	0	0	0
Hereditary nephritis	0	0	0	0	0	0	0	0
Unknown	63	50	55	53	62	49	54	38
Others	18	14	10	10	6	5	12	8

Year	20	01	20	02	20	03	20	04	20	05	20	06
	No.	%										
New transplant patients	161	100	168	100	160	100	189	100	161	100	126	100
Glomerulonephritis	42	26	53	32	54	34	62	33	46	29	47	37
Diabetes Mellitus	23	14	16	10	26	16	32	17	27	17	21	17
Hypertension	17	11	24	14	25	16	51	27	38	24	26	21
Obstructive uropathy	3	2	2	1	2	1	4	2	3	2	4	3
ADPKD	1	1	3	2	5	3	4	2	3	2	1	1
Drugs/toxic nephropathy	0	0	0	0	2	1	2	1	0	0	0	0
Hereditary nephritis	0	0	0	0	0	0	1	1	0	0	0	0
Unknown	61	38	68	40	58	36	83	44	48	30	38	30
Others	23	14	15	9	12	8	27	14	18	11	15	12

13.3 TRANSPLANT PRACTICES

In 2006, commercial transplants from China constituted 57% of all new renal transplantation (dropped from 65% in2005). While live donor transplantation made up 21% in 2006, local cadaveric transplants contributed 19% (increased from 6% in 2005) of all new renal transplantation in 2006 (Table 13.3.1).

Table 13.3.1: Type of Renal Transplantation, 1997-2006

Year	19	97	19	98	19	99	20	00
	No.	%	No.	%	No.	%	No.	%
Commercial Cadaver	80	66	51	52	62	51	80	56
Commercial Live Donor	7	6	4	4	4	3	9	6
Live Donor (genetically related)	27	22	27	27	40	33	21	15
Live Donor (emotionally related)	0	0	2	2	5	4	6	4
Cadaver	8	7	15	15	10	8	27	19
Total	122	100	99	100	121	100	143	100

Year	20	01	20	02	20	03	20	04	20	05	20	06
	No.	%										
Commercial Cadaver	82	51	102	61	112	70	141	76	104	65	68	57
Commercial Live Donor	6	4	11	7	3	2	5	3	8	5	2	2
Live Donor (genetically related)	32	20	30	18	25	16	21	11	37	23	22	18
Live Donor (emotionally related)	4	2	3	2	5	3	2	1	3	2	4	3
Cadaver	37	23	22	13	15	9	17	9	9	6	23	19
Total	161	100	168	100	160	100	186	100	161	100	119	100

^{*}Commercial Cadaver (China, India, other oversea) *Commercial live donor (living unrelated) *Cadaver (local)

Table 13.3.2: Biochemical data, 2004-2006

Biochemical parameters	Summary	2004	2005	2006
Creatinine, umol/L	N	1550	1635	1559
	Mean	132	133.7	135.3
	SD	63.8	65.4	80.4
	Median	120	120	119
	Minimum	38	35	13.2
	Maximum	817	763	1152
Hb, g/dL	N	1550	1635	1559
	Mean	12.9	12.8	12.7
	SD	1.9	1.9	1.9
	Median	12.9	12.9	12.8
	Minimum	4.9	5.5	3.3
	Maximum	19.7	19	19.8
Albumin, g/L	N	1550	1635	1559
	Mean	39.4	39.5	39.5
	SD	1	0.5	0.7
	Median	39.4	39.4	39.4
	Minimum	22	34	29
	Maximum	50	46	48
Calcium, mmol/L	N	1550	1635	1559
	Mean	2.4	2.3	2.3
	SD	0.2	0.2	0.2
	Median	2.3	2.3	2.3
	Minimum	1.1	1.2	1.1
	Maximum	3.3	3.3	3.1

Table 13.3.2: Biochemical data, 2004-2006 (cont'd)

Biochemical parameters	Summary	2004	2005	2006
Phosphate, mmol/L	N	1550	1635	1559
	Mean	1.1	1.1	1.1
	SD	0.2	0.2	0.2
	Median	1.1	1.1	1.1
	Minimum	0.3	0.3	0.4
	Maximum	2.7	3.3	3.5
Alkaline Phosphate (ALP), U/L	N	1550	1635	1559
	Mean	79.5	78.9	78.1
	SD	46.5	46.6	42.6
	Median	73	73	70
	Minimum	10	18	6.3
	Maximum	994	831	700
ALT, U/L	N	1550	1635	1559
	Mean	31.4	30.8	29.9
	SD	32.6	30.9	30.7
	Median	25	24	21
	Minimum	4	4	4
	Maximum	563	613	433
Total cholesterol, mmol/L	N	1550	1635	1559
	Mean	5.5	5.4	5.3
	SD	1.1	1	1.1
	Median	5.4	5.4	5.4
	Minimum	2.6	2.1	2
	Maximum	20	13.1	14.7
LDL cholesterol, mmol/L	N	1550	1635	1559
,	Mean	3.1	3	3
	SD	0.7	0.8	0.8
	Median	3	3	3
	Minimum	1	0.9	1
	Maximum	8.5	9.2	11.1
HDL cholesterol, mmol/L	N	1550	1635	1559
TIDE GROUDGE, TIME	Mean	1.6	1.6	1.6
	SD	0.4	0.5	0.5
	Median	1.6	1.6	1.6
	Minimum	0.2	0.2	0.2
	Maximum	4.3	5.6	5.8
Systolic Blood Pressure, mmHg	N	1550	1635	1559
dystolic blood i ressure, mining	Mean	132.2	133.3	130.8
	SD	15.9	16.9	16
	Median	130	130	130
	Minimum	80	80	66
	Maximum	200	220	210
Diastolic Blood Pressure, mmHg	N	1550	1635	1559
Diastolic blood Flessule, Illing		80.3		79
	Mean		80.5	
	SD Madian	9.6	9.2	9.8
	Median	80	80	80
	Minimum	40	50	30
	Maximum	121	127	120

Cyclosporine/prednisolone based triple therapy has remained the backbone of maintenance immunosuppressive therapy. In year 2006, 76% of renal transplant recipients were on cyclosporine while 97% were on prednisolone. Only 17% were on tacrolimus. However, 48% of the recipients were on MMF as opposed to 33% on azathioprine.

Table 13.3.3: Medication data, 2004-2006

Medication data		Sir	ngle drug	treatme	ent			Com	bined dru	ug treat	ment	
	200	04	200	05	20	06	20	04	20	05	20	06
	N	%	N	%	N	%	N	%	N	%	N	%
All patients	1416	100	1562	100	1470	100	1416	100	1562	100	1470	100
(i) Immunosuppressiv	re drug(s)) treatm	ent									
Prednisolone	13	1	12	1	7	0	1394	98	1528	98	1433	97
Azathioprine	0	0	1	0	0	0	603	43	605	39	492	33
Cyclosporin A	4	0	4	0	5	0	1135	80	1221	78	1111	76
Tacrolimus (FK506)	0	0	0	0	0	0	185	13	224	14	252	17
Mycophenolate Mofetil (MMF)	1	0	0	0	0	0	524	37	682	44	704	48
Rapamycin	0	0	0	0	0	0	5	0	8	1	6	0
Others	1	0	0	0	0	0	16	1	10	1	18	1
(ii) Non-Immunosupp	ressive d	rug(s) tr	eatment									
Beta blocker	104	7	105	7	77	5	650	46	667	43	594	40
Calcium channel blocker	188	13	195	12	199	14	795	56	822	53	784	53
ACE inhibitor	35	2	60	4	39	3	272	19	342	22	292	20
AIIRB	11	1	20	1	27	2	76	5	160	10	139	9
Anti-lipid	74	5	67	4	154	10	567	40	601	38	672	46
Other anti- hypertensive	5	0	5	0	11	1	130	9	158	10	159	11

In 2006, 65% of the recipients had hypertension as co-morbidity before transplantation while another 22% developed hypertension post transplantation (Table 13.4.1). Among these patients, only 30% were on monotherapy while the rest were on multiple drug treatment. For those on combination therapy, majority was on calcium channel blockers (53%) and beta blockers (40%). Only 20% were on ACE inhibitors while another 9% were on AIIRBs.

13.4 TRANSPLANT OUTCOMES

13.4.1 Post-transplant complications

Table 13.4.1: Post transplant complications, 2004-2006

Post transplant complications			n develop less of co transpla	omplica	tion after		C	omplica	ation dev transpla	•	only afte	er
	20	04	20	05	20	06	20	04	20	05	20	06
	N	%	N	%	N	%	N	%	N	%	N	%
All patients	1558	100	1637	100	1559	100	1558	100	1637	100	1559	100
Diabetes (either as Primary Renal Disease or co-morbid)	520	33	532	32	527	34	128	8	123	8	125	8
Cancer	3	0	2	0	1	0	17	1	19	1	20	1
Cardiovascular disease + cerebrovascular disorder	208	13	209	13	207	13	83	5	45	3	45	3
Hypertension	1010	65	1047	64	1016	65	397	25	439	27	346	22

In 2006, 34% of the prevalent renal transplant recipients had diabetes mellitus before transplantation (either as primary renal disease or co-morbidity), while another 8% of them developed diabetes mellitus post transplantation (PTDM).

 $^{^{\}star}$ Hypertension: BP systolic > 140 and BP diastolic >90 OR have either Beta blocker / Calcium channel blocker / ACE inhibitor / AIIRB / Other anti-hypertensive

13.4.2 Deaths and Graft loss

In 2006, 49 (3%) transplant recipients died and 30 (2%) lost their grafts. These rates of transplant death and graft loss have remained constant for the last 10 years (Table 13.4.2). Infection, cardiovascular disease and death at home were among the commonest causes of death for the last decade and in 2006, they accounted for 41%, 19% and 13% of the causes of death respectively (Table 13.4.3). Death secondary to cancer has become more common from 2001 to 2004, whereby cancer death accounted for 13% to 18% of all causes of death during that period. However, in 2006, death due to cancer has dropped to 7% of all deaths. Renal allograft rejection accounted for 50-78% of graft losses for the last 10 years (Table 13.4.4).

Table 13.4.2: Transplant Patients Death Rate and Graft Loss, 1997-2006

Year	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006
No. at risk	1054	1096	1143	1211	1288	1377	1463	1545	1636	1703
Transplant death	31	26	25	30	37	32	37	41	41	49
Transplant death rate %	3	2	2	2	3	2	3	3	3	3
Graft loss	38	49	36	32	40	38	41	44	20	30
Graft loss rate %	4	4	3	3	3	3	3	3	1	2
Acute rejection	0	0	0	0	0	0	3	19	14	17
Acute rejection rate %	0	0	0	0	0	0	0	1	1	1
All losses	69	75	61	62	77	70	81	104	75	96
All losses rate %	7	7	5	5	6	5	6	7	5	6

^{*}Graft loss=graft failure

Figure 13.4.2(i): Transplant Recipient Death Rate, 1975-2006

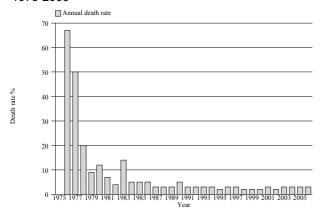
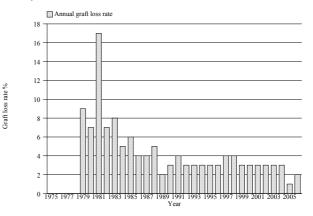


Figure 13.4.2(ii): Transplant Recipient Graft Loss Rate, 1975-2006



^{*}All losses=death / graft loss (acute rejection happens concurrently with graft failure/ death)

Table 13.4.3: Causes of Death in Transplant Recipients, 1997-2006

Year	19	1997	1998	38	1999	6(2000	00	2001	7	2002	2	2003	တ္	2004	4	2005	5	2006	9
	Š.	%	Š.	%	Š.	%	No.	%	o N	%	O	%	O	%	No.	%	Š.	%	Š	%
Cardiovascular	4	13	က	11	4	13	10	29	7	16	2	16	6	23	4	6	2	12	10	19
Died at home	7	9	4	15	9	19	_	က	2	12	2	16	2	13	9	13	4	10	7	13
Infection	4	45	10	37	7	23	12	35	20	47	6	28	7	28	7	24	22	52	22	41
Graft failure	_	က	0	0	0	0	7	9	0	0	0	0	0	0	7	4	0	0	0	0
Cancer	0	0	က	7	ဗ	10	7	9	9	4	4	13	9	15	80	18	4	10	4	7
Liver disease	7	9	7	7	ဗ	10	_	က	_	7	က	6	7	2	က	7	က	7	4	7
Accidental death	0	0	0	0	~	ဗ	_	က	_	7	_	က	0	0	0	0	0	0	0	0
Others	4	13	7	7	2	16	က	6	7	2	က	0	2	13	10	22	က	7	4	7
Unknown	4	13	က	7	7	9	7	9	_	7	7	9	7	2	_	7	_	7	က	9
TOTAL	31	100	27	100	31	100	34	100	43	100	32	100	40	100	45	100	42	100	54	100

Table 13.4.4: Causes of Graft Failure, 1997-2006

Year	1997	97	1998	98	19(666	2000	00	2001	21	2002)2	2003)3	2004	4	2005)5	2006	9(
	Š	%	Š.	%	o N	%	Š.	%	Š.	%	o N	%	ė.	%	Š.	%	No.	%	o N	%
Rejection	21	54	28	53	23	64	19	29	25	61	22	55	22	20	33	70	18	78	23	99
Calcineurin toxicity	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	_	က
Other drug toxicity	_	က	0	0	0	0	0	0	0	0	0	0	0	0	_	7	0	0	0	0
Ureteric obstruction	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
Infection	0	0	_	7	0	0	_	က	7	2	0	0	7	2	_	7	~	4	က	6
Vascular causes	4	10	က	9	_	က	က	6	_	7	0	0	က	7	4	6	2	6	က	6
Recurrent/de novo renal disease	_	က	~	7	0	0	0	0	7	2	7	2	~	7	~	7	0	0	~	က
Others	2	13	2	6	0	0	7	9	0	0	4	10	-	7	0	0	~	4	7	9
Unknown	7	18	15	28	12	33	7	22	7	27	12	30	15	34	7	15	_	4	7	9
TOTAL	39	100	53	100	36	100	32	100	41	100	40	100	44	100	47	100	23	100	35	100

13.5 Patient and Graft Survival

Table 13.5.1: Patient survival, 1993-2006

Interval (years)	No.	% Survival	SE
1	1778	95	1
3	1342	91	1
5	971	88	1
10	349	80	1

^{*} No.=Number at risk SE=standard error

Figure 13.5.1: Patient survival, 1993-2006

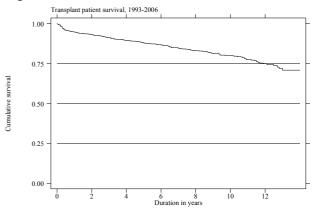
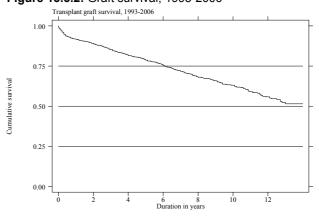


Table 13.5.2: Graft survival, 1993-2006

Interval (years)	No.	% survival	SE
1	1778	92	1
3	1342	85	1
5	971	79	1
10	349	63	1

^{*} No.=Number at risk SE=standard error

Figure 13.5.2: Graft survival, 1993-2006



The overall transplant patient survival rate from 1993 to 2006 was 95%, 91%, 88% and 80% at 1 year, 3 years, 5 years and 10 years respectively, while the overall graft survival rate was 92%, 85%, 79% and 63% respectively.

Outcomes of renal transplantation from the four donor groups are shown in Figures 13.5.3 and 13.5.4 and demonstrate substantially different patient and graft survival rates. Living donor grafts maintained the best patient and graft survival rates. The 1, 3, 5 and 10 year patient survival rate for recipients of living donor grafts were 96%, 95%, 94% and 90% respectively. The graft survival rates also differed between these 4 groups; living and commercial cadaver donor graft had the best outcomes.

Table 13.5.3: Patient survival by type of transplant, 1993-2006

Type of Transplant		Commercial Cadaver			Commercial Live Donor			Live Donor			Cadaver	
Interval (years)	No.	% Survival	SE	No.	% Survival	SE	No.	% Survival	SE	No.	% Survival	SE
1	935	96	1	285	96	1	398	96	1	131	84	2
3	659	92	1	240	90	1	323	95	1	100	79	3
5	428	87	1	203	87	2	252	94	1	71	76	4
10	117	83	2	126	72	3	100	90	2	7	70	5

^{*} No.=Number at risk SE=standard error

Table 13.5.4: Graf survival by type of transplant, 1993-2006

Type of Transplant		Commercial Cadaver			Commercial Live Donor			Live Donor			Cadaver	
Interval (years)	No.	% Survival	SE	No.	% Survival	SE	No.	% Survival	SE	No.	% Survival	SE
1	935	94	1	285	93	1	398	92	1	131	76	3
3	659	89	1	240	82	2	323	88	2	100	68	4
5	428	82	1	203	73	3	252	84	2	71	64	4
10	117	70	2	126	52	3	100	69	3	7	48	8

^{*} No.=Number at risk SE=standard error

Figure 13.5.3: Patient survival by type of transplant, 1993-2006

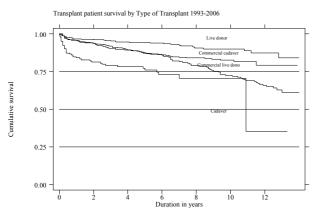
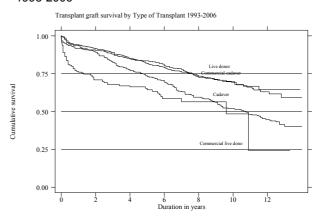


Figure 13.5.4: Graf survival by type of transplants, 1993-2006



The patient and graft survival rates for 1993-1998 cohort and 1999-2005 cohort were compared. Patient survival rate for living related donor renal transplants has remained excellent and unchanged for these two cohorts (Figure 13.5.5).

Figure 13.5.5: Patient survival by year of transplant (Living related transplant, 1993-2006)

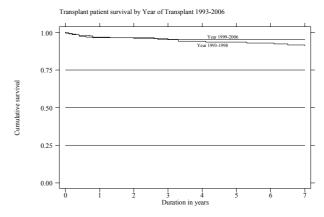


Table 13.5.5: Patient survival by year of transplant (Living related transplant, 1993-2006)

Year of Transplant		1993-1998			1999-2006	
Interval (years)	No.	% Survival	SE	No.	% Survival	SE
1	181	97	1	218	96	1
3	169	95	2	155	95	1
5	159	93	2	94	95	1
7	147	91	2	37	95	1

^{*} No.=Number at risk SE=standard error

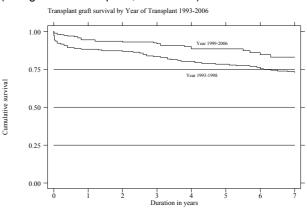
Interestingly, the risk of graft failure for living related donor renal transplantation improved for the 1999-2006 cohort compared to the 1993-1998 cohort (Table & Figure 13.5.6). One possible explanation, among others, is the increasing use of newer immunosuppressive agents such as MMF and FK506 in recent years.

Table 13.5.6: Graf survival by year of transplant (Living related transplant, 1993-2006)

Year of Transplant		1993-1998			1999-2006	
Interval (years)	No.	% Survival	SE	No.	% Survival	SE
1	181	88	2	218	94	1
3	169	83	3	155	92	2
5	159	78	3	94	89	2
7	147	73	3	37	83	3

^{*} No.=Number at risk SE=standard error

Figure 13.5.6: Graft survival by year of transplant (Living related transplant, 1993-2006)



Interestingly, our data showed that commercial cadaveric transplants have excellent patient and graft survival rates, which are comparable to living related donor transplants for both 1993-1998 and 1999-2006 cohorts (Figure 13.5.7 and 13.5.8).

Table 13.5.7: Patient survival by year of transplant (Commercial cadaver transplant, 1993-2006)

`		•			,	
Year of Transplant		1993-1998		1	999-2006	
Interval (years)	No.	% Survival	SE	No.	% Survival	SE
1	288	94	1	648	96	1
3	275	92	2	384	92	1
5	247	87	2	181	87	2
7	225	84	2	50	85	2

^{*} No.=Number at risk SE=standard error

Table 13.5.8: Graft survival by year of transplant (Commercial cadaver transplant, 1993-2006)

Year of Transplant		1993-1998	3	1	1999-2006	
Interval (years)	No.	% Survival	SE	No.	% Survival	SE
1	288	93	1	648	95	1
3	275	89	2	384	89	1
5	247	80	2	181	83	2
7	225	73	3	50	78	3

^{*} No.=Number at risk SE=standard error

Figure 13.5.7: Patient survival by year of transplant (Commercial cadaver transplant, 1993-2006)

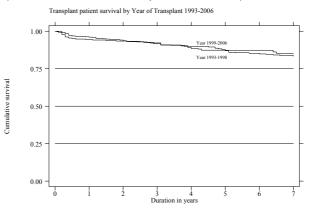
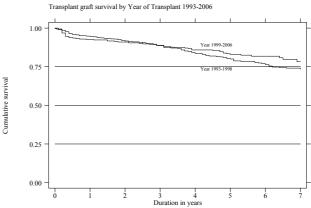


Figure 13.5.8: Graft survival by year of transplant (Commercial cavader transplant, 1993-2006)



13.6 CARDIOVASCULAR RISK IN RENAL TRANSPLANT RECIPIENTS

13.6.1 Risk factors for Ischaemic Heart Disease.

In year 2006, 85.5% of recipients were hypertensive, 22.9% had diabetes and 57.1% had renal insufficiency fulfilling the criteria for CKD III and above. A majority had 2 or more cardiovascular risk factors with 10.2% having 3 major risk factors.

Table 13.6.1: Risk factors for IHD in renal transplant recipients at year 2004, 2005, and 2006

	2004	2005	2006
Diabetes	27 (1.9)	19 (1.3)	21 (1.4)
Hypertension**	504 (34.3)	513 (33.5)	452 (31.5)
CKD	121 (8.2)	142 (9.3)	169 (11.8)
Diabetes + Hypertension**	146 (9.9)	157 (10.2)	144 (10.0)
Diabetes + CKD	21 (1.4)	20 (1.3)	18 (1.3)
CKD + Hypertension**	530 (36.1)	540 (35.2)	484 (33.8)
Diabetes + CKD + Hypertension**	120 (8.2)	141 (9.2)	146 (10.2)

^{**} Hypertension: BP systolic > 140 and BP diastolic >90

Figure 13.6.1(a): Venn Diagram for Pre and Post Transplant Complications (in %) at year 2004

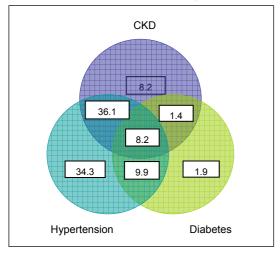


Figure 13.6.1(c): Venn Diagram for Pre and Post Transplant Complications (in %) at year 2006

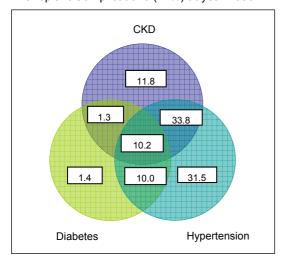
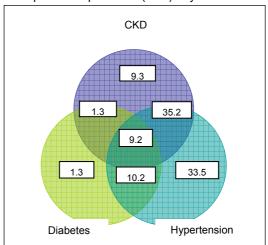


Figure 13.6.1(b): Venn Diagram for Pre and Post Transplant Complications (in %) at year 2005



OR have either Beta blocker / Calcium channel blocker/ ACE inhibitor/ AIIRB / Other anti-hypertensive drugs

 $GFR(mL/min/1.73m2) = 1.2*(140-age(year))* \ weight(kg) \ / \ creatinine(\mu mol/L) \ if \ male$

GFR(mL/min/1.73m2) = $0.85*(1.2*(140-age(year))*weight(kg) / creatinine(\mu mol/L))$ if female.

CKD stage III - GFR, 30-60

CKD stage IV - GFR, 15-30

CKD stage V – GFR, < 15

13.6.2 Blood Pressure classification according to JNC VI criteria, 2004, 2005, and 2006

In 2006, 5.8% had stage II systolic hypertension while another 1.2% had stage III systolic hypertension despite being on treatment. Four percent had stage II diastolic hypertension while another 0.3% had stage III diastolic hypertension despite being on treatment.

Table 13.6.2(a): Systolic BP, 2004 - 2006

Year	2004	2005	2006
	No. (%)	No. (%)	No. (%)
Systolic BP <120	207 (13.4)	233 (14.3)	241 (15.5)
Systolic BP <130	341 (22.0)	318 (19.4)	393 (25.2)
Systolic BP 130-139	459 (29.6)	475 (29.1)	463 (29.7)
Systolic BP 140-159	418 (27.0)	452 (27.6)	352 (22.6)
Systolic BP 160-179	102 (6.6)	133 (8.1)	91 (5.8)
Systolic BP >=180	23 (1.5)	24 (1.5)	19 (1.2)

Table 13.6.2(b): Diastolic BP, 2004 - 2006

Year	2004	2005	2006
	No. (%)	No. (%)	No. (%)
Diastolic BP<80	513 (33.1)	522 (31.9)	600 (38.5)
Diastolic BP<85	602 (38.8)	657 (40.2)	580 (37.2)
Diastolic BP 85-89	48 (3.1)	73 (4.5)	73 (4.7)
Diastolic BP 90-99	319 (20.6)	308 (18.8)	241 (15.5)
Diastolic BP 100-109	56 (3.6)	65 (4.0)	61 (3.9)
Diastolic BP ≥110	12 (0.8)	10 (0.6)	4 (0.3)

Figure 13.6.2(a): Systolic BP, 2004 - 2006

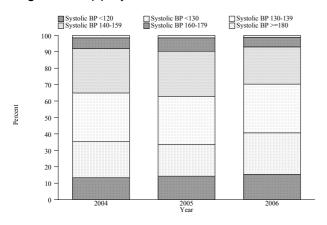


Figure 13.6.2(b): Diastolic BP, 2004 - 2006

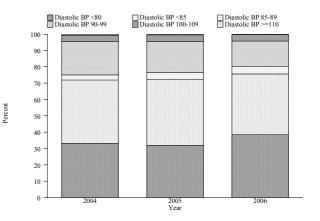
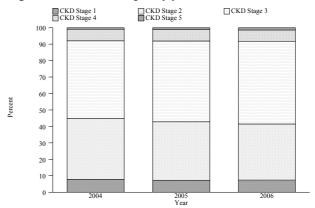


Table 13.6.3 shows classification of renal function according to CKD classification. Estimated GFR is calculated using the Cockroft and Gault equation. In 2006, 50.4% had CKD III while another 8.2% had CKD IV or V.

Table 13.6.3: CKD stages, 2004 - 2006

	2004	2005	2006
	No. (%)	No. (%)	No. (%)
CKD stage 1	120 (7.8)	118 (7.3)	114 (7.4)
CKD stage 2	571 (37.1)	578 (35.5)	528 (34.1)
CKD stage 3	727 (47.2)	800 (49.2)	779 (50.4)
CKD stage 4	109 (7.1)	112 (6.9)	105 (6.8)
CKD stage 5	13 (0.8)	19 (1.2)	21 (1.4)

Figure 13.6.3: CKD stages by year

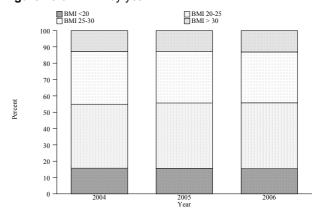


In year 2006, 31.1% were overweight while another 13.1% were obese with BMI above 30.

Table 13.6.4: BMI, 2004 - 2006

Year	2004	2005	2006	
	No. (%)	No. (%)	No. (%)	
BMI <20	243 (15.7)	255 (15.6)	241 (15.5)	
BMI 20-25	609 (39.3)	655 (40.1)	629 (40.3)	
BMI 25-30	500 (32.3)	514 (31.4)	485 (31.1)	
BMI > 30	198 (12.8)	211 (12.9)	204 (13.1)	

Figure 13.6.4: BMI by year



In year 2006, 23% had LDL cholesterol \geq 3.4 mmol/L, 58.9% had total cholesterol >5.2 while 6.5% had HDL cholesterol <1.

Table 13.6.5(a): LDL, 2004 - 2006

	2004	2005	2006			
	No. (%)	No. (%)	No. (%)			
LDL < 2.6	282 (18.2)	418 (25.6)	487 (31.2)			
LDL 2.6-3.4	944 (60.9)	862 (52.7)	713 (45.7)			
LDL >= 3.4	324 (20.9)	355 (21.7)	359 (23.0)			

Figure 13.6.5(a): LDL by year

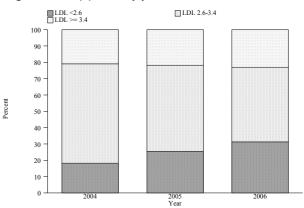


Table 13.6.5(b): Total Cholesterol, 2004 - 2006

4Year	2004	2005	2006
	No. (%)	No. (%)	No. (%)
Total Cholesterol <4.1	113 (7.3)	159 (9.7)	159 (10.2)
Total Cholesterol 4.1-5.1	413 (26.6)	455 (27.8)	482 (30.9)
Total Cholesterol 5.2-6.2	751 (48.5)	774 (47.3)	677 (43.4)
Total Cholesterol 6.3- 7.2	197 (12.7)	173 (10.6)	171 (11.0)
Total Cholesterol > 7.2	76 (4.9)	74 (4.5)	70 (4.5)

Figure 13.6.5(b): Total Cholesterol by year

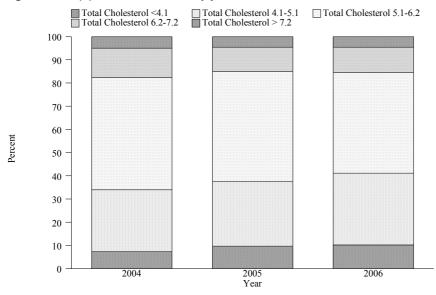
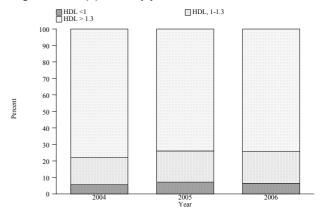


Table 13.6.5(c): HDL, 2004 - 2006

Year	2004	2005	2006	
	No. (%)	No. (%)	No. (%)	
HDL <1	87 (5.6)	118 (7.2)	101 (6.5)	
HDL 1-1.3	255 (16.5)	308 (18.8)	299 (19.2)	
HDL >1.3	1208 (77.9)	1209 (73.9)	1159 (74.3)	

Figure 13.6.5(c): HDL by year



Majority of patients were on more than one anti-hypertensive drug with 34% on 2 anti-hypertensives while 18% required 3. In 2005, despite being on treatment, a substantial number of patients had SBP \geq 160 (11%) and DBP \geq 90 (25%) (Table 13.6.6d and 13.6.6e)

Table 13.6.6(a): Treatment for hypertension, 2004 – 2006

Year	No.	% on anti- hypertensives	% on 1 anti-hypertensive drug	% on 2 anti- hypertensives	% on 3 anti- hypertensives
2004	1550	87	30	34	18
2005	1635	85	28	30	19
2006	1559	86	34	26	17

Table 13.6.6(b): Distribution of Systolic BP without anti-hypertensives, 2004 – 2006

Year	No.	Mean	SD	Median	LQ	UQ	% Patients <u>></u> 160 mmHg
2004	179	126.3	13.6	130	120	130	4
2005	229	126.9	15	130	120	137	3
2006	184	124.1	14.4	120	118.5	130	4

Table 13.6.6(c): Distribution of Diastolic BP without anti-hypertensives, 2004 – 2006

Year	No.	Mean	SD	Median	LQ	UQ	% Patients <u>></u> 90 mmHg
2004	179	78.9	9.1	80	73	80	17
2005	229	79	9	80	70	80	18
2006	184	76.6	10.2	80	70	80	11

Table 13.6.6(d): Distribution of Systolic BP on anti-hypertensives, 2004 – 2006

Year	No.	Mean	SD	Median	LQ	UQ	% Patients ≥ 160 mmHg
2004	1312	133.1	16.3	130	120	140	9
2005	1350	134.5	17.3	130	120	143	11
2006	1321	131.7	16.3	130	120	140	8

Table 13.6.6(e): Distribution of Diastolic BP on anti-hypertensives, 2004 – 2006

Year	No.	Mean	SD	Median	LQ	UQ	% Patients <u>></u> 90 mmHg
2004	1312	80.5	9.9	80	74	90	27
2005	1350	80.8	9.4	80	76	90	25
2006	1321	79.3	9.9	80	70	86	22

APPENDIX 1: DATA MANAGEMENT

Introduction

Data integrity of a register begins from the data source, data collection tools, data verification and data entry process. Registry data is never as perfect as the clinical trail data. Caution should be used when interpreting the results.

Data source

The initial phase of the data collected in the Register covered all Renal Replacement Therapy (RRT) patients in the Ministry of Health program since its inception in the early 1970s. The Register subsequently received the data from other sectors of RRT providers like the private, non-government organization (NGO), armed forces and the universities.

The Register continues to actively ascertain new RRT centres in the country. The mechanism of ascertainment is through feedback from the dialysis related company, current Source Data Provider (SDP) and public propagandas. This will gradually and eventually result in a complete RRT centre database. The identified RRT centre is invited to participate in data collection.

Participation in the National Renal Registry which was entirely voluntary prior to 2006 is now made compulsory by the Private Health Care Facilities and Services Act 1996 and its Regulations 2006 which was implemented in 1st Maly 2006. This however only applies to private and NGO centres and data submission from centres managed by the Ministry of Health, Defence or the Universities is still voluntary. RRT centres which have expressed interest in participating will be recruited as SDP.

In the year 2006, there were 38 new haemodilaysis centres. Thus, this is an average of 3 new centres per month. Within the same year 6 centres had ceased operation. This may be partly due to the Private Health Care Facilities and Services Act 1996 and its Regulations 2006. The number of RRT centres is shown in the table below. The participating rate for government centres was 100%. The data submission rate of 83.87% and 66.15% for PD and Transplant centres are due to centres reporting that they provide these services but do not have patients on regular follow-up.

	At December 2006 Known centres (N)	Agreed to Participate (N)	Submitting data in 2006 (N)	Submitting annual returns (N)	Submitted data (%)
Haemodialysis	423	415	385	332	92.77
Peritoneal Dialysis	31	31	26	26	83.87
Transplant	65	65	48	41	66.15
All modality	519	511	459	399	80.93

Data collection

The data collection tools are designed to mimic the data capture format in the patient case notes to facilitate the data transcription and minimise transcription error. All the SDPs are provided with instructions on data collection and submission to the Register.

The Register collects the RRT patients' demographic details, clinical data, dialysis treatment data, transplant data, peritonitis data and outcome data. The Register holds individual patient's identifiable data that allow complete follow-up despite patient transfers from one centre to another or change of modality which are especially common among the RRT patients. These patients are monitored and tracked through from the time they were registered and commenced their RRT treatment till their death. For those patients who were lost to follow-up, the Register will verify their final outcome with the National Vital Registration System. Patient Profiles are submitted to the Register throughout the year. The identity of patients in the database is not released publicly or in the registry reports.

Centre-specific reports are generated and forwarded to SDP on a quarterly basis. This has generated increased feedback from SDP and improved the patient ascertainment rate and the accuracy of the data transmitted to the Register.

At the end of each year, centres submit their patients' information related to dialysis and drug treatment, clinical and laboratory measurements for the year. Work related rehabilitation and Quality of life Assessment was performed for all patients during the last clinic follow-up.

The Register also conducts an annual centre survey on the staffing and facility profile. The survey questionnaire provides summary information about the number of patients on various treatments. This acts as the basis to calculate the patient ascertainment rate.

Database System

The Register initial database was created in DBASE IV in a single computer environment. It was then upgraded to Microsoft Access as a client server application. Currently the NRR data system is a Pentium Xeon 2.4 with dual processors, with a total of 1GB RAM memory and 72GB of RAID-5 (Redundant Array of Independent Disks, level 5). In view of capacity ability, performance and security issues of Microsoft Access, it was subsequently migrated to SQL Server 2000 in the year 2004.

Data management personnel

The data management personnel in the Register office are trained base on the standard operating procedures (SOP). The data entry process is also designed to enhance data quality. Quality assurance procedures are in place at all stages to ensure the quality of data.

Visual review, Data entry and de-duplication verification, Data Editing

On receiving the case report form (CRF) submitted by SDP, visual review is performed to check for obvious error or missing data in the compulsory fields. Data entry will not be performed if a critical variable on the CRF is missing or ambiguous. The CRF is returned to the SDP for verification.

After passing the duplicate check, the data is than entered and coded where required. Edit checks are performed against pre-specified validation rules to detect missing values, out of range values or inconsistent values. Any data discrepancy found is verified against the source CRF and resolved within the Register office where possible. Otherwise the specific data query report will be generated and forwarded to the SDP to clarify and resolve the data discrepancy.

Data coding, data cleaning / data analysis

Most of the data fields have auto data coding. Those data in text fields will be manually coded by the Register manager. A final edit check run is performed to ensure that data is clean. All queries are resolved before dataset is locked and exported to the statistician for analysis

Limitation:

NRR data submission is still paper base. The majority of the RRT centres do not have electronic patient information system. Computer literacy among staff is still low.

The data submission to the Register is still mainly on voluntary basis using the standard data collection tools. Some SDP choose not to participate in data collection on the patient treatment data for various reasons. We sincerely hope with the enforcement of the Private Health Care Facilities and Services Act 1996 and its Regulations 2006 which was implemented in 1st May 2006, participation rate from private and NGO centres shall improve in the coming years.

Data release and publication policy

One of the primary objectives of the Registry is to make data available to the renal community. There are published data in the registry's annual report in the website: http://www.msn.org.my/nrr. This report is copyrighted. However it may be freely reproduced without the permission of the National Renal Registry. Acknowledgment would be appreciated. Suggested citation is: YN Lim, TO Lim (Eds). Fourteenth Report of the Malaysian Dialysis and Transplant Registry 2006. Kuala Lumpur 2007

A distinction is made between use of NRR results (as presented in NRR published report) and use of NRR data in a publication. The former is ordinary citation of published work. NRR, of course encourages such citation whether in the form of presentation or other write-ups. The latter constitutes original research publication. NRR position is as follows:

- The NRR does not envisage independent individual publication based entirely on NRR published results, without further analyses or additional data collection.
- NRR however agrees that investigator shall have the right to publish any information or material arising in part out of NRR work. In other words, there must be additional original contribution by the investigator in the work intended for publication.
- NRR encourages the use of its data for research purpose. Any proposed publication or presentation
 (e.g. manuscript, abstract or poster) for submission to journal or scientific meeting that is based in part
 or entirely on NRR data should be sent to the NRR prior to submission. NRR will undertake to
 comment on such documents within 4 weeks. Acknowledgement of the source of the data would also
 be appreciated.
- Any formal publication of a research based in part or entirely on NRR data in which the input of NRR
 exceeded that of conventional data management and provision will be considered as a joint
 publication by investigator and the appropriate NRR personnel.

Participating centre is now able to down load own centre's data from the secured web-site from link from www.msn.org.my/nrr. Any party who wish to request data for a specific purpose that requires computerrun should make such requests in writing (by e-mail, fax, or classic mail) accompanied by a Data Release Application Form and signed Data Release Agreement Form. Such request will require approval by the Advisory Board before the data can be released.

Distribution of report

The Malaysian Society of Nephrology has made a grant towards the cost of running the registry and the report printing to allow distribution to all members of the association and the source data producers. The report will also be distributed to relevant Health Authorities and international registries.

Further copies of the report can be made available with donation of RM60.00 to defray the cost of printing. The full report is also available in the registry web *site* <u>www.msn.org.my/nrr</u>.

APPENDIX II: ANALYSIS SETS, STATISTICAL METHODS AND DEFINITIONS

ANALYSIS SETS

This refers to the sets of cases whose data are to be included in the analysis. Four analysis sets were defined:

1. Dialysis patients notification between 1997 and 2006

This analysis set consists of patients commencing dialysis between 1997 and 2006. This analysis set was used for the analysis in Chapter 1, 2 and 3. Sections 2.3.3 & 2.3.4 were tabulated based on Dialysis patients' notification and Annual return.

Patients with age commencing dialysis less than 20 years old between 1997 and 2006 was used for the analysis in Chapter 5.

Since 1993, the NRR conducted an annual survey on all dialysis patients to collect data on dialysis and drug treatment, clinical and laboratory measurements. All available data were used to describe the trends in these characteristics. However, in the early years, the data collected from annual survey were relatively incomplete. Hence, for any analysis in relation to these characteristics, we used only data from 1997 onwards when the data were more complete. Remaining missing data in this analysis set was imputed using first available observation carried backward or last observation carried forward. This analysis set was used for the analysis in Chapters 6 to 12.

2. Rehabilitation outcomes

Chapter 4 analysis set was confined to the relevant population. Hence we exclude the following groups.

- i. Age less than or equal to 21 years
- ii. Age more than or equal to 55 years
- iii. Homemaker
- iv. Full time student
- v. Retired

3. Centre Survey data

Section 2.2 in the report was based on annual centre survey data between 1997 to 2006 rather than individual patient data reported to the Registry.

4.Peritonitis data

Analysis was confined to CAPD patients who were on peritoneal dialysis from 31st Dec 1999. This analysis set was used for the analysis in Section 12.4.

STATISTICAL METHODS

Population treatment rates (new treatment or prevalence rates)

Treatment rate is calculated by the ratio of the count of number of new patients or prevalent patients in a given year to the mid-year population of Malaysia in that year, and expressed in per million-population. Results on distribution of treatment rates by state are also expressed in per million-population since states obviously vary in their population sizes.

Death rate calculation

Annual death rates were calculated by dividing the number of deaths in a year by the estimated mid-year patient population.

Odds ratio

The odds of an event is the probability of having the event divided by the probability of not having it. The odds ratio is used for comparing the odds of 2 groups. If the odds in group 1 is O1 and group 2 is O2, then odds ratio is O1/O2. Thus the odds ratio expresses the relative probability that an event will occur when 2 groups are compared.

With multiple factors, logistic regression model was used to estimate the independent effect of each factor, expressed as odds ratio, on the event of interest.

Risk ratio

The relative measure of the difference in risk between the exposed and unexposed populations in a cohort study. The relative risk is defined as the rate of disease among the exposed divided by the rate of the disease among the unexposed. A relative risk of 2 means that the exposed group has twice the disease risk as the unexposed group.

Multinomial logistic regression model was used to estimate the independent effect of each factor, expressed as risk ratio, on the event of interest. Sero-converters are defined as patient with anti-HCV negative at his first dialysis year convert to anti-HCV positive in the subsequent year.

Survival analysis

The unadjusted survival probabilities were calculated using the Kaplan-Meier method, in which the probability of surviving more than a given time can be estimated for members of a cohort of patients without accounting for the characteristics of the members of that cohort.

In order to estimate the difference in survival of different subgroups of patients within the cohort, a stratified proportional hazards model (Cox) was used where appropriate. The results from Cox model are interpreted using a hazard ratio. Adjusted survival probabilities are with age, gender, primary diagnosis and time on RRT used as adjusting risk factors. For diabetics compared with non-diabetics, for example, the hazard ratio is the ratio of the estimated hazards for diabetics relative to non-diabetics, where the hazard is the risk of dying at time t given that the individual has survival until this time. The underlying assumption of a proportional hazards model is that the ratio remains constant throughout the period under consideration.

Technique failure is defined as occurrence of death or transfer to another modality of dialysis. Similarly, graft failure is defined as occurrence of death or returned to dialysis.

The analysis is confined to all dialysis patients regardless they switched modalities. For example, if the patient is on CAPD for the first 2 years and then switches to HD, the survival duration on HD is calculated as if he is a new patient on HD. In addition, the calculation of survival on CAPD of that patient is censured as if he has been lost to follow-up.

Analysis of trend of intermediate results

For summarizing intermediate results like continuous laboratory data, we have calculated summary statistics like mean, standard deviation, median, lower quartile, upper quartile and the cumulative frequency distribution graph is plotted over year. Cumulative distribution plot shows a listing of the sample values of a variable on the X axis and the proportion of the observations less than or greater than each value on the Y axis. An accompanying table gives the Median (50% of values are above or below it), upper quartile (UQ, 25% of values above and 75% below it), lower quartile (LQ, 75% of values above and 25% below it). Other percentiles can be read directly off the cumulative distribution plot. The table also shows percent of observations above or below a target value, or with an interval of values; the target value or interval obviously vary with the type of laboratory data. For example, interval of values for prescribed KT/V is >1.3 and that for haemoglobin is <10, 10-11 and >11 g/l. The choice of target value is guided by published clinical practice guidelines, for example, the DOQI guideline; or otherwise they represent consensus of the local dialysis community.

Centre survey data

In contrast to other results reported in this report, Section 2.2 was based on centre survey data rather than individual patient data reported to the Registry. This is to provide up-to-date information on patient and centre census in the country and thus overcome the inevitable time lag between processing individual patient data and subsequent reporting of results. The survey was conducted in the month of December 2006. Centre response rate to survey was 100%. Standard error estimates are not reported because no sample was taken. Results on distribution by state are also expressed in per million-population since states obviously vary in their population sizes. State population data are based on 2006 census projection. It is very difficult to estimate the amount of cross boundary patient flow; this source of error is therefore not accounted for in computing states estimates. However, we minimize the bias by combining states (Selangor and Wilayah Persekutuan, Kedah and Perlis) based on geographical considerations. HD treatment capacity is derived by assuming on average patients underwent 3 HD sessions per week and a centre can maximally operate 2.5 shifts per day. A single HD machine can therefore support 5 patients' treatment. Obviously HD treatment capacity is calculated only for centre HD. The ratio of the number of centre HD capacity to number of centre HD patient is a useful measure of utilization of available capacity.

Centre variation

To compare the variation of the intermediate results between centres, graph describing intermediate results in each centre are presented. The 95% confidence intervals have been calculated using the normal approximation of the Poisson to show the variation of proportion in centres. Lower quartile and upper quartile are instead plotted in comparison of variation in median among centres. In the analysis, centres with less than ten patients were combined in a pooled centre. An accompanying table gives the summary statistics like minimum, 5th percentile, lower quartile, median, upper quartile, 95th percentile and maximum value among centres over year.

Centres with intermediate results for <10 patients were combined into one composite centre.

Peritonitis rate

The occurrence of peritonitis is expressed as number of episode per patient-month of observation; peritonitis rate in short. Relapse peritonitis is defined as peritonitis caused by the same organism occurring within 6 weeks of diagnosis of previous peritonitis.

Funnel plot

Analysis confined to new dialysis patients from year 1997-2002. The figure is included to assess whether survival probability adjusted to age 60 and diabetes of each centre is likely to be different from the national average. Centres with patients less 10 will be excluded from the analysis.

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