

# Clinical Practice Guidelines

MOH/P/PAK/279.14 (GU)

## Management of Autism Spectrum Disorder in Children and Adolescents



Ministry of Health Malaysia



Malaysian Psychiatric Association



Malaysian Child and Adolescent Psychiatry Association



Academy of Medicine Malaysia

## Published by:

Malaysian Health Technology Assessment Section (MaHTAS)  
Medical Development Division, Ministry of Health Malaysia  
Level 4, Block E1, Precinct 1  
Federal Government Administrative Centre  
62590, Putrajaya, Malaysia

## Copyright

The copyright owner of this publication is MaHTAS. The content may be reproduced in any number of copies and in any format or medium provided that a copyright acknowledgement to MaHTAS is included. The content is not to be changed, sold, used to promote or endorse any product or service, nor used in an inappropriate or misleading context.

ISBN: 978-967-0769-01-1

Available on the following websites:

<http://www.moh.gov.my>

<http://www.acadmed.org.my>

<http://www.psychiatry-malaysia.org>

## STATEMENT OF INTENT

These clinical practice guidelines (CPG) are designed to be a guide for clinical practice, based on the best available evidence at the time of development. Adherence to these guidelines may not necessarily guarantee the best outcome in every case. Every healthcare provider is responsible for the management of his / her unique patient based on the clinical picture presented by the patient and the management options available locally.

These guidelines were issued in July 2014 and will be reviewed in 2018 or earlier if new evidence becomes available.

## TABLE OF CONTENTS

No.	Title	Page
	Levels of Evidence and Grades of Recommendation	iii
	Guidelines Development and Objectives	iv
	Guidelines Development Group	vii
	Review Committee	ix
	External Reviewers	xi
	Algorithm: Management of Children with Autism Spectrum Disorder	xiii
1	Introduction	1
2	Risk Factors	1
3	Screening	3
4	Assessment and Diagnosis	5
5	Comorbidities and Other Difficulties	8
6	Investigations	10
	a. Audiological Evaluation	10
	b. Other Investigations	10
7	Treatment	11
	7.1 Non-Pharmacological Treatment	12
	a Applied Behaviour Analysis	12
	b Speech, Language and Communication Intervention	13
	c Occupational Therapy	15
	d Other Interventions / Programmes	16
	e Parental Education and Support	17
	f Cognitive Behaviour Therapy	17
	g Treatment and Education of Autistic and Related Communication Handicapped Children	18
	h Diet	18
	7.2 Pharmacotherapy	19
	a Atypical antipsychotics	19
	b Antidepressants	20
	c Other medications	21

<b>No.</b>	<b>Title</b>	<b>Page</b>
	7.3 Traditional and Complementary Medicine	22
8	Social Welfare Services	23
9	Monitoring and Preparation of Transition to Adult Services	23
10	Implementing the Guidelines	23
	References	26
	Appendices	
	Appendix 1 : Examples of Search Strategy	33
	Appendix 2 : Clinical Questions	34
	Appendix 3 : Modified Checklist for Autism in Toddlers (M-CHAT)	35
	Appendix 4 : Diagnostic Criteria for Autism Disorder in the Diagnostic and Statistical Manual of Mental Disorders, Fourth Edition, Text Revision (DSM-IV-TR)	40
	Appendix 5 : Diagnostic Criteria for Autism Spectrum Disorder in the Diagnostic and Statistical Manual of Mental Disorders, Fifth Edition (DSM-5)	42
	Appendix 6 : International Classification of Disease (ICD) 10	46
	Appendix 7 :	
	• Signs and Symptoms of Possible Autism in Preschool Children (or Equivalent Mental Age)	49
	• Signs and Symptoms of possible Autism in Primary School Children (Aged 5–11 Years or Equivalent Mental Age)	51
	• Signs and Symptoms of Possible Autism in Secondary School Children (Older Than 11 Years or Equivalent Mental Age)	53
	Appendix 8 : Diagnostic Tools	55
	List of Abbreviations	58
	Acknowledgement	60
	Disclosure Statement	60
	Source of Funding	60

## Levels of Evidence

Level	Study design
I	Evidence from at least one properly randomised controlled trial (RCT)
II-1	Evidence obtained from well-designed controlled trials without randomisation
II-2	Evidence obtained from well-designed cohort or case-control analytic studies, preferably from more than one centre or group
II-3	Evidence from multiple time series with or without the intervention. Dramatic results in uncontrolled experiments (such as the results of the introduction of penicillin treatment in the 1940s) could also be regarded as this type of evidence
III	Opinions of respected authorities based on clinical experience; descriptive studies and case reports; or reports of expert committees

**Source: US / Canadian Preventive Services Task Force 2001**

## Grades of Recommendation

A	At least one meta-analysis, systematic review, or RCT, or evidence rated as good and directly applicable to the target population
B	Evidence from well conducted clinical trials, directly applicable to the target population, and demonstrating overall consistency of results; or evidence extrapolated from meta-analysis, systematic review, or RCT
C	Evidence from expert committee reports, or opinions and /or clinical experiences of respected authorities; indicates absence of directly applicable clinical studies of good quality

**Source: Modified From The Scottish Intercollegiate Guidelines Network (SIGN)**

Note: The grades of recommendation relate to the strength of the evidence on which the recommendation is based. It does not reflect the clinical importance of the recommendation.

## Guidelines Development and Objectives

### Guidelines Development

The members of the Development Group (DG) for this Clinical Practice Guidelines (CPG) are staff of the Ministry of Health (MOH) and Ministry of Higher Education Malaysia. A multidisciplinary review committee (RC) was actively involved during this CPG development process.

A systematic literature search was carried out using the following databases: Guidelines International Network (G-I-N), Medline via Ovid, PUBMED, Cochrane Database of Systematic Reviews (CDSR) and International Health Technology Assessment websites. A search strategy to cover all aspects on management of autism spectrum disorders (ASD) was developed in the Medline database and adapted to other databases. Search strategies were a combination of MeSH and keyword searches including abbreviations (refer to Appendix 1 on an example of Search Strategy). The search was restricted to human studies, literature published in the English language and the last ten years. If the evidence was insufficient, the period of publication was extended for another ten years. In addition, the reference lists of all retrieved literature and guidelines were searched to further identify relevant studies. All searches were conducted from May 2012 till November 2013. Literature searches were repeated for all clinical questions at the end of the CPG development process. The aim was to identify and include any further relevant papers published before 28 February 2014. Future CPG updates will consider evidence published after this cut-off date. The details of the search strategy can be obtained upon request from the CPG Secretariat.

Reference was also made to other CPGs on ASD such as i) Guidelines on Treatment, Diagnosis and Clinical Intervention for Children and Young People with Autism Spectrum Disorder (SIGN 98-2007), ii) AMS-MOH Clinical Practice Guidelines 1/2010 - Autism Spectrum Disorders in Pre-school Children, iii) Autism Spectrum Disorder by the Ministry of Health New Zealand 2008, and iv) Autism diagnosis in children and young people: Recognition, referral and diagnosis of children and young people on the autism spectrum (NICE, 2011). These CPGs were evaluated using the Appraisal of Guidelines for Research and Evaluation (AGREE) II prior to them being used as references.

A total of 27 clinical questions were developed under different sections. Members of the DG were assigned individual questions within these sections. (Refer to Appendix 2) The DG members met 23 times throughout the development of these guidelines. All literatures retrieved were appraised by at least two DG members using Critical Appraisal Skill Programme checklist,

presented in evidence tables and further discussed in each DG meetings. All statements and recommendations formulated after that were agreed upon by both the DG and RC. Where evidence was insufficient, the recommendations were made by consensus of the DG and RC. These CPGs are based largely on the findings of systematic reviews, meta-analyses and clinical trials, with local practices taken into consideration.

The evidence used in these guidelines was graded using the US / Canadian Preventive Services Task Force Level of Evidence (2001), while the grading of recommendation was modified from grades of recommendation of the Scottish Intercollegiate Guidelines Network.

On completion, the draft was sent for review by external reviewers. It was also posted on the MOH Malaysia official website for feedback by any interested parties. The draft was finally presented to the Technical Advisory Committee for CPG, the HTA and CPG Council of the MOH Malaysia for review and approval.

## Objectives

The aim of this CPG is to provide evidence-based recommendations in the management of ASD, particularly on the detection, assessment and intervention of the condition in children and adolescents.

## Clinical Questions

Refer to Appendix 2

## Target Population

Children and adolescents with ASD who are less than 18 years of age

## Target Group / User

This document is intended to guide health professionals and relevant stakeholders in primary and secondary / tertiary care of autism including:

- i Primary Care Providers (Family Medicine Specialists / Medical Officers / General Practitioners)
- ii Psychiatrists / Child and Adolescent Psychiatrists
- iii Paediatricians
- iv Psychologists
- v Audiologists
- vi Speech-Language Therapists
- vii Occupational Therapists
- viii Social Workers
- ix Nurses
- x Pharmacists
- xi Policy Makers / Programme Managers
- xii Educators

## Healthcare Settings

Outpatient, inpatient and community settings

## Guidelines Development Group

### Chairperson

Dr. Farahidah Md Dai  
Head of Department and  
Consultant Child & Adolescent Psychiatrist  
Hospital Sultanah Aminah, Johor Bahru  
Johor

### Members (Alphabetical Order)

Ms. Ang Wei Nei  
Pharmacist  
Hospital Selayang  
Selangor

Ms. Masrita Mohd Rosli  
Occupational Therapist  
Hospital Selayang  
Selangor

Dr. Azah Abdul Samad  
Family Medicine Specialist  
Klinik Kesihatan Tanglin  
Kuala Lumpur

Dr. Mohd Aminuddin Mohd Yusof  
Head of Clinical Practice Guidelines Unit  
Health Technology Assessment Section  
Medical Development Division, MOH  
Putrajaya

Ms. Ee Su Im  
Occupational Therapist  
Hospital Kuala Lumpur  
Kuala Lumpur

Ms. Nor Asmawati Mohamad Ali Abdul Rahman  
Medical Social Worker  
Hospital Umum Sarawak  
Sarawak

Dr. Eni Rahaiza Muhd Ramli  
Consultant Child &  
Adolescent Psychiatrist  
Hospital Taiping  
Perak

Dr. Norharlina Bahar  
Child & Adolescent Psychiatrist  
Hospital Selayang  
Selangor

Dr. Juriza Ismail  
Senior Lecturer and  
Consultant Developmental Paediatrician  
Universiti Kebangsaan Malaysia  
Medical Centre  
Kuala Lumpur

Dr. Norhayati Nordin  
Hospital Director and  
Consultant Child & Adolescent Psychiatrist  
Hospital Mesra, Bukit Padang  
Sabah

Dr. Manveen Kaur Harbajan Singh  
Senior Lecturer and Psychiatrist  
University of Malaya  
Kuala Lumpur

Dr. Nurdiana Abdullah  
Senior Lecturer and  
Primary Care Physician  
University of Malaya  
Kuala Lumpur

Ms. Nurshahira Razali  
Speech-Language Therapist  
Hospital Kuala Lumpur  
Kuala Lumpur

Datin Dr. Sheila Marimuthu  
Consultant Paediatrician and  
Adolescent Specialist  
Hospital Kuala Lumpur  
Kuala Lumpur

Dr. V. Paranthaman  
Family Medicine Specialist  
Klinik Kesihatan Jelapang  
Perak

Dr. Selvasingam Ratnasingam  
Consultant Child & Adolescent Psychiatrist  
Hospital Umum Sarawak  
Sarawak

Dr. Ramli Mohd Ali  
Consultant Child & Adolescent Psychiatrist  
Hospital Kuala Lumpur  
Kuala Lumpur

Ms. Sin Lian Thye  
Nursing Officer  
(Coordinator / Information Specialist)  
Health Technology Assessment Section  
Medical Development Division, MOH  
Putrajaya

Dr. Ranjini S. Sivanesom  
Consultant Developmental Paediatrician  
Hospital Kuala Lumpur  
Kuala Lumpur

Ms. Siti Suriani Che Hussin  
Audiologist  
Hospital Kuala Lumpur  
Kuala Lumpur

Ms. Rozila Sumardi  
Speech-Language Therapist  
Hospital Sungai Buloh  
Selangor

Dr. Toh Teck Hock  
Paediatrician and  
Head of Clinical Research Centre  
Hospital Sibul  
Sarawak

Ms. See Geok Lan  
Clinical Psychologist  
Hospital Rehabilitation Cheras  
Kuala Lumpur

Dr. Yusni Yusuff  
Consultant Child & Adolescent Psychiatrist  
Hospital Sultan Abdul Halim  
Kedah

## Review Committee

The draft of these guidelines was reviewed by a panel of experts from both public and private sectors. They were asked to comment primarily on the comprehensiveness and accuracy of the interpretation of evidence supporting the recommendations in the guidelines.

## Chairperson

Dr. Toh Chin Lee  
Head of Department and  
Senior Consultant Child & Adolescent Psychiatrist  
Department of Psychiatry & Mental Health  
Hospital Selayang  
Selangor

## Members (Alphabetical Order)

Associate Professor Dr. Aili Hanim Hashim  
Lecturer and  
Consultant Child & Adolescent Psychiatrist  
Department of Psychological Medicine  
Faculty of Medicine, University of Malaya  
Kuala Lumpur

Dr. Baizurah Mohd Hussain  
Head of Department Pathology (Patient Advocate)  
Department of Pathology  
Hospital Ampang  
Selangor

YBhg. Dato' Dr. Amar-Singh HSS  
Head of Department and  
Senior Consultant Paediatrician  
Hospital Raja Permaisuri Bainun  
Perak

Dr. Leethuman Ramanathan  
Head of Department and  
Consultant Physician (Patient Advocate)  
Department of Medicine  
Hospital Taiping  
Perak

Dr. Rajini Sarvanathan  
Consultant Developmental Paediatrician  
Sunway Medical Centre  
Selangor

Dr. Rekha Balachandran  
Consultant Otorhinolaryngologist (Patient Advocate)  
Hospital Raja Permaisuri Bainun  
Perak

Dr. Wong Woan Yiing  
Secretary, National Early Childhood Intervention Council and  
Consultant Paediatrician  
KPJ Ipoh Specialist Hospital  
Perak

## External Reviewers

The following external reviewers provided feedback on the draft:-

Dr. Andy Shih  
Senior Vice President of Scientific Affairs  
Autism Speaks  
United States of America

Ms. Amelia Inbam Neelagandan  
Speech-Language Therapist  
Hospital Queen Elizabeth  
Sabah, Malaysia

Professor Dr. Frances Page Glascoe  
Professor of Paediatrics  
Vanderbilt University  
United States of America

Associate Professor Dr. Jill Sewell  
Deputy Director  
Centre for Community Child Health  
Royal Children's Hospital  
Melbourne, Australia

Professor Dr. Mohd Jamil Yaacob  
Consultant Child & Adolescent Psychiatrist  
AIMST University  
Kedah, Malaysia

Clinical Associate Professor Dr. John Wray  
State Child Development Centre  
West Perth, Australia

Dr. Lai Fong Hwa  
Consultant Child & Adolescent Psychiatrist  
Hospital Pulau Pinang  
Pulau Pinang, Malaysia

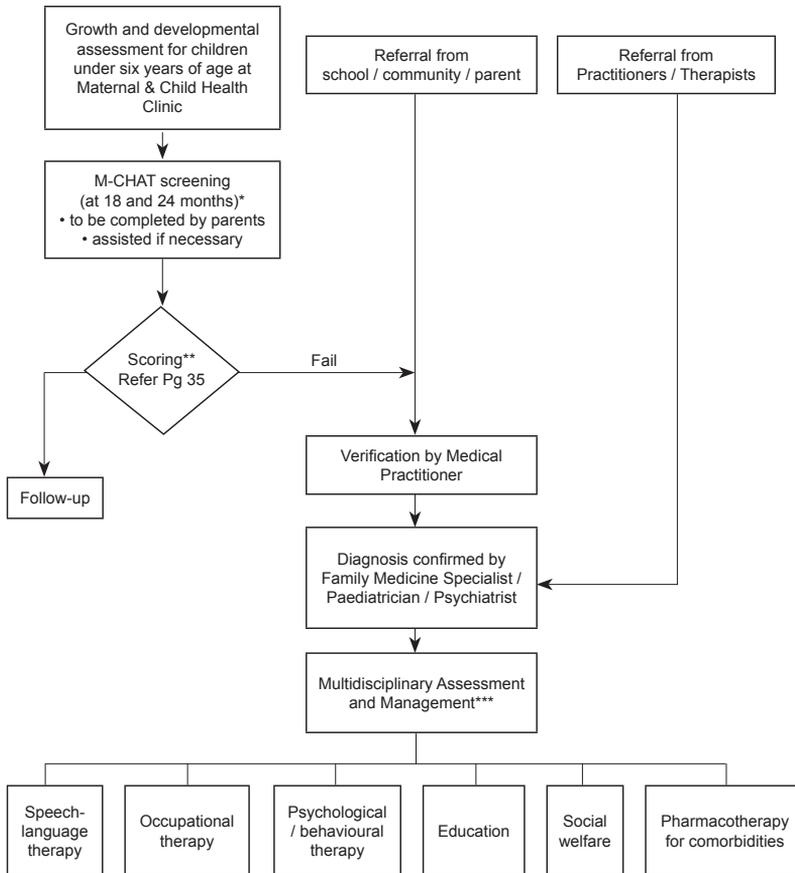
Ms. Masne Kadar  
Occupational Therapist and Lecturer  
Universiti Kebangsaan Malaysia  
Kuala Lumpur, Malaysia

Dr. Nazrila Hairizan Nasir  
Family Medicine Specialist  
Klinik Kesihatan Pandamaran  
Selangor, Malaysia

Ms. Nor Azizah Nurdin  
Freelance Speech-Language Therapist  
Selangor, Malaysia

Professor Dr. Wong Swee Lan  
Department of Population Medicine  
Faculty of Medicine and Health Sciences  
University Tunku Abdul Rahman  
Kuala Lumpur, Malaysia

## Algorithm on Management of Children with Autism Spectrum Disorder



**Early intervention programme (EIP) is strongly advocated**

\* M-CHAT may be used to screen children up until 30 months of age if the child misses the earlier screening

\*\*Regardless of the screening result, children suspected of ASD by the family or other care provider should be referred for evaluation

\*\*\*Multidisciplinary Assessment and Management Team may include:

- Family Medicine Specialist
- Paediatrician
- Psychiatrist
- Clinical Psychologist
- Occupational Therapist
- Speech-Language Therapist
- Audiologist
- Medical Social Worker

## 1. Introduction

Autism Spectrum Disorder (ASD) is a neurodevelopmental disorder characterised by impairments in communication, behaviour and social functioning beginning in childhood. There is no local epidemiological study on ASD prevalence in Malaysia. However, in a feasibility study on the use of Modified Checklist for Autism in Toddlers (M-CHAT) among children of 18 to 36 months of age in child health clinics by Ministry of Health Malaysia, the prevalence of ASD in Malaysia was approximately 1.6 in 1,000.<sup>1</sup> In the USA, the overall prevalence of ASD is 14.7 per 1,000 (one in 68) children aged 8 years and the prevalence shows an increasing trend over the years.<sup>2</sup>

Clinicians have been seeing an increasing number of children with speech delay and social communication difficulties who require further assessment. Children with speech delay and behavioural issues are frequently placed in special education classes without an accurate diagnosis. While waiting for diagnosis to be established, these children should be referred for the relevant interventions.

Early diagnosis and prompt intervention of children with ASD is crucial for the best outcome. The importance of early recognising and initiation of early referral to optimise the child's potential must be emphasised.

This has prompted the need to produce a CPG for the wide usage of healthcare providers in Malaysia as well as to provide assistance to parents of children with ASD. It is hoped that this CPG can give a clear picture to primary care providers to direct these children in an efficient manner. Referral pathways to specialists and various interventionists have been outlined in this CPG in order to facilitate this process.

## 2. Risk Factors

The aetiology of ASD is unclear. It is multi-factorial which includes both genetic vulnerability and environmental factors. Only risk factors deemed important by the DG and RC are discussed here.

### a. Advancing Parental Age

Parental age is among the most consistently studied risk factor and the risk for ASD increases with parental age.

- Maternal age:
  - >35 years old vs 25 - 29 years old (OR = 1.31, 95% CI 1.19 to 1.45)<sup>3, level II-2</sup>
  - >40 years old vs <30 years old (OR = 2.1, 95% CI 1.48 to 2.86)<sup>4, level II-2</sup>

- Paternal age: compared to  $\leq 29$  years old,<sup>5, level II-2</sup>
  - 49 years old (OR = 1.42, 95% CI 1.07 to 1.87)
  - $\geq 50$  years old (OR = 2.21, 95% CI 1.26 to 3.88)
  - $\geq 55$  years old (OR = 4.36, 95% CI 2.09 to 9.09)
- First born of mother aged  $>35$  years old and father aged  $>40$  years old vs parents aged 25 - 29 years old, adjusted OR = 3.1, 95% CI 2.0 to 4.7.6,<sup>level II-2</sup>

**b. Prematurity** ( $<37$  weeks of gestation) in particular those born  $<33$  weeks are at risk of developing ASD (OR = 5.4, 95% CI 1.1 to 27.7).<sup>4, level II-2</sup>

**c. Neonatal encephalopathy** has OR for ASD ranging from 3.06 to 5.59.<sup>4, level II-2</sup>

#### **d. Genetic Risk**

The adjusted relative recurrence risk of ASD is increased with increasing genetic relatedness:<sup>3, level II-2</sup>

- Monozygotic twins = 153.0 (95% CI 56.7 to 412.8)
- Dizygotic twins = 8.2 (95% CI 3.7 to 18.1)
- Full siblings = 10.3 (95% CI 9.4 to 11.3)
- Maternal half-siblings = 3.3 (95% CI 2.6 to 4.2)
- Paternal half-siblings = 2.9 (95% CI 2.2 to 3.7)
- Cousins = 2.0 (95% CI 1.8 to 2.2)

The RR of ASD for infants with multiple older affected siblings is significantly 2.2 times higher than those who had only one older affected sibling.<sup>7, level II-2</sup>

#### **e. Peri-conceptual supplement**

Use of folic acid supplement in mothers around the time of conception (four weeks before and eight weeks after pregnancy)<sup>8, level II-2</sup> or peri-conceptual prenatal vitamin intake (containing more iron, vitamin B6, vitamin B12 and 800  $\mu\text{g}$  of folic acid as compared to ordinary multivitamins)<sup>9, level II-2</sup> appears to offer some benefit especially in those who are genetically susceptible.

#### **f. Immunisation (Vaccination)**

Based on a Cochrane systematic review of 10 studies, no significant association was found between MMR immunisation and autism.<sup>10, level II-2</sup> In another study, increasing exposure to antibody-stimulating proteins and polysaccharides in vaccines during the first two years of life was not associated with risk of developing ASD.<sup>11, level II-2</sup>

Immunisation is not associated with the development of autism spectrum disorder.

### RECOMMENDATION 1

- Screening for autism spectrum disorder (ASD) should be emphasised in children with the following high risk factors:-
  - Increased parental age
    - Maternal age >40 years old
    - Paternal age >50 years old
    - First born of mother aged >35 years old and father aged >40 years old
  - Prematurity of <33 weeks of gestation
  - Neonatal encephalopathy
  - Family history of ASD

(Grade C)

## 3. Screening

Screening for ASD is important as early detection enables early intervention and better outcomes. There are limited studies on the effectiveness of screening tools for ASD in children. Three systematic reviews (SRs) found that Checklist for Autism in Toddler (CHAT), Modified Checklist for Autism in Toddlers (M-CHAT) and Social Communication Questionnaire (SCQ) performed better in the screening of young children for ASD. The primary studies were inadequate with small sample sizes, lack of blinding between screening and diagnosis, and largely did not follow up children with negative results.<sup>12, level II-1; 13, level II-1; 14, level II-1</sup>

It is important to be aware that false positive or false negative results from any use of screening tests may cause unnecessary parental anxiety or delay correct diagnosis. The ultimate decision about the need for referral and further assessment should be made on clinical grounds.

### a. Modified Checklist for Autism in Toddlers (M-CHAT)

Modified Checklist for Autism in Toddlers (M-CHAT) [Appendix 3] is a 23-item questionnaire on child behaviour and development reported by parents for children aged 18 and 24 months of age.<sup>15, level III</sup> It has been translated into Malay and Chinese languages for local use in healthcare facilities in Malaysia. Training is required for interpretation of the result.

The specificity of M-CHAT was reported as 98% and follow-up of a subset of children at age 3.5 years resulted in a sensitivity of 100%.<sup>14, level II-1</sup> In a recent SR, the sensitivity, specificity and PPV were found to be 70% to 92%, 27% to 43% and 5.8% to 76% respectively. M-CHAT was also better at detecting

autism in children aged 24 months versus (vs) 18 months and those in a high risk group in early intervention programme centres vs a low risk group in the routine baby clinic. M-CHAT is recommended for use at 18 months to assist with early identification of ASD, and 24 months, to identify those toddlers who have regression.<sup>12, level II-2</sup>

## **b. Social Communication Questionnaire (formerly known as Autism Screening Questionnaire)**

Social Communication Questionnaire (SCQ) is a parent-rated questionnaire on children aged above four years. It evaluates the social interaction, communication, language and stereotypic behaviours for possible autism or other ASD.<sup>16, level III</sup>

SCQ was better in detecting ASD in individuals over seven years of age (sensitivity of 86% to 90% and specificity of 78% to 86%) compared to children aged 2 to 3 years old (sensitivity 47% to 54% and specificity 89% to 92%).<sup>13, level II-1</sup>

## **c. Other Screening Tools**

Other screening tools used are:

### **i) General / broadband developmental screening tools:**

- Child Behaviour Checklist (CBCL)<sup>13, level II-1; 17, level III; 18, level III</sup>
- Infant-Toddler Checklist (ITC)<sup>19, level III</sup>
- Parents' Evaluation of Developmental Status (PEDS)<sup>20, level II-2</sup>

### **ii) Screening tools specific for ASD:**

- Checklist for Autism in Toddlers (CHAT)<sup>12, level II-1; 14, level II-1; 21, level II-2</sup>
- Checklist for Autism in Toddlers for Chinese Children (CHAT-23)<sup>13, level II-1; 17, level III; 18, level III</sup>
- Modified Checklist for Autism in Toddlers, Revised with Follow-up<sup>22, level III</sup>
- Gilliam Autism Rating Scale / Gilliam Autism Rating Scale Second Edition (GARS / GARS-2)<sup>13, level II-1; 17, level III; 18, level III</sup>
- Social Responsiveness Scale (SRS)<sup>13, level II-1; 17, level III; 18, level III</sup>
- Autism Spectrum Screening Questionnaire (ASSQ)<sup>13, level II-1; 17, level III; 18, level III</sup>
- Asperger Syndrome Diagnostic Scale (ASDS)<sup>13, level II-1; 17, level III; 18, level III</sup>

## RECOMMENDATION 2

- Modified Checklist for Autism in Toddlers (M-CHAT) may be used as a screening tool for autism spectrum disorder (ASD) among children of 18 months and repeat at 24 months if the child passes the earlier M-CHAT. **(Grade C)**
- M-CHAT may be used to screen children up till the age of 30 months if the child misses the earlier screening. **(Grade C)**
- Regardless of the screening result, children suspected of ASD at any age by the family or other care providers should be referred for evaluation. **(Grade C)**

## 4. Assessment and Diagnosis

The diagnosis of ASD is made either by using criteria from the Diagnostic and Statistical Manual of Mental Disorders [DSM-IV-TR, 2000 (Appendix 4) and DSM-5, 2013 (Appendix 5)] or 10th-Revision of International Classification of Diseases [ICD-10] (Appendix 6).

### 4.1 Initial Assessment – History, Developmental History and Physical Examination

Children with ASD may show early symptoms manifested by behavioural abnormalities. These include poor eye contact, impairment in visual tracking of an object, atypical response to name, less social smiling and delayed expressive and receptive language. The possibility of ASD should be considered if there are concerns about development or behaviour.<sup>23</sup> Repetitive behaviour, restricted interests and limited ability to imitate are also early symptoms of ASD.

Health care providers should take into consideration cultural and socioeconomic factors that may affect assessment.<sup>24, level III</sup>

#### a. Signs and symptoms of ASD<sup>23</sup> (refer to Appendix 7)

#### b. A complete history should include:

- Concerns by parents / carers
- Symptoms suspicious of ASD (refer to Appendix 7).
- Developmental history
- Behavioural problems and interaction with others
- Medical history including prenatal and perinatal histories
- Psychiatric history – to assess co-existing mental disorders e.g. depression, anxiety
- Family history
- Social history including schooling, home life, physical environment, social needs
- Medication and allergy history

### c. Physical examination should include:

- Presence of dysmorphic features
- Presence of congenital anomalies
- Stigmata of neurofibromatosis or tuberous sclerosis
- Speech / communication skills and developmental assessment
- Hearing / visual assessment
- Signs of physical abuse / self-harm

Other differential diagnoses for ASD to be considered during assessment are shown in Table 1:

**Table 1: Differential diagnoses for autism<sup>23</sup>**

- 
- **Neurodevelopmental disorders:**
    - specific language delay or disorder
    - intellectual disability or global developmental delay
    - developmental coordination disorder (DCD)
  - **Mental and behavioural disorders:**
    - attention deficit hyperactivity disorder (ADHD)
    - mood disorder
    - anxiety disorder
    - attachment disorders
    - oppositional defiant disorder (ODD)
    - conduct disorder
    - obsessive compulsive disorder (OCD)
    - psychosis
  - **Conditions in which there is developmental regression:**
    - Rett syndrome
    - epileptic encephalopathy
  - **Other conditions:**
    - severe hearing impairment
    - severe visual impairment
    - maltreatment
    - selective mutism
- 

## 4.2 Diagnosis

The diagnosis of ASD encompasses difficulties in each of the three domains: social relatedness, communication / play, and restricted interests and activities with onset by three years of age (DSM-IV-TR).

The ICD and DSM categorical system classifications have led to development of the Autism Diagnostic Interview-Revised (ADI-R) and the Diagnostic Interview for Social and Communication Disorders (DISCO). Clinical experience and professional training are crucial in order to establish the diagnosis. Observation in different settings such as school and home may be helpful.

Besides ADI-R and DISCO, other ASD-diagnostic instruments that can be used to facilitate assessment are Autism Diagnostic Observation Schedule (ADOS) and Childhood Autism Rating Scale (CARS). ADOS is a reliable diagnostic instrument to supplement clinical history. Refer to Appendix 8 for more details.

#### **a. Autism Diagnostic Interview-Revised (ADI-R)**

This is a structured interview conducted with parents / carers of individuals for assessment of ASD. It is used for diagnostic purpose for anyone with a mental age of at least 18 months. The interview measures behaviour in the areas of reciprocal social interaction, communication and language, as well as patterns of behaviour.

#### **b. Autism Diagnostic Observation Schedule (ADOS)**

This is a semi-structured assessment of communication, social interaction and play (or imaginative use of materials) for individuals suspected of having ASD. The ADOS consists of four modules, each of which is appropriate for children and adults with differing developmental and language levels, ranging from nonverbal to verbally-fluent.

#### **c. Diagnostic Interview for Social and Communication Disorders (DISCO)**

This is a detailed, semi-structured interview to be used with the parents / carers to identify the impairment of social interaction, social communication and social imagination together with the associated repetitive behaviour, and all other features that may be found in ASD.

#### **d. Childhood Autism Rating Scale (CARS)**

This is a 15-item behaviour observation rating scale to identify and differentiate children with ASD from typically developing children or others with developmental disabilities.

### **RECOMMENDATION 3**

- Diagnosis of autism spectrum disorder should be made clinically based on comprehensive history and observation. Diagnostic tools may be used to assist in the clinical diagnosis. **(Grade C)**

### 4.3 Clinical Progression

Diagnosis of ASD based on clinical evaluation by trained and experienced clinicians is relatively stable.<sup>25, level II-3</sup> Improvement in symptoms reported by parents do not translate into changes in diagnosis.<sup>26, level II-3</sup> The severity levels for ASD vary by context and fluctuate over time.

## 5. Comorbidities and Other Difficulties

Children with ASD can experience a wide range of difficulties with emotional, attentional, activity, thought, behavioural and medical problems. Diagnosis of comorbid disorders is of major importance as they may cause significant clinical impairment in children with ASD. It is crucial that all comorbid conditions are appropriately assessed and managed.

### a. Intellectual Disability

Extreme autistic traits are significantly associated with intellectual disability and poor academic performance ( $p < 0.001$ ).<sup>27, level III</sup> Among those with ASD, about half have intellectual disability.<sup>28, level III</sup>

### b. Attention Deficit Hyperactivity Disorder (ADHD)<sup>29, level III</sup>

Prevalence of ADHD in children with ASD is 53%, with the following subtypes:-

- 22% - hyperactivity / impulsivity
- 46% - inattentive
- 32% - combined

Compared to children with ASD alone, those with comorbid ADHD:-

- are younger, with children between 5 and 7 years presenting more symptoms of hyperactivity ( $p < 0.05$ )
- have a lower mean IQ ( $p = 0.01$ )
- are on medication more often ( $p < 0.05$ )
- do not show differences in gender ( $p = 0.59$ ) and type of ASD diagnosis ( $p = 0.11$ )

### c. Sleep Problems

Sleep problems occur in 44% - 83% of school aged children with ASD. Children with ASD significantly have:<sup>30, level-III; 31, level II-2</sup>

- at least one sleep problem
- sleep onset problems
- night waking

These findings are supported by polysomnography.<sup>31, level II-2</sup> Cognitive or adaptive development did not predict severity of sleep problems in the ASD group.<sup>30, level III</sup>

#### **d. Epilepsy**

The prevalence of epilepsy in autism ranges from 7% - 46%. It is increased with greater intellectual disability, symptomatic autism, age, history of cognitive / developmental regression, use of psychotropics medications and abnormality of electroencephalography (EEG).<sup>32, level III; 33, level III</sup>

The median age of onset for epilepsy is 14 years.<sup>33, level III</sup> Epileptiform abnormalities on EEG occur in 6% - 60% of autistic children.<sup>32, level III</sup> The presence of epilepsy in children with ASD is significantly associated with psychiatric disorders.<sup>34, level III</sup>

#### **e. Gastrointestinal Problems**

Children with ASD are five times more common to have feeding problems than those without ASD (OR 5.11, 95% CI 3.74 to 6.97).<sup>35, level II-2</sup> Types of feeding problems are food selectivity (54%),<sup>35, level II-2; 36, level II-2</sup> food refusal (21%), behavioural rigidity during meals (17%) or their combinations (7%). Children with ASD tend to have a significantly lower consumption of calcium and protein compared to children without ASD and a higher level of nutritional inadequacies.<sup>35, level II-2</sup>

The overall incidence of gastrointestinal symptoms in children with ASD does not differ from the general population, although there is an increased incidence of constipation in those with ASD.<sup>37, level II-2</sup>

#### **f. Motor Coordination**

Substantial motor coordination deficits occur in children with ASD across all age groups. Handwriting is very important for academic progress and social communication development. Adolescents with ASD are known to have poor handwriting. The main predictors are intellectual ability ( $p = 0.006$ ) and motor skill ( $p = 0.005$ ).<sup>38, level III</sup>

## **g. Other Psychiatric Disorders**<sup>34, level III</sup>

Prevalence of psychiatric disorders in children with ASD is as follows:

- 70.8% have at least one current psychiatric disorder (95% CI 58.2 to 83.4) with 57% having multiple diagnosis
- 62.8% (95% CI 49.8 to 75.9) have ADHD, emotional or behavioural disorders (oppositional defiant or conduct)
- 24.7% (95% CI 14.1 to 35.3) have Tourette syndrome, chronic tics, trichotillomania, enuresis, or encopresis (neuropsychiatric disorders)
- 41.9% (95% CI 26.8 to 57.0) have anxiety or phobic disorder
- 1.4% (95% CI 0 to 3.0) have depressive disorders
- 30% (95% CI 14.9 to 45.0) have oppositional or conduct disorder

There is no substantial evidence on the prevalence of psychosis in children with ASD.

## **6. Investigations**

### **a. Audiological Evaluation**

Most children with ASD present with speech and language delay. Therefore, audiological evaluation is an important component of initial assessment in order to rule out hearing impairment. Standard behavioural audiometric procedures are difficult to apply in children with ASD. Behavioural response in audiometric test is less reliable ( $\geq 15$  dB) in younger children with ASD.<sup>39, level III</sup>

The electrophysiological tests used to evaluate hearing impairment are:

- transient evoked otoacoustic emissions (TEOAE)
- auditory brainstem evoked response (ABR)
- acoustic reflexes (AR)

### **b. Other Investigations**

ASD is predominantly a clinical diagnosis. Children with ASD generally do not require intensive investigation. Investigations may be carried out in some children with ASD to establish underlying pathology, exclude treatable conditions and identify comorbid conditions.

- **Electroencephalography (EEG)**

There is insufficient evidence to support the use of EEG in the investigation of children with ASD without clinical seizures.<sup>40, level II-2; 41, level II-2</sup>

## ● Genetic and Metabolic Investigations

Genetic and metabolic studies are not routinely done in children with ASD as the association of inherited metabolic disorders and ASD is low.<sup>42, level II-2</sup> These tests are done usually when there is a suspicion of syndromes such as Fragile X syndrome or when there is dysmorphism or macrocephaly and/or association with severe intellectual disability or global developmental delay. These children have to be referred to a paediatrician or a geneticist for further evaluation.<sup>43, level II-2</sup>

## ● Neuroimaging / Brain Imaging

Brain imaging are not routinely done in patients with ASD as a meta-analysis did not show any difference in the brains of children with ASD and controls.<sup>44, level 1</sup> Brain imaging is usually considered in selective cases where syndromes or neurological conditions are suspected and is usually done at tertiary levels.<sup>45, level III</sup>

## ● Food Allergy

There is no significant association between ASD and food allergies.<sup>46, level II-2; 47, level II-2</sup>

## ● Mineral Analysis

There is insufficient evidence to support the use of mineral analysis in ASD.<sup>48, level II-2; 49, level II-2; 50, level II-2</sup>

### RECOMMENDATION 4

- Audiological assessment should be performed in children with or suspected of having autism spectrum disorder (ASD). **(Grade C)**
- Other investigation\* should not be done routinely in children with ASD unless indicated. **(Grade C)**

\* Refer to preceding texts

## 7. Treatment

Ideally, children with ASD should be managed by a multidisciplinary team as stated in the algorithm (page xiii). Early Intervention Programmes (EIP) are advocated strongly, especially for children below the age of three to improve outcomes. However, all children should be offered EIP upon diagnosis. Policy makers play an important role in the implementation of services for children with ASD.

Children with autism spectrum disorder should be managed by a multidisciplinary team consisting of:

- family medicine specialist
- paediatrician
- psychiatrist / child and adolescent psychiatrist
- clinical / educational psychologist or counsellor
- occupational therapist
- speech-language therapist
- medical social worker
- education officers

#### **RECOMMENDATION 5**

- Parents or carers should actively participate in any intervention offered to children with autism spectrum disorder. **(Grade C)**

### **7.1. Non-pharmacological Treatment**

#### **a. Applied Behaviour Analysis (ABA)**

Applied Behaviour Analysis (ABA) is the application of behavioural principles to everyday situations, that will over time increase or decrease targeted behaviours. This intervention is widely used in the management of children with ASD, and recognised as a safe and effective intervention. Although there are a number of ABA approaches, the Lovaas method is the most well-known and extensively researched.<sup>51, level I</sup>

Lovaas therapy is superior to standard care or regular instruction. In comparison to special education, Lovaas therapy significantly improves among others, social communication skills, language and daily living skills.<sup>51, level I</sup>

There is inadequate evidence to pin-point specific behavioural intervention approaches to be most effective for individual child with ASD. Lovaas therapy and early intensive behavioural intervention variants, and the Early Start Denver Model result in improvements in cognitive performance, language skills, and adaptive behaviour skills in some children.<sup>52, level I</sup>

## RECOMMENDATION 6

- Applied Behaviour Analysis should be considered in the management of children with autism spectrum disorder. **(Grade A)**

### b. Speech, Language and Communication Interventions

Children with ASD may have limited or no speech, poor joint attention and pragmatic skills, as well as difficulty in understanding and interacting with others. Those who receive speech and language therapy between two to three years of age show improvement in expressive language at four years of age ( $p < 0.05$ ).<sup>53, level II-2</sup>

It is important for family members to be active participants in speech therapy sessions. Parents' involvement in therapy of ASD children:-

- increases the number of communication acts and use of communication means ( $p < 0.05$ )<sup>54, level II-2</sup>
- decreases autism behaviour and increases typical communication ( $p < 0.05$ )<sup>55, level I</sup>

The various types of communication interventions for children with ASD are discussed below.

#### ● **Naturalistic Approach**

Responsive Education and Prelinguistic Milieu Teaching (RPMT) facilitates intentional communication during the prelinguistic period in children with developmental delay. Whereas Reciprocal Imitation Training (RIT) is an imitation intervention developed for young children with ASD.

RPMT significantly improves social communication and language learning in children with ASD compared to Picture Exchange Communication System (PECS) in terms of:-

- object exchange turns and requesting<sup>56, level I</sup>
- initiating joint attention<sup>56, level I</sup>
- increasing in object interest<sup>57, level I</sup>

RIT improves elicited and spontaneous imitation of objects, as well as gestures in young children with autism (<5 years old) with a greater play repertoire compared to controls ( $p < 0.05$ ).<sup>58, level I</sup>

- **Augmentative and alternative Communication (AAC)**

Augmentative and Alternative Communication (AAC) is another communication methods that can help children with ASD who have limited speech or no speech to communicate. It is divided into aided and unaided. Unaided AAC consists of nonverbal means of natural communication including gestures and facial expressions. Aided AAC requires some additional external support such as a communication board with visual-graphic symbols using pictures, line drawings or printed words.

AAC improves communication skills (IRD = 0.99, 84% CI 0.98 to 0.99), social skills (IRD = 0.90, 84% CI 0.84 to 0.95), spelling (IRD = 0.79, 84% CI 0.76 to 0.82) and challenging behaviour (IRD = 0.80, 84% CI 0.76 to 0.84) in individuals with ASD.<sup>59, level III</sup>

Picture Exchange Communication System (PECS) is a six-phased aided AAC. It is designed to facilitate functional and spontaneous communication skills. Its use in children with ASD helps them to develop speech.

PECS improves communication skills in individuals with ASD in terms of:-

- non-imitative spoken communication acts ( $p = 0.03$ )<sup>60, level I</sup>
- number of different non-imitative words ( $p = 0.04$ )<sup>60, level I</sup>
- functional communication (IRD = 0.65, 84% CI 0.59 to 0.73)<sup>61, level III</sup>
- challenging behaviour (IRD = 0.45, 84% CI 0.48 to 0.73)<sup>61, level III</sup>
- socialisation (IRD = 0.73, 84% CI 0.53 to 0.93)<sup>61, level III</sup>

PECS increases requesting skills in children with autism based on Percentage of Non-overlapping Data (PND) score.<sup>62, level I</sup> It also increases frequency and number of non-imitative spoken communication compared to RPMT in pre-schoolers with ASD ( $p < 0.05$ ).<sup>56, level I</sup> PECS has larger effects than other picture-based AAC systems in targeted behaviours which include communication and social skills (IRD = 0.99, 84% CI 0.98 to 0.99).<sup>61, level III</sup>

PECS training is effective for pre-school children ( $p < 0.001$ ), those with single ASD diagnosis ( $p = 0.005$ ) and completed all six phases ( $p = 0.005$ ).<sup>61, level III</sup>

A speech generating device (SGD) is an electronic device that produces speech for the user. It is an aided AAC and often used with graphic symbols. SGD with enhanced milieu teaching and signing increase requesting skills in children with autism based on PND score.<sup>62, level I</sup> SGD has larger effects than other picture-based AAC systems in targeted behaviours which include communication and social skills, although it is not statistically significant.<sup>61, level III</sup>

There is no retrievable evidence on unaided AAC.

- **Video Modelling**

Video modelling is a mode of teaching that uses video recording to provide a visual model of the targeted skill to children with ASD. Types of video modelling include video self-modelling and “video modelling with other as model”.

Video modelling and video self-modelling moderately improve social communication skills, functional skills and behavioural functioning in ASD children.<sup>63, level III</sup> “Video modelling with other as model” improves play skills, independent living and social-communicative skills in ASD children ( $p < 0.05$ ).<sup>64, level III</sup>

Two to three years old children with autism spectrum disorder who receive speech and language therapy show improvement in expressive language at four years of age.

#### **RECOMMENDATION 7**

- Children with autism spectrum disorder should receive speech, language and communication interventions as needed. **(Grade A)**

### **c. Occupational Therapy**

ASD may affect the child’s self-care ability, play, academic performance and social activities at home, school and in the community. Sensory processing dysfunction or sensory integration dysfunction are also affected in 50% - 90% of children with ASD.<sup>65; 66; 67; 68</sup> Children with ASD have motor impairment in motor skill development when compared to their typically developing peers.<sup>69; 70; 71</sup>

Occupational therapy provides assessment and intervention to maximise activities of daily living. The interventions provided by occupational therapists include social skills, self-help skills, sensory integration therapy, perceptual motor skills, sensory-motor skills, as well as behavioural and developmental interventions.<sup>72</sup>

- **Social Skills and Self-help Skills**

Social skills and self-help skills interventions improve skill deficits in children with ASD.<sup>73, level III</sup> For social skills intervention, the effectiveness was supported by a Cochrane SR which showed improvement in overall social competence (ES = 0.47, 95% CI 0.16 to 0.78) and friendship quality (ES = 0.41, 95% CI 0.02 to 0.81) in children with ASD.<sup>74, level I</sup>

- **Sensory Integration Therapy (SIT)**

Sensory Integration Therapy (SIT) is often used by occupational therapists to manage children with ASD.<sup>75</sup> The intervention typically involves provision of sensory input by using appropriate modalities.

SIT has shown significant improvements in sensory processing, motor skills, social functioning and autistic mannerisms compared to fine motor intervention,<sup>76, level I</sup> as well as reduction in stereotypic behaviours, off task behaviours and touch aversion compared to no treatment.<sup>51, level I</sup> Yet in a recent SR, SIT showed no consistent positive effect as a treatment for children with ASD, with many of the studies included having serious methodological flaws.<sup>77, level III</sup>

- **Joint Attention Intervention**

Joint attention refers to the child's capacity to coordinate attention with a social partner around an object or event.<sup>78</sup> Joint attention intervention in children with ASD improves joint attention in teacher-child play ( $p = 0.036$ ) and joint engagement in mother-child play ( $p = 0.015$ ).<sup>79, level I</sup>

- **Perceptual Motor Training**

Perceptual motor training involves movement related skills such as eye hand coordination. In a quasi-experimental study, perceptual motor training increased the attention span of children with ASD ( $p < 0.05$ ).<sup>80, level II-1</sup>

### **RECOMMENDATION 8**

- Occupational therapy should be offered to children with autism spectrum disorder. **(Grade A)**

#### **d. Other Interventions / Programmes**

- **Social Stories**

Social stories help children with ASD to understand the subtle differences in interpersonal communication for effective and appropriate interaction. It can be conducted by trained occupational therapists, speech-language therapists, psychologists, teachers or caregivers. Social stories improve total social skills and social isolation ( $p < 0.005$ ),<sup>81, level I</sup> and appropriate social behaviours (mean PND = 56) in children with ASD. Social stories also reduce inappropriate behaviours (mean PND = 87).<sup>82, level III</sup> Standard and directive social story interventions are equally effective in eliciting, generalising and maintaining the targeted social skills in children with autism ( $p < 0.001$ ).<sup>83, level I</sup>

- **Developmental, Individual-Difference, Relationship-Based (DIR) / Floortime**

In this therapy, adults help children to develop their circles of communication by playing with them at their developmental level and building on their strengths. DIR / Floortime™ intervention at an average of 14.2 hours per week for a year significantly improves developmental skills and reduces autistic symptoms in 47% children with ASD.<sup>84, level I</sup>

- **Music Therapy**

In a RCT, children with autism were randomised to receive music therapy and play in different sequences. Generally, music therapy improved joint attention in both groups ( $p = 0.01$ ). However, joint attention improved significantly when play was conducted before music therapy while eye contact events occurred significantly longer when music therapy was instituted before play.<sup>85, level I</sup>

#### **e. Parent Education and Support**

Managing children with ASD can be a huge challenge to their families. Children need parental involvement and support in acquiring social and communication skills. Parents without proper training may contribute to the failure of children applying learnt skills outside the classrooms. Thus, parental involvement is important to ensure the continuity of intervention.

A SR showed that parental training improved parent-child interaction (shared attention: SMD = 0.41, 95% CI 0.14 to 0.68 and parent synchrony: SMD = 0.90, 95% CI 0.56 to 1.23). Besides that, improvement was also found in child's language comprehension (MD = 36.26, 95% CI 1.31 to 71.20) as a result of parental training.<sup>86, level 1</sup>

Children's functional verbal utterances increase following parental training ( $p < 0.05$ ) and are maintained at follow-up (2 - 4 months). The relationship between parent fidelity to the training and child's communication improves from pre-training to follow-up ( $p < 0.05$ ).<sup>87, level III</sup>

#### **RECOMMENDATION 9**

- Parental training should be offered to parents of children with autism spectrum disorder. **(Grade A)**

#### **f. Cognitive Behaviour Therapy (CBT)**

Cognitive Behaviour Therapy (CBT) is used in children with ASD who have anxiety and mood disorders.

The application of modified CBT is effective in reducing anxiety in children with Asperger subtype but not in other subtypes of ASD.<sup>88, level I</sup> CBT modalities that are effective in the treatment of anxiety in ASD include Coping Cat,<sup>89, level I</sup>; 90, level II-3 Building Confidence CBT Program<sup>91, level I</sup>; 92, level I and modules specifically developed for ASD children.<sup>93, level I</sup> However, most of the primary studies were done on children with high verbal intelligence.

#### **RECOMMENDATION 10**

- Cognitive Behaviour Therapy may be offered to children with autism spectrum disorder who have high verbal intelligence and anxiety disorder. **(Grade A)**

#### **g. Treatment and Education of Autistic and Related Communication Handicapped Children (TEACCH)**

This is a structured teaching system based on a close collaboration between parents and professionals. It can be used for children with ASD of all ages and skill levels.

There is no significant difference between TEACCH and standard care in imitation skills and eye-hand integration,<sup>51, level I</sup> although there is moderate to large effect sizes (ES) in social functioning (ES = 0.65, 95% CI 0.15 to 1.15) and maladaptive behaviour (pooled ES = -0.92, 95% CI -1.51 to -0.33)<sup>94, level I</sup>

Treatment and Education of Autistic and Related Communication Handicapped Children (TEACCH) may be useful in the management of children with autism spectrum disorder.

#### **h. Diet**

Gluten and casein-free diet, and dimethylglycine are not effective in the management of autism.<sup>95, level I</sup>; 96, level I

Omega-3 supplements have no significant effect on social interaction, communication, stereotypy or hyperactivity.<sup>97, level I</sup>

Diet-related intervention has no significant benefit in children with autism spectrum disorder.

## 7.2. Pharmacotherapy

In ASD, medications are used in the treatment of comorbid disorders. Conventional antipsychotics such as haloperidol are used less frequently due to its high incidence of severe adverse reactions.

### a. Atypical Antipsychotics (AAP)

The use of atypical antipsychotics in ASD is preferred due to the reduced propensity of causing extrapyramidal symptoms.

#### ● Risperidone

Low dose risperidone (up to 2.5 mg per day in children weighing from 20 - 45 kg and up to 3.5 mg per day in those weighing over 45 kg) may be beneficial in some features of ASD.<sup>98, level I</sup> A meta-analysis of three RCTs in children with ASD suggested that short-term use of risperidone significantly improved:-

- irritability (MD = -8.09, 95% CI -12.99 to -3.19)
- social withdrawal / lethargy (MD = -1.00, 95% CI -5.03 to -0.97)
- hyperactivity (MD = -8.98, 95% CI -12.01 to -5.94)
- stereotypy (MD = -1.71, 95% CI -2.97 to -0.45)
- inappropriate speech (MD = -1.93, 95% CI -3.79 to -0.07)

However, there was a higher risk of weight gain (MD = 1.78, 95% CI 1.15 to 2.41) in the treatment group compared to those on placebo.<sup>99, level I</sup>

#### ● Aripiprazole

A meta-analysis of two RCTs in children with ASD suggested that aripiprazole up to 15 mg per day might be efficacious in treating:-

- irritability (MD = -6.17, 95% CI -9.07 to -3.26)
- hyperactivity (MD = -7.93, 95% CI -10.98 to -4.88)
- stereotypy (MD = -2.66, 95% CI -3.55 to -1.77)
- inappropriate speech (MD = -1.43, 95% CI -2.60 to -0.27)

Adverse effects such as weight gain (MD = 1.13, 95% CI 0.71 to 1.54), sedation (MD = 4.28, 95% CI 1.58 to 11.60), drooling (MD = 9.64, 95% CI 1.29 to 72.10) and tremor (MD = 10.26, 95% CI 1.37 to 76.63) may occur more often in the treatment group compared to those on placebo.<sup>100, level I</sup>

- **Other AAPs**

- **Olanzapine**

Olanzapine up to 20 mg per day improves CGI-I score in pervasive developmental disorder (PDD) at eight weeks ( $p = 0.012$ ). However, clinically significant weight gain is more likely to occur in individuals treated with the medication ( $p = 0.028$ ).<sup>101, level I</sup>

- **Paliperidone**

Paliperidone up to 12 mg per day is efficacious for irritability in individuals with autism ( $p = 0.0002$ ). The safety profile is acceptable except for raised prolactin level in males ( $p = 0.0001$ ).<sup>102, level II-3</sup>

- **Quetiapine**

Low dose quetiapine ( $\leq 150$  mg per day) is efficacious in reducing aggression ( $p = 0.028$ ) and improving sleep quality ( $p = 0.014$ ) in children with ASD at eight weeks. No significant difference in body weight was observed and the adverse effects were mild.<sup>103, level II-3</sup>

- **Ziprasidone**

Ziprasidone up to 160 mg per day is efficacious in reducing irritability ( $p = 0.05$ ) and hyperactivity ( $p = 0.01$ ) in autism at six weeks. The serious side effect of QTc prolongation of ziprasidone may offset the arguably minimal benefits associated with its use. ( $p = 0.04$ ).<sup>104, level II-3</sup>

## **b. Antidepressants**

- **Selective Serotonin Reuptake Inhibitors (SSRI)**

A Cochrane SR explored the efficacy of various selective serotonin reuptake inhibitors (SSRIs) in the treatment of children with ASD. There was no evidence of SSRIs effect in children on core symptoms and severity of the disorder. There was also emerging evidence of harm such as seizure, poor appetite and weight loss.<sup>105, level I</sup>

- **Tricyclic Antidepressants (TCA)**

A Cochrane SR concluded that there was limited and conflicting evidence of effect and the side effects of TCA as a treatment option in children with ASD. For example, a study in 1993 favoured clomipramine

against placebo but a later study in 2001 showed non-significant finding with substantial side effects. Further research is required before TCAs can be recommended for treatment of children with ASD.<sup>106, level I</sup>

### **c. Other Medications**

- **Methylphenidate**

Methylphenidate was found to be more efficacious than placebo among subjects who have PDD in a crossover RCT followed by open-label continuation of three months duration. Parent-rated and teacher-rated Aberrant Behaviour Checklist hyperactivity subscale scores during crossover phase were significantly lower in different methylphenidate dosages and the response was maintained in the continuation phase. Reported side effects were loss of appetite, sleep difficulties and irritability.<sup>107, level I</sup>

- **Atomoxetine**

A SR showed that there was limited evidence to suggest the efficacy of atomoxetine in children with ASD and ADHD symptoms. Further research is required before it can be strongly recommended for the treatment of children with such condition.<sup>108, level I</sup>

- **Valproate**

In a RCT, divalproate sodium significantly reduced irritability compared to the placebo. The findings were robust even after controlling for intelligence quotient (IQ) differences.<sup>109, level I</sup> It was well tolerated. Most side effects were mild to moderate in severity, resolved with small changes in dosing and did not require a discontinuation of medication.

***Except for risperidone and aripiprazole, the drugs listed above are for off-label use.***

- **Melatonin**

Melatonin is an endogenous neurohormone produced predominantly in the pineal gland. It is commonly used for insomnia in children and has a favourable side-effect profile.<sup>110</sup>

In a meta-analysis, melatonin was more efficacious than placebo in the treatment of ASD in terms of:<sup>111, level I</sup>

- increased sleep duration by 44 minutes (Hedge's  $g = 1.07$ , 95% CI 0.49 to 1.65)
- shorter sleep onset latency by 39 minutes (Hedge's  $g = 2.46$ , 95% CI 1.96 to 2.98)

The findings above were supported by a recent RCT whereby a combination of CBT and melatonin was statistically most efficacious in reducing insomnia symptoms compared to either modality or placebo when used alone.<sup>112, level I</sup>

In terms of safety, mild side effects reported in the meta-analysis were headache, diarrhoea and dizziness.<sup>111, level I</sup>

### RECOMMENDATION 11

- Children with autism spectrum disorder may be offered:
  - atypical antipsychotics as a short-term treatment for irritability. **(Grade A)**
  - methylphenidate for hyperactivity. **(Grade A)**
  - atomoxetine for hyperactivity. **(Grade C)**
  - melatonin for sleep difficulties. **(Grade A)**

## 7.3 Traditional and Complementary Medicine

Any other treatment modalities not described above are grouped under traditional and complementary medicine. The following treatment modalities have poor evidence, no effectiveness and / or some harmful effects in children with ASD:-

- chelation<sup>113, level II-3</sup>
- secretin<sup>114, level I</sup>
- fatty acids<sup>97, level I</sup>
- vitamin B6-magnesium<sup>115, level I</sup>
- vitamin B12<sup>116, level I</sup>
- acupuncture<sup>117, level I</sup>
- hyperbaric oxygen therapy<sup>118, level I</sup>

There is no retrievable evidence for vitamin A, vitamin C, trimethylglycine, cupping, ayurvedic medicine or homeopathy in the treatment of children with ASD.

Traditional and Complementary Medicine could not be recommended to children with autism spectrum disorder because of insufficient evidence and potential harmful effects.

## 8. Social Welfare Service

Children with ASD should be referred to the Department of Social Welfare at their respective local districts. This will enable the child to be registered for benefits such as:

- placement for special needs education if warranted
- welfare support including financial allowances from the department
- free services in the public sector

The registration form should be completed by the medical practitioner / medical officer at the point of diagnosis.

## 9. Monitoring and Preparation of Transition to Adult Services

Children with ASD require clinical monitoring for physical growth, behaviour and development, associated medical problems, as well as other comorbidities, sexuality and safety. The main goal is for children with ASD to become independent and achieve their potential in all aspects of life, leading to a higher quality of life when they exit the school system. However, there is a lack of reliable and valid measures to evaluate progress and change of a child's behaviour and functioning over time after the diagnosis of ASD.<sup>119, level II-2</sup>

Transition for adolescents with ASD should be discussed and planned early by all involved in their management. ASD children have varying intellectual and functional abilities, hence the transition into adulthood has to be planned early according to their individual abilities.<sup>120, level III</sup>

Vocational training, post-secondary education, day care activities and supervised employment are options for these young people.<sup>121, level II-1</sup>

Care for children and adolescents with ASD should be continued into adult health services. There is a need for establishment of this service to support adolescents when they enter adulthood.

## 10. Implementing the Guidelines

Implementation of this CPG is the responsibility of each healthcare provider. Mechanisms should be in place to review care provided against the guidelines recommendations. The reasons for any differences should be assessed and addressed where appropriate. Local arrangements should then be made to implement the national guidelines in individual hospitals and clinics.

## **a. Facilitating and Limiting Factors**

The facilitating factors in implementing these CPG are:-

### **i. Dissemination of CPG**

- Availability and dissemination of CPG to health care providers (hard and soft copies)
- Conferences and updates on child behaviour and development

### **ii. Implementation of CPG**

- Developmental screening including M-CHAT
- Public awareness via World Autism Day
- Accessibility to relevant multidisciplinary teams
- Active involvement of government and non-governmental organisations

The limiting factors in the implementation are:

- Limited awareness and knowledge of detection and subsequent referral of children with ASD
- Variation of practice and treatment at different levels of care
- Limited financial and human resources
- Lack of training at all levels of healthcare providers
- Lack of networking between government and private practitioners

## **b. Potential Resource Implications**

- Widespread distribution of CPG to healthcare personnel via printed copies
- Reinforce training of healthcare providers via regular seminars and workshops
- Establish ASD registry in Malaysia
- Develop a multidisciplinary team in secondary and tertiary care levels.

To enhance the utilisation of these CPG on the Management of Autism Spectrum Disorder in Children and Adolescents, the following clinical audit indicators for quality management are proposed:-

$$\text{Percentage of children aged 18 and 24 months screened with M-CHAT} = \frac{\text{Number of children aged 18 and 24 months screened with M-CHAT in a year}}{\text{Number of children aged 18 and 24 months attended child health clinic in the same period}} \times 100\%$$

$$\text{Percentage of children with failed M-CHAT referred for further assessment*} = \frac{\text{Number of children with failed M-CHAT referred for further assessment in a year}}{\text{Number of children failed M-CHAT in the same period}} \times 100\%$$

\*Assessment by MDT

## References

1. Family Health Division. Prosiding Mesyuarat Membincangkan Hasil Kajian Saringan dan Pengendalian Masalah Autisme. Kuala Lumpur 2006.
2. Developmental Disabilities Monitoring Network Surveillance Year 2010 Principal Investigators; Centers for Disease Control and Prevention (CDC). Prevalence of autism spectrum disorder among children aged 8 years - autism and developmental disabilities monitoring network, 11 sites, United States, 2010. *MMWR Surveill Summ.* 2014, 63(2):1-21.
3. Sandin S, Hultman CM, Kolvezon A, et al. Advancing maternal age is associated with increasing risk for autism: a review and meta-analysis. *J Am Acad Child Adolesc Psychiatry.* 2012, 51(5):477-486.
4. Guinchat V, Thorsen P, Laurent C, et al. Pre-, peri- and neonatal risk factors for autism. *Acta Obstet Gynecol Scand.* 2012, 91(3):287-300.
5. Hultman CM, Sandin S, Levine SZ, et al. Advancing paternal age and risk of autism: new evidence from a population-based study and a meta-analysis of epidemiological studies. *Mol Psychiatry.* 2011, 16(12):1203-1212.
6. Durkin MS, Maenner MJ, Newschaffer CJ, et al. Advanced parental age and the risk of autism spectrum disorder. *Am J Epidemiol.* 2008, 168(11):1268-1276.
7. Ozonoff S, Young GS, Carter A, et al. Recurrence risk for autism spectrum disorders: A Baby Siblings Research Consortium Study. *Pediatrics.* 2011, 128:e488-495.
8. Surén P, Roth C, Bresnahan M, et al. Association between maternal use of folic acid supplements and risk of autism spectrum disorders in children. *JAMA.* 2013, 309(6):570-577.
9. Schmidt RJ, Hansen RL, Hartiala J, et al. Prenatal vitamins, one-carbon metabolism gene variants, and risk for autism. *Epidemiology.* 2011, 22(4):476-485.
10. Demicheli V, Rivetti A, Debalini MG, et al. Vaccines for measles, mumps and rubella in children. *Cochrane Database Syst Rev.* 2012, 2:CD004407.
11. DeStefano F, Price CS, Weintraub ES. Increasing exposure to antibody stimulating proteins and polysaccharides in vaccines is not associated with the risk of autism. *J Paediatrics.* 2013, 163(2):561-567.
12. Sunita, Bilszta JLC. Early identification of autism: A comparison of the Checklist for Autism in Toddlers and the Modified Checklist for Autism in Toddlers. *J Paediatr Child Health.* 2013, 49(6):438-444.
13. Norris M, Lecavalier L. Screening accuracy of level 2 autism spectrum disorder rating scales: a review of selected instruments. *Autism.* 2010, 14(4):263-284.
14. Mawlea E, Griffiths P. Screening for autism in pre-school children in primary care: systematic review of English language tools. *Int J Nurs Stud.* 2006, 43(5):623-636
15. Robins DL, Fein D, Barton ML, et al. The Modified Checklist for Autism in Toddlers: An initial study investigating the early detection of autism and pervasive developmental disorders. *J Autism Dev Disord.* 2001, 31(2):131-144.
16. Berument SK, Rutter M, Lord C, et al. Autism screening questionnaire: diagnostic validity. *Br J Psychiatry.* 1999, 175:444-445.
17. Wong V, Hui LH, Lee WC, et al. A modified screening tool for autism (Checklist for Autism in Toddlers [CHAT-23]) for Chinese children. *Pediatrics.* 2004, 114(2):e166-176.

18. Ooi YP, Rescorla L, Ang RP, et al. Identification of autism spectrum disorders using the child behaviour checklist in Singapore. *J Autism Dev Disord*. 2011, 41(9):1147-1156.
19. Wetherby AM, Brosnan-Maddox S, Peace V, et al. Validation of the Infant-Toddler Checklist as a broadband screener for autism spectrum disorders from 9 to 24 months of age. *Autism*. 2008, 12(5):487-511.
20. Ozonoff S, Young GS, Steinfeld MB, et al. How early do parent concerns predict later autism diagnosis? *J Dev Behav Pediatrics*. 2009, 30(5):367-375.
21. Baron-Cohen S, Allen J, Gillberg C. Can autism be detected at 18 months? The needle, the haystack and the CHAT. *Br J Psychiatry*. 1992, 161:839-843.
22. Robins DL, Casagrande K, Barton M, et al. Validation of the modified checklist for autism in toddlers, revised with follow-up (M-CHAT-R/F). *Pediatrics*. 2014, 133(1):37-45.
23. National Institute for Health and Care Excellence. Autism diagnosis in children and young people: Recognition, referral and diagnosis of children and young people on the autism spectrum, 2011, National Clinical Guideline, Number 128 2011.
24. Volkmar F, Siegel M, Woodbury-Smith M, et al. Practice parameter for the assessment and treatment of children and adolescents with autism spectrum disorder. *J Am Acad Child Adolesc Psychiatry*. 2014, 53(2):237-257.
25. Kleinman JM, Ventola PE, Pandey J, et al. Diagnostic stability in very young children with autism spectrum disorders. *J Autism Dev Disord*. 2008, 38(4):606-615.
26. Soke GN, Philofsky A, Diguseppi C, et al. Longitudinal changes in scores on the Autism Diagnostic Interview–Revised (ADI-R) in pre-school children with autism: Implications for diagnostic classification and symptom stability. *Autism*. 2011, 15(5):545-562.
27. Hoekstra RA, Happé F, Baron - Cohen S, et al. Association between extreme autistic traits and intellectual disability: insights from a general population twin study. *Br J Psychiatry*. 2009, 195(6):531-536.
28. Charman T, Pickles A, Simonoff E, et al. IQ in children with autism spectrum disorders: data from the Special Needs and Autism Project (SNAP). *Psychol Med*. 2011, 41(3):619-627.
29. Sinzig J, Walter D, Doepfner M. Attention deficit / hyperactivity disorder in children and adolescents with autism spectrum disorder: symptom or syndrome? *J Atten Disord*. 2009, 13(2):117-126.
30. Krakowiak P, Goodlin-Jones B, Hertz-Picciotto I, et al. Sleep problems in children with autism spectrum disorders, developmental delays, and typical development: a population-based study. *J Sleep Res*. 2008, 17(2):197-206.
31. Miano S, Bruni O, Elia M, et al. Sleep in children with autistic spectrum disorder: a questionnaire and polysomnographic study. *Sleep Med*. 2007, 9(1):64-70.
32. Sansa G, Carlson C, Doyle W, et al. Medically refractory epilepsy in autism. *Epilepsia*. 2011, 52(6):1071-1075.
33. Hara H. Autism and epilepsy: a retrospective follow-up study. *Brain Dev*. 2007, 29(8):486-490.

34. Simonoff E, Pickles A, Charman T, et al. Psychiatric disorders in children with autism spectrum disorders: prevalence, comorbidity, and associated factors in a population-derived sample. *J Am Acad Child Adolesc Psychiatry*. 2008, 47(8):921-929.
35. Sharp WG, Berry RC, McCracken C, et al. Feeding problems and nutrient intake in children with autism spectrum disorders: a meta-analysis and comprehensive review of the literature. *J Autism Dev Disord*. 2013, 43(9):2159-2173.
36. Mari-Bauset S, Zazpe I, Mari-Sanchis A, et al. Food selectivity in autism spectrum disorders: A systematic review. *J Child Neurol*. 2013, [Epub ahead of print].
37. Ibrahim SH, Voigt RG, Katusic SK, et al. Incidence of gastrointestinal symptoms in children with autism: a population-based study. *Pediatrics*. 2009, 124(2):680-686.
38. Fuentes CT, Mostofsky SH, Bastian AJ. Perceptual reasoning predicts handwriting impairments in adolescents with autism. *Neurology*. 2010, 75(20):1825-1829.
39. Tharpe AM, Bess FH, Sladen DP, et al. Auditory characteristics of children with autism. *Ear and Hearing*. 2006, 27(4):430-441.
40. Fong CY, Baird G, Wraige E. Do children with autism and developmental regression need EEG investigation in the absence of clinical seizures? *Arch Dis Child*. 2008, 93:998-999.
41. Kagan-Kushnir T, Roberts SW, Snead OC. Screening electroencephalograms in autism spectrum disorders: evidence-based guideline. *J Child Neurol*. 2005, 20(3):197-206.
42. Schiff M, Benoist JF, Aïssaoui S, et al. Should metabolic diseases be systematically screened in nonsyndromic autism spectrum disorders? *PLoS One*. 2011, 6(7):e219-232.
43. Battaglia A, Carey JC. Etiologic yield of autistic spectrum disorders: a prospective study. *Am J Med Genet C Semin Med Genet*. 2006, 142C(1):3-7.
44. Via E, Radua J, Cardoner N, et al. Meta-analysis of gray matter abnormalities in autism spectrum disorder: should Asperger disorder be subsumed under a broader umbrella of autistic spectrum disorder? *Arch Gen Psychiatry*. 2011, 68(4):409-418.
45. Boddaert N, Zilbovicius M, Philippe A, et al. MRI findings in 77 children with nonsyndromic autistic disorder. *PLoS One*. 2009, 4(2):e4415.
46. Bakkaloglu B, Anlar B, Anlar FY, et al. Atopic features in early childhood autism. *Eur J Paediatr Neurol*. 2008, 12(6):476-479.
47. Jyonouchi H, Geng L, Cushing-Ruby A, et al. Impact of innate immunity in a subset of children with autism spectrum disorders: a case control study. *J Neuroinflammation*. 2008, 21(5):52.
48. Fido A, Al-Saad S. Toxic trace elements in the hair of children with autism. *Autism*. 2005, 9(3):290-298.
49. De Palma G, Catalani S, Franco A, et al. Lack of correlation between metallic elements analyzed in hair by ICP-MS and autism. *J Autism Dev Disord*. 2012, 42(3):342-353.
50. Yorbik O, Kurt I, Haşimi A, et al. Chromium, cadmium, and lead levels in urine of children with autism and typically developing controls. *Biol Trace Elem Res*. 2010, 135(1-3):10-15.
51. Ospina MB, Krebs Seida J, Clark B, et al. Behavioural and developmental interventions for autism spectrum disorder: a clinical systematic review. *PLoS One*. 2008, 3(11):e3755.
52. Warren Z, McPheeters ML, Sathe N. A systematic review of early intensive intervention for autism spectrum disorders. *Pediatrics*. 2011, 127(5):e1303-1301.
53. Stone WL, Yoder PJ. Predicting spoken language level in children with autism spectrum disorders. *Autism*. 2001, 5(4):341-361.

54. Fernandes FD, Cardoso C, Sassi FC, et al. Language therapy and autism: results of three different models. *Pro Fono Revista de Atualizacao Cientifica*. 2008, 20(4):267-272.
55. Tamanaha AC, Perissinoto J. Comparison of the evolutionary process of children with autism spectrum disorders in different language therapeutic interventions. *J Soc Bras Fonoaudiol*. 2011, 23(1):8-12.
56. Yoder P, Stone WL. Randomized comparison of two communication interventions for preschoolers with autism spectrum disorders. *J Consult Clin Psychol*. 2006, 74(3):426-435.
57. McDuffie AS, Lieberman RG, Yoder PJ. Object interest in autism spectrum disorder: a treatment comparison. *Autism*. 2012, 16(4):398-405.
58. Ingersoll B. Pilot randomized controlled trial of Reciprocal Imitation Training for teaching elicited and spontaneous imitation to children with autism. *J Autism Dev Disord*. 2010, 40(9):1154-1160.
59. Ganz JB, Earles-Vollrath TL, Heath AK, et al. A meta-analysis of single case research studies on aided augmentative and alternative communication systems with individuals with autism spectrum disorders. *J Autism Dev Disord*. 2012, 42(1):60-74.
60. Flippin M, Reszka S, Watson LR. Effectiveness of the Picture Exchange Communication System (PECS) on communication and speech for children with autism spectrum disorders: a meta-analysis. *Am J Speech Lang Pathol*. 2010, 19(2):178-195.
61. Ganz JB, Davis JL, Lund EM, et al. Meta-analysis of PECS with individuals with ASD: investigation of targeted versus non-targeted outcomes, participant characteristics, and implementation phase. *Res Dev Disabil*. 2012, 33(2):406-418.
62. Schlosser RW, Wendt O. Effects of augmentative and alternative communication intervention on speech production in children with autism: a systematic review. *Am J Speech Lang Pathol*. 2008, 17(3):212-230.
63. Bellini S, Akullian J. A meta-analysis of video modeling and video self-modeling interventions for children and adolescents with autism spectrum disorders. *Exceptional Children*. 2007, 73:264-287.
64. Mason RA, Ganz JB, Parker RI, et al. Moderating factors of video-modeling with other as model: a meta-analysis of single-case studies. *Res Dev Disabil*. 2012, 33(4):1076-1086.
65. Baranek GT, David FJ, Poe MD, et al. Sensory experiences questionnaire: discriminating sensory features in young children with autism, developmental delays, and typical development. *J Child Psychol Psychiatry*. 2006, 47(6):591-601.
66. Leekam SR, Nieto C, Libby SJ, et al. Describing the sensory abnormalities of children and adults with autism. *J Autism Dev Disord*. 2007, 37(5):894-910.
67. Tomchek SD, Dunn W. Sensory processing in children with and without autism: a comparative study using the short sensory profile. *Am J Occup Ther*. 2007, 61(2):190-200.
68. Baker AE, Lane A, Angley MT, et al. The relationship between sensory processing patterns and behavioural responsiveness in autistic disorder: a pilot study. *J Autism Dev Disord*. 2008, 38(5):867-875.
69. Berkeley SL, Zittel II, Pitney IV, et al. Locomotor and object control skills of children diagnose with autism. *Adapt Phys Activ Q*. 2001, 18(4):405-416.
70. Dziuk MA, Gidley Larson JC, Apostu A, et al. Dyspraxia in autism: association with motor, social, and communicative deficits. *Dev Med Child Neurol*. 2007, 49(10):734-739.

71. Breslin CM, Rudisill ME. The effect of visual supports on performance of the TGMD-2 for children with autism spectrum disorder. *Adapt Phys Activ Q.* 2011, 28(4):342-353.
72. Case-Smith J, Arbesman M. Evidence-based review of interventions for autism used in or of relevance to occupational therapy. *Am J Occup Ther.* 2008, 62(4):416-429.
73. Flynn L, Healy O. A review of treatments for deficits in social skills and self-help skills in autism spectrum disorder. *Res in Autism Spectr Disord.* 2012, 6 (1):431-441.
74. Reichow B, Steiner AM, Volkmar F. Social skills groups for people aged 6 to 21 with autism spectrum disorders (ASD). *Cochrane Database Syst Rev.* 2012, 7:CD008511.
75. Ayres AJ, Tickle LS. Hyper-responsivity to touch and vestibular stimuli as a predictor of positive response to sensory integration procedures. *Am J Occup Ther,* 1980, 34:375-381.
76. Pfeiffer BA, Koenig K, Kinnealey M, et al. Effectiveness of sensory integration interventions in children with autism spectrum disorders: a pilot study. *Am J Occup Ther.* 2011, 65(1):76-85.
77. Lang R, O'Reilly M, Healy O, et al. Sensory integration therapy for autism spectrum disorders: a systematic review. *Res Autism Spectr Disord.* 2012, 6(3):1004-1018.
78. Mundy P, Sigman M. Joint attention, social competence, and developmental psychopathology In D. Cicchetti & D. Cohen (Eds.) *Developmental psychopathology, 2nd ed., Vol.1: Theory and Methods* (pp. 293 - 332). 2006
79. Kaale A, Smith L, Sponheim E. A randomized controlled trial of preschool-based joint attention intervention for children with autism. *J Child Psychol Psychiatry.* 2012, 53(1):97-105.
80. Afshari J. The effect of perceptual-motor training on attention in the children with autism spectrum disorders. *Res Autism Spectr Disord.* 2012, 6:1331-1336.
81. Karkhaneh M, Clark B, Ospina MB, et al. Social Stories™ to improve social skills in children with autism spectrum disorder: a systematic review. *Autism.* 2010, 14(6):641-662.
82. Kokina A, Kern L. Social story interventions for students with autism spectrum disorders: a meta-analysis. *J Autism Dev Disord.* 2010, 40(7):812-826.
83. Quirnbach LM, Lincoln AJ, Feinberg-Gizzo MJ, et al. Social stories: mechanisms of effectiveness in increasing game play skills in children diagnosed with autism spectrum disorder using a pretest posttest repeated measures randomized control group design. *J Autism Dev Disord.* 2009, 39(2):299-232.
84. Pajareya K, Nopmaneejumruslers K. A pilot randomized controlled trial of DIR/ Floortime™ parent training intervention for pre-school children with autistic spectrum disorders. *Autism.* 2011, 15(5):563-577.
85. Kim J, Wigram T, Gold C. The effects of improvisational music therapy on joint attention behaviors in autistic children: a randomized controlled study. *J Autism Dev Disord.* 2008, 38(9):1758-1766.
86. Oono IP, Honey EJ, McConachie H. Parent-mediated early intervention for young children with autism spectrum disorders (ASD). *Cochrane Database Syst Rev.* 2013, CD009774.pub009772.
87. Coolican J, Smith IM, Bryson SE. Brief parent training in pivotal response treatment for preschoolers with autism. *J Child Psychol Psychiatry.* 2010, 51(12):1321-1330.
88. Lang R, Regester A, Lauderdale S, et al. Treatment of anxiety in autism spectrum disorders using cognitive behaviour therapy: a systematic review. *Dev Neurorehabil.* 2010, 13(1):53-63.
89. McNally Keehn RH, Lincoln AJ, Brown MZ, et al. The Coping Cat program for children with anxiety and autism spectrum disorder: a pilot randomized controlled trial. *J Autism Dev Disord.* 2013, 43(1):57-67.

90. Viecili MA. Effects of A modified cognitive behavior therapy program to treat anxiety in children with autism spectrum disorders. ProQuest Dissertations & Theses. Toronto, Ontario: York University; 2011.
91. Drahota A, Wood JJ, Sze KM, et al. Effects of cognitive behavioral therapy on daily living skills in children with high-functioning autism and concurrent anxiety disorders. *J Autism Dev Disord.* 2011,41(3):257-265.
92. Wood JJ, Drahota A, Sze K, et al. Brief report: effects of cognitive behavioral therapy on parent-reported autism symptoms in school-age children with high-functioning autism. *J Autism Dev Disord.* 2009, 39(11):1608-1612.
93. Sung M, Ooi YP, Goh TJ, et al. Effects of cognitive-behavioral therapy on anxiety in children with autism spectrum disorders: a randomized controlled trial. *Child Psychiatry Hum Dev.* 2011, 42(6):634-649.
94. Virues-Ortega J, Julio FM, Pastor-Barriuso R. The TEACCH program for children and adults with autism: a meta-analysis of intervention studies. *Clin Psychol Rev.* 2013, 33(8):940-953.
95. Millward C, Ferriter M, Calver S, et al. Gluten- and casein-free diets for autistic spectrum disorder. *Cochrane Database Syst Rev.* 2008, 2:CD003498.
96. Kern JK, Miller VS, Cauller PL, et al. Effectiveness of N,N-dimethylglycine in autism and pervasive developmental disorder. *J Child Neurology.* 2001, 16(3):169-173.
97. James S, Montgomery P, Williams K. Omega-3 fatty acids supplementation for autism spectrum disorders (ASD). *Cochrane Database Syst Rev.* 2011(11):CD007992.
98. McCracken JT, McGough J, Shah B, et al. Risperidone in children with autism and serious behavioral problems. *N Engl J Med.* 2002, 347(5):314-321.
99. Jesner OS, Aref-AdibM, Coren E. Risperidone for autism spectrum disorder. *Cochrane Database of Syst Rev.* 2007, (1):CD005040.
100. Ching H, Pringsheim T. Aripiprazole for autism spectrum disorders (ASD). *Cochrane Database Syst Rev.* 2012, 5:CD009043.
101. Hollander E, Wasserman S, Swanson E, et al. A double-blind placebo-controlled pilot study of olanzapine in childhood/adolescent Pervasive Developmental disorder. *J Child Adolesc Psychopharmacol.* 2006, 16(5):541-548.
102. Stigler KA, Mullett JE, Erickson CA, et al. Paliperidone for irritability in adolescents and young adults with autistic disorder. *Psychopharmacology.* 2012, 223(2):237-245.
103. Golubchik P, Sever J, Weizman A. Low-dose quetiapine for adolescents with autistic spectrum disorder and aggressive behavior: open-label trial. *Clin Neuropharmacol.* 2011, 34(6):216-219.
104. Malone RP, Delaney MA, Hyman SB, et al. Ziprasidone in adolescents with autism: an open-label pilot study. *J Child Adolesc Psychopharmacol.* 2007, 17(6):779-790.
105. Williams K, Wheeler DM, Silove N, et al. Selective serotonin reuptake inhibitors (SSRIs) for autism spectrum disorders (ASD). *Cochrane Database Syst Rev.* 2010 (8):CD004677.
106. Hurwitz R, Blackmore R, Hazell P, et al. Tricyclic antidepressants for autism spectrum disorders (ASD) in children and adolescents. *Cochrane Database Syst Rev.* 2012, (3):CD008372.
107. Research Units on Pediatric Psychopharmacology Autism Network. Randomized, controlled, crossover trial of methylphenidate in pervasive developmental disorders with hyperactivity. *Arch Gen Psychiatry.* 2005, 62(11):1266-1274.
108. Ghanizadeh A. Atomoxetine for treating ADHD symptoms in autism: A systematic review. *J Atten Disord.* 2013, 17(635-640).

109. Hollander E, Chaplin W, Soorya L, et al. Divalproex sodium vs placebo for the treatment of irritability in children and adolescents with autism spectrum disorders. *Neuropsychopharmacology*. 2010, 35(4):990-998.
110. Johnson KP, Malow BA. Assessment and pharmacologic treatment of sleep disturbance in autism. *Child Adolesc Psychiatr Clin N Am*. 2008, 17:773-785.
111. Rossignol D, Frye R. Melatonin in autism spectrum disorders: a systematic review and meta-analysis. *Dev Med Child Neurol*. 2011, 53(9):783-792.
112. Cortesi F, Giannotti F, Sebastiani T, et al. Controlled-release melatonin, singly and combined with cognitive behavioural therapy, for persistent insomnia in children with autism spectrum disorders: a randomized placebo-controlled trial. *J Sleep Res*. 2012, 21(6):700-709.
113. Davisa TN, O'Reilly M, Kangb S, et al. Chelation treatment for autism spectrum disorders: A systematic review. *Res Autism Spectr Disord*. 2013, 7(1):49-55.
114. Williams K, Wray JA, Wheeler DM. Intravenous secretin for autism spectrum disorders (ASD). *Cochrane Database Syst Rev*. 2012, (4):CD003495.
115. Nye C, Brice A. Combined vitamin B6-magnesium treatment in autism spectrum disorder. *Cochrane Database Syst Rev*. 2005, (4):CD003497.
116. Bertoglio K, Jill James S, Deprey L, et al. Pilot study of the effect of methyl B12 treatment on behavioral and biomarker measures in children with autism. *J Altern Complement Med*. 2010, 16(5):555-560.
117. Cheuk DKL, Wong V, Chen WX. Acupuncture for autism spectrum disorders (ASD). *Cochrane Database Syst Rev*. 2011, (9):CD007849.
118. Ghanizadeh A. Hyperbaric oxygen therapy for treatment of children with autism: a systematic review of randomized trials. *Med Gas Res*. 2012, 2:13.
119. Magiati I, Moss J, Yates R, et al. Is the Autism Treatment Evaluation Checklist (ATEC) a useful tool for monitoring progress in children with Autism Spectrum Disorders? *J Intellect Disabil Res*. 2011, 55(3):302-312.
120. Taylor JL, Seltzer MM. Employment and post-secondary educational activities for young adults with autism spectrum disorders during the transition to adulthood. *J Autism Dev Disord*. 2011, 41(5):566-574.
121. Taylor JL, McPheeters ML, Sathe NA, et al. A systematic review of vocational interventions for young adults with autism spectrum disorders. *Pediatrics*. 2012, 130(3):531-538.
122. Le Couteur A, Haden G, Hammal D, et al. Diagnosing autism spectrum disorders in pre-school children using two standardised assessment instruments: the ADI-R and the ADOS. *J Autism Dev Disord*. 2008, 38(2):362-367.
123. Maljaars J, Noens I, Scholte E, et al. Evaluation of the criterion and convergent validity of the Diagnostic Interview for Social and Communication Disorders in young and low-functioning children. *Autism*, 2012, 16(5):487-497.
124. Chlebowski C, Green JA, Barton ML, et al. Using the childhood autism rating scale to diagnose autism spectrum disorders. *J Autism Dev Disord*. 2010, 40(7):787-799.
125. Molloy CA, Murray DS, Akers R, et al. Use of the Autism Diagnostic Observation Schedule (ADOS) in a clinical setting. *Autism*. 2011, 15(2):143-162.
126. Lord C, Risi S, DiLavore PS, et al. Autism from 2 to 9 years of age. *Arch Gen Psychiatry*. 2006, 63(6):694-701.

## Examples of Search Strategy

The following MeSH terms or free text terms were used either singly or in combination, search was limited to English, human, 2001 to current and children 0 -18 years old

**TCM**

1. Autistic Disorder/
2. (Autistic adj1 disorder\*).tw.
3. Autism\*.tw.
4. 1 or 2 or 3
5. Vitamin B 12/
6. (b12 adj1 vitamin).tw.
7. cyanocobalamin.tw.
8. cobalamin.tw.
9. 5 or 6 or 7 or 8
10. 4 and 9
11. Vitamin B 6/
12. Vitamin B6.tw.
13. Vitamin B 6.tw.
14. Pyridoxine.tw.
15. Pyridoxamine.tw.
16. Picolines.tw.
17. /or 11-16
18. Vitamin A/
19. vitamin a.tw.
20. vitamin a1.tw.
21. all trans retinol.tw.
22. Retinol.tw.
23. 11-cis-retinol.tw.
24. /or 18 -23
25. Fish Oils/
26. (fish adj1 oil\*).tw.
27. liver oils fish.tw.
28. fish liver oils.tw.
29. oils fish liver.tw.
30. /or 25-29
31. Zinc/
32. Zinc.tw.
33. 31 or 32
34. Melatonin/
35. Melatonin.tw.
36. 34 or 35
37. 4 and 17
38. 4 and 24
39. 4 and 30
40. 4 and 33
41. 4 and 36
42. trimethylglycine.mp.
43. Trimethylglycine.tw.
44. Magnesium/
45. Magnesium.tw.
46. 42 or 43 or 44 or 45
47. 4 and 46
48. Hyperbaric Oxygenation/
49. therapies hyperbaric oxygen.tw.
50. hyperbaric oxygenation\*.tw.
51. hyperbaric oxygen therap\*.tw.
52. oxygen therapies hyperbaric.tw.
53. oxygenation hyperbaric.tw.
54. therap\* hyperbaric oxygen.tw.
55. oxygen therap\* hyperbaric.tw.
56. HBO.t.w.
57. /or 48 - 56
58. 4 and 57
59. Ascorbic Acid/
60. (acid adj1 ascorbic).tw.
61. (sodium adj1 ascorbate).tw.
62. magnorbin.tw.
63. (ferrous adj1 ascorbate).tw.
64. acid l-ascorbic.tw.
65. magnesium ascorbicum.tw.
66. l-ascorbic acid.tw.

67. vitamin c.tw.
68. l ascorbic acid.tw.
69. ascorbic acid monosodium salt.tw.
70. hydrin.tw.
71. ascorbic acid monosodium salt.tw.
72. hydrin.tw.
73. /or59 -72
74. 4 and 73
75. Chelation Therapy/
76. (Chelation adj 1 therap\*).tw.
77. 75 or 76
78. 4 and 77

**Speech Language Therapy**

1. Autistic Disorder/
2. autism\*.tw.
3. (Autistic adj1 disorder\*).tw.
4. 1 or 2 or 3
5. Speech Therapy/
6. Language Therapy/
7. Speech language therap\*.tw.
8. (Speech adj1 therap\*).tw.
9. (Language adj1 therap\*).tw.
10. (Language adj1 train\*).tw.
11. 5 or 6 or 7 or 8 or 9 or 10
12. 4 and 11
13. Augmentative communication\*.tw.
14. Alternative communication\*.tw.
15. Augmentative alternative communication.tw.
16. AAC.tw.
17. /or 13-16
18. 4 and 17
19. Communication Aids for Disabled/
20. Augmentative communication\*.tw.
21. Alternative communication\*.tw.
22. Augmentative alternative communication.tw.
23. AAC.tw.
24. AAC intervention\*.tw.
25. /or 19-24
26. 4 and 25
27. Picture Exchange Communication System\*.tw.
28. PECS.tw.
29. 27 or 28
30. 4 and 29
31. naturalistic behavior?\*.tw.
32. natural language teach\*.tw.
33. 31 or 32
34. 4 and 33
35. milieu therap\*.tw.
36. milieu teach\*.tw.
37. 35 or 36
38. 4 and 37
39. incidental teach\*.tw.
40. 4 and 39
41. facilitate teach\*.tw.
42. facilitate communication\*.tw.
43. facilitated communication\*.tw.
44. 41 or 42 or 43
45. 4 and 44
46. discrete trial train\*.tw.
47. discrete trial teach\*.tw.
48. discrete trial instruct\*.tw.
49. discrete trial intervention\*.tw.
50. 46 or 47 or 48 or 49
51. 4 and 50

52. Video instruct\*.tw.
53. Video model\*.tw.
54. Video self-model\*.tw.
55. Point-of-view video\*.tw.
56. /or 52- 55
57. 4 and 56

**Behaviour intervention**

1. Autism\*.tw.
2. (Autistic adj1 disorder\*).tw.
3. (Conditioning adj1 therap\*).tw.
4. Behavior?r intervention\*.tw.
5. (Behavior?r adj1 modification\*).tw.
6. (Therap\* adj1 behavior?r\*).tw.
7. Behavior Therapy/
8. Autistic Disorder/
9. Applied behavior analysis.tw.
10. 1 or 2 or 8
11. 3 or 4 or 5 or 6 or 7 or 9
12. 10 and 11

**Occupational therapy**

1. Autistic Disorder/
2. Autism\*.tw.
3. (autistic adj1 disorder\*).tw.
4. 1 or 2 or 3
5. Sensory processing\*.tw.
6. 4 and 5
7. Fine motor.mp.
8. Occupational Therapy/
9. (occupational adj1 therap\*).tw.
10. 7 or 8 or 9
11. 4 and 10
12. Toilet training\*.tw.
13. (toilet \* adj1 training\*).tw.
14. 12 or 13
15. 4 and 14

**Diet intervention**

1. Autistic Disorder/
2. Autism\*.tw.
3. (Autistic adj1 disorder\*).tw.
4. Diet\*.tw.
5. 1 or 2 or 3
6. 4 and 5

## Clinical Questions

1. **Risk Factors**
  - What are the risk factors for ASD?
2. **Comorbidities**
  - What are the common comorbidities of ASD?
3. **Screening**
  - What are the effective screening tools for diagnosis or to detect ASD?
4. **Assessment and Diagnosis**
  - What is the initial assessment – history, developmental history, physical examination?
  - Is ADOS accurate in diagnosing ASD?
  - Are DISCO, CARS and ADI-R useful for diagnosing ASD?
5. **Investigation**
  - What are the effective hearing tests which can be used in children with ASD to detect deafness/ hearing loss?
  - Is mineral analysis a useful investigation tool in children with ASD?
  - Do children with ASD need to be assessed for food allergies?
  - Do children with ASD need routine EEG?
  - What is the role of neuroimaging, metabolic and genetic testing in the diagnosis of ASD?
6. **Treatment**
  - Is speech-language therapy effective in treating children with ASD?
  - Is social stories effective in treating children with ASD?
  - Is behavioural intervention effective in treating ASD?
  - Is occupational therapy effective for children with ASD?
  - Is parental education intervention effective in treating children with ASD?
  - Is cognitive behavioural intervention effective in treating children with ASD?
  - Is TEACCH an effective and safe intervention for children with ASD?
  - Is dietary intervention effective in children with ASD?
  - Is pharmacological therapy (antipsychotics, SSRI, TCA, methylphenidate, divalproate sodium) effective and safe for children with ASD?
  - Is TCM (chelation, secretin, fatty acids, melatonin, vitamin B12, HBOT, vitamin B6- Mg) safe and efficacious treatment for ASD?
7. **Monitoring and Preparation of Transition to Adult services?**
  - What are the services available for children with ASD in the transition into adulthood?

### Modified Checklist for Autism in Toddlers (M-CHAT)

Name:		Rated by: Mother / Father / Carer / Nurse / Doctor
Date of Birth:	Age:	Date of Evaluation:
IC/MIC:		Location (Clinic):

#### Notes:

- Source: Robins D, Fein D, Barton M, Green JA. *The Modified Checklist for Autism in Toddlers (M-CHAT): an initial investigation in the early detection of autism and pervasive developmental disorders.* *J Autism Dev Disord.* 2001, 31:131–144.
- Translated into Malay by Ministry of Health for use in screening
- Valid for children aged 16 - 30 months
- Clinical use should proceed with caution, given that the current scoring system is designed to maximize sensitivity (i.e., identify as many children with autism spectrum disorders (ASD) as possible, which results in a number of false positive cases).
- The M-CHAT is not designed to detect all possible developmental disorders. Any parent who has concerns about their child should see their child's doctor regardless on the child's score on the M-CHAT.

#### Instructions:

Sila jawab soalan berikut (item 1 - 23). Cuba menjawab semua soalan. Sekiranya anda mengalami masalah untuk menjawab mana-mana soalan, sila rujuk kepada jururawat.

*Please fill out the following (item 1-23) about how your child usually is. Please try to answer every question. Contact the Nurses shall you have any queries. If the behaviour is rare (e.g., you have seen it once or twice), please answer as if the child does not do it.*

**Jawab SEMUA soalan. Bulatkan jawapan yang sesuai.  
Answer ALL questions. Circle the appropriate answer.**

1.	Adakah anak anda seronok apabila ditimang, dibuai atau dihentut atas kaki / paha dan sebagainya? <i>Does your child enjoy being swung, bounced on your knee, etc?</i>	Ya / Yes	Tidak / No
2.	<b>Adakah anak anda menunjukkan minat terhadap kanak-kanak lain? (contohnya bergaul, bermain, berkawan)</b> <i>Does your child take an interest in other children?</i>	<b>Ya / Yes</b>	<b>Tidak / No</b>
3.	Adakah anak anda suka memanjat, contohnya tangga, kerusi, meja dan lain-lain? <i>Does your child like climbing on things, such as up stairs?</i>	Ya / Yes	Tidak / No
4.	Adakah anak anda seronok bermain “cak-cak” atau main sorok-sorok? <i>Does your child enjoy playing peek-a-boo or hide and seek?</i>	Ya / Yes	Tidak / No
5.	Adakah anak anda pernah bermain-main olok-olok / berlakon, contohnya menelefon, bermain anak patung atau bermain masak-masak dan sebagainya? <i>Does your child ever pretend for example to talk on the phone or take care of dolls or pretend other things?</i>	Ya / Yes	Tidak / No
6.	Adakah anak anda pernah menunjuk menggunakan jari telunjuk untuk meminta sesuatu? <i>Does your child ever use his / her index finger to point, to ask for something?</i>	Ya / Yes	Tidak / No
7.	<b>Adakah anak anda pernah menunjuk / menggunakan jari telunjuk terhadap sesuatu yang menarik minatnya?</b> <i>Does your child ever use his / her index finger to point, to indicate interest in something?</i>	<b>Ya / Yes</b>	<b>Tidak / No</b>

8.	Bolehkah anak anda bermain dengan alat permainan yang kecil dengan betul, selain dari memasukkannya ke dalam mulut, membelek-belek atau menjatuhkan permainan itu? (contohnya kiub, kereta kecil, dan lain-lain) <i>Can your child play properly with small toys without just mouthing, fiddling or dropping them?</i>	Ya / Yes	Tidak / No
9.	<b>Pernakah anak anda membawa objek / benda dan menunjukkannya kepada anda?</b> <b><i>Does your child ever bring objects over to you (parent) to show you something?</i></b>	<b>Ya / Yes</b>	<b>Tidak / No</b>
10.	Adakah anak anda bertentang mata dengan anda lebih daripada dua saat? <i>Does your child look you in the eye for more than a second or two?</i>	Ya / Yes	Tidak / No
11.	Pernakah anak anda kelihatan seperti sangat sensitif / terganggu terhadap bunyi bising (contohnya: menutup telinga)? <i>Does your child ever seem oversensitive to noise? (e.g. plugging ears)</i>	Ya / Yes	Tidak / No
12.	Adakah anak anda senyum bila melihat anda atau membalas senyuman anda? <i>Does your child smile in response to your face or your smile?</i>	Ya / Yes	Tidak / No
13.	Adakah anak anda meniru perlakuan anda (contohnya meniru mimik muka anda dan sebagainya)? <b><i>Does your child imitate you? (e.g. if you make a face will your child imitate it?)</i></b>	<b>Ya / Yes</b>	<b>Tidak / No</b>
14.	Adakah anak anda bertindak balas apabila namanya dipanggil? <b><i>Does your child respond to his / her name when you call?</i></b>	<b>Ya / Yes</b>	<b>Tidak / No</b>
15.	Sekiranya anda menunjuk pada alat permainan yang jauh dari anda, adakah anak anda akan melihat kepada alat permainan tersebut? <b><i>If you point at a toy across the room, does your child look at it?</i></b>	<b>Ya / Yes</b>	<b>Tidak / No</b>

16.	Bolehkah anak anda berjalan? <i>Does your child walk?</i>	Ya / Yes	Tidak / No
17.	Adakah anak anda akan melihat pada benda yang sedang anda lihat? <i>Does your child look at things you are looking at?</i>	Ya / Yes	Tidak / No
18.	Adakah anak anda membuat pergerakan jari yang ganjil / pelik dekat mukanya? <i>Does your child make unusual finger movements near his / her face?</i>	Ya / Yes	Tidak / No
19.	Adakah anak anda cuba menarik perhatian anda terhadap aktiviti yang dilakukannya? <i>Does your child try to attract your attention to his / her own activity?</i>	Ya / Yes	Tidak / No
20.	Pernahkah anda terfikir bahawa anak anda ada masalah pendengaran? <i>Have you ever wondered if your child is deaf?</i>	Ya / Yes	Tidak / No
21.	Adakah anak anda dapat memahami percakapan orang? <i>Does your child understand what people say?</i>	Ya / Yes	Tidak / No
22.	Adakah anak anda kadang-kala kelihatan termenung atau merayau / berjalan tanpa tujuan? <i>Does your child sometimes stare at nothing or wander with no purpose?</i>	Ya / Yes	Tidak / No
23.	Adakah anak anda memandangi ke muka anda untuk melihat reaksi / tindakbalas anda apabila ia menghadapi sesuatu yang baru atau luar biasa? <i>Does your child look at your face to check your reaction when faced with something unfamiliar?</i>	Ya / Yes	Tidak / No

### **Outcome of M-CHAT**

Critical items failed: \_\_\_\_\_ Total items failed: \_\_\_\_\_

M-CHAT: Passed / Failed

1. Tidak / No	6. Tidak / No	11. Ya / Yes	16. Tidak / No	21. Tidak / No
<b>2. Tidak / No</b>	<b>7. Tidak / No</b>	12. Tidak / No	17. Tidak / No	22. Ya / Yes
3. Tidak / No	8. Tidak / No	<b>13. Tidak / No</b>	18. Ya / Yes	23. Tidak / No
4. Tidak / No	<b>9. Tidak / No</b>	<b>14. Tidak / No</b>	19. Tidak / No	
5. Tidak / No	10. Tidak / No	<b>15. Tidak / No</b>	20. Ya / Yes	

**Scoring:**

- *The bold items are critical: i.e. **2, 7, 9, 13, 14, 15***
- *A child requires referral (i.e. fail M-CHAT) for further evaluation if he / she fulfills the following:*
  - *2 or more of critical items*
  - *3 or more of any items*

***Not all children who fail the checklist will meet criteria for a diagnosis on the autism spectrum disorder (ASD). However children who fail the checklist should be evaluated in more depth by the relevant specialist.***

**Diagnostic Criteria for Autistic Disorder in the Diagnostic and  
Statistical Manual of Mental Disorders, Fourth Edition, Text Revision  
(DSM-IV-TR)**

The essential features of Autistic Disorder are the presence of markedly abnormal or impaired development in social interaction and communication and a markedly restricted repertoire of activity and interests. Manifestations of the disorder vary greatly depending on the developmental level and chronological age of the individual. Autistic Disorder is sometimes referred to as *early infantile autism*, *childhood autism*, or *Kanner's autism*.

- A. A total of six (or more) items from 1, 2, and 3, with at least two from 1, and one each from 2 and 3:
1. Qualitative impairment in social interaction, as manifested by at least two of the following:
    - a. marked impairment in the use of multiple nonverbal behaviours such as eye-to-eye gaze, facial expression, body postures, and gestures to regulate social interaction
    - b. failure to develop peer relationships appropriate to developmental level
    - c. a lack of spontaneous seeking to share enjoyment, interests, or achievements with other people (for example, by a lack of showing, bringing, or pointing out objects of interest)
    - d. lack of social or emotional reciprocity
  2. Qualitative impairments in communication as manifested by at least one of the following:
    - a. delay in or total lack of, the development of spoken language (not accompanied by an attempt to compensate through alternative modes of communication such as gestures or mime)
    - b. in individuals with adequate speech, marked impairment in the ability to initiate or sustain a conversation with others
    - c. stereotyped and repetitive use of language or idiosyncratic language
    - d. lack of varied, spontaneous make-believe play or social imitative play appropriate to developmental level
  3. Restricted, repetitive, and stereotyped patterns of behaviour, interests, and activities, as manifested by at least one of the following:
    - a. encompassing preoccupation with one or more stereotyped and restricted patterns of interest that is abnormal either in intensity or focus
    - b. apparently inflexible adherence to specific, non-functional routines or rituals

- c. stereotyped and repetitive motor mannerisms (for example, hand or finger flapping or twisting, or complex whole-body movements)
  - d. persistent preoccupation with parts of objects
- B. Delays or abnormal functioning in at least one of the following areas, with onset prior to age 3 years: (1) social interaction, (2) language as used in social communication, or (3) symbolic or imaginative play.
- C. The disturbance is not better accounted for by Rett's Disorder or Childhood Disintegrative Disorder.

(Adapted from the Diagnostic and Statistical Manual of Mental Disorder, Fourth Edition, Text Revision: DSM-IV-TR. Washington D.C.: American Psychiatric Association; 2000)

**Diagnostic Criteria for Autism Spectrum Disorder in the Diagnostic and Statistical Manual of Mental Disorder, Fifth Edition (DSM-5)**

Autism Spectrum Disorder 299.00 (F84.0)

Diagnostic Criteria

- A. Persistent deficits in social communication and social interaction across multiple contexts, as manifested by the following, currently or by history (examples are illustrative, not exhaustive; see text):
1. Deficits in social-emotional reciprocity, ranging, for example, from abnormal social approach and failure of normal back-and-forth conversation; to reduced sharing of interests, emotions, or affect; to failure to initiate or respond to social interactions.
  2. Deficits in nonverbal communicative behaviours used for social interaction, ranging, for example, from poorly integrated verbal and nonverbal communication; to abnormalities in eye contact and body language or deficits in understanding and use of gestures: to a total lack of facial expressions and nonverbal communication.
  3. Deficits in developing, maintaining, and understanding relationships, ranging, for example, from difficulties adjusting behaviour to suit various social contexts; to difficulties in sharing imaginative play or in making friends; to absence of interest in peers.

*Specify* current severity: Severity is based on social communication impairments and restricted, repetitive patterns of behaviour (refer to table below).

- B. Restricted, repetitive patterns of behaviour, interests, or activities, as manifested by at least two of the following, currently or by history (examples are illustrative, not exhaustive; see text):
1. Stereotyped or repetitive motor movements, use of objects, or speech (e.g., simple motor stereotypies, lining up toys or flipping objects, echolalia, idiosyncratic phrases).
  2. Insistence on sameness, inflexible adherence to routines, or ritualized patterns of verbal or nonverbal behaviour (e.g., extreme distress at small changes, difficulties with transitions, rigid thinking patterns, greeting rituals, need to take same route or eat same food every day).
  3. Highly restricted, fixated interests that are abnormal in intensity or focus (e.g., strong attachment to or preoccupation with unusual objects, excessively circumscribed or perseverative interests).

4. Hyper- or hypo-reactivity to sensory input or unusual interest in sensory aspects of the environment (e.g., apparent indifference to pain / temperature, adverse response to specific sounds or textures, excessive smelling or touching of objects, visual fascination with lights or movement).

*Specify* current severity: Severity is based on social communication impairments and restricted, repetitive patterns of behaviour (refer to table below).

- C. Symptoms must be present in the early developmental period (but may not become fully manifest until social demands exceed limited capacities, or may be masked by learned strategies in later life).
- D. Symptoms cause clinically significant impairment in social, occupational, or other important areas of current functioning.
- E. These disturbances are not better explained by intellectual disability (intellectual developmental disorder) or global developmental delay. Intellectual disability and autism spectrum disorder frequently co-occur; to make comorbid diagnoses of autism spectrum disorder and intellectual disability, social communication should be below that expected for general developmental level.

Note: Individuals with a well-established DSM-IV diagnosis of autistic disorder, Asperger's disorder, or pervasive developmental disorder not otherwise specified should be given the diagnosis of autism spectrum disorder. Individuals who have marked deficits in social communication, but whose symptoms do not otherwise meet criteria for autism spectrum disorder, should be evaluated for social (pragmatic) communication disorder.

Specify if;

**With or without accompanying intellectual impairment**

**With or without accompanying language impairment**

**Associated with a known medical or genetic condition or environmental factor (Coding note: Use additional code to identify the associated medical or genetic condition.)**

**Associated with another neurodevelopmental, mental, or behavioural disorder (Coding note: Use additional code[s] to identify the associated neuro-developmental, mental, or behavioural disorder[s].)**

**With catatonia** (refer to the criteria for catatonia associated with another mental disorder, pp. 119-120, for definition) (Coding note: Use additional code 293.89 [F06.1] catatonia associated with autism spectrum disorder to indicate the presence of the comorbid catatonia).

### Severity levels for autism spectrum disorder

Severity level	Social communication	Restricted, repetitive behaviours
<b>Level 3</b>  "Requiring very substantial support"	Severe deficits in verbal and nonverbal social communication skills cause severe impairments in functioning, very limited initiation of social interactions, and minimal response to social overtures from others. For example, a person with few words of intelligible speech who rarely initiates interaction and, when he or she does, makes unusual approaches to meet needs only and responds to only very direct social approaches	Inflexibility of behaviour, extreme difficulty coping with change, or other restricted/repetitive behaviours markedly interfere with functioning in all spheres. Great distress/difficulty changing focus or action.

<p><b>Level 2</b></p> <p>“Requiring substantial support”</p>	<p>Marked deficits in verbal and nonverbal social communication skills; social impairments apparent even with supports in place; limited initiation of social interactions; and reduced or abnormal responses to social overtures from others. For example, a person who speaks simple sentences, whose interaction is limited to narrow special interests, and who has markedly odd nonverbal communication.</p>	<p>Inflexibility of behaviour, difficulty coping with change, or other restricted/ repetitive behaviours appear frequently enough to be obvious to the casual observer and interfere with functioning in a variety of contexts. Distress and/or difficulty changing focus or action.</p>
<p><b>Level 1</b></p> <p>“Requiring support”</p>	<p>Without supports in place, deficits in social communication cause noticeable impairments. Difficulty initiating social interactions, and clear examples of atypical or unsuccessful responses to social overtures of others. May appear to have decreased interest in social interactions. For example, a person who is able to speak in full sentences and engages in communication but whose to-and-fro conversation with others fails, and whose attempts to make friends are odd and typically unsuccessful.</p>	<p>Inflexibility of behaviour causes significant interference with functioning in one or more contexts. Difficulty switching between activities. Problems of organization and planning hamper independence.</p>

(Adapted from the Diagnostic and Statistical Manual of Mental Disorder, Fifth Edition: DSM-5. Washington D.C.: American Psychiatric Association; 2014)

**International Classification of Diseases (ICD) 10****F84 Pervasive Developmental Disorders****F84.0 Childhood autism**

- A. Presence of abnormal or impaired development before the age of three years, in at least one out of the following areas:
1. receptive or expressive language as used in social communication;
  2. the development of selective social attachments or of reciprocal social interaction;
  3. functional or symbolic play.
- B. Qualitative abnormalities in reciprocal social interaction, manifest in at least one of the following areas:
1. failure adequately to use eye-to-eye gaze, facial expression, body posture and gesture to regulate social interaction;
  2. failure to develop (in a manner appropriate to mental age, and despite ample opportunities) peer relationships that involve a mutual sharing of interests, activities and emotions;
  3. A lack of socio-emotional reciprocity as shown by an impaired or deviant response to other people's emotions; or lack of modulation of behaviour according to social context, or a weak integration of social, emotional and communicative behaviours.
- C. Qualitative abnormalities in communication, manifest in at least two of the following areas:
1. a delay in, or total lack of development of spoken language that is not accompanied by an attempt to compensate through the use of gesture or mime as alternative modes of communication (often preceded by a lack of communicative babbling);
  2. relative failure to initiate or sustain conversational interchange (at whatever level of language skills are present) in which there is reciprocal to and from responsiveness to the communications of the other person;
  3. stereotyped and repetitive use of language or idiosyncratic use of words or phrases;
  4. abnormalities in pitch, stress, rate, rhythm and intonation of speech.
- D. Restricted, repetitive, and stereotyped patterns of behaviour, interests and activities, manifest in at least two of the following areas:
1. an encompassing preoccupation with one or more stereotyped and restricted patterns of interest that are abnormal in content or focus; or one or more interests that are abnormal in their intensity and circumscribed nature although not abnormal in their content or focus;

2. apparently compulsive adherence to specific, non-functional, routines or rituals;
  3. stereotyped and repetitive motor mannerisms that involve either hand or finger flapping or twisting, or complex whole body movements;
  4. preoccupations with part-objects or non-functional elements of play materials (such as their odour, the feel of their surface, or the noise or vibration that they generate);
  5. distress over changes in small, non-functional, details of the environment.
- E. The clinical picture is not attributable to the other varieties of pervasive developmental disorder; specific developmental disorder of receptive language (F80.2) with secondary socio-emotional problems; reactive attachment disorder (F94.1) or disinhibited attachment disorder (F94.2); mental retardation (F70 - F72) with some associated emotional or behavioural disorder; schizophrenia (F20) of unusually early onset; and Rett's syndrome (F84.2).

#### **F84.1 Atypical autism**

- A. Presence of abnormal or impaired development at or after age three years (criteria as for autism except for age of manifestation).
- B. Qualitative abnormalities in reciprocal social interaction or in communication, or restricted, repetitive and stereotyped patterns of behaviour, interests and activities (criteria as for autism except that it is not necessary to meet the criteria in terms of number of areas of abnormality).
- C. The disorder does not meet the diagnostic criteria for autism (F84.0). Autism may be atypical in either age of onset (F84.11) or phenomenology (84.12), these two types being differentiated with a fifth character for research purposes. Syndromes that are atypical in both respects should be coded F84.12.

#### **F84.5 Asperger's syndrome**

- A. A lack of any clinically significant general delay in spoken or receptive language or cognitive development.

Diagnosis requires that single words should have developed by two years of age or earlier and that communicative phrases be used by three years of age or earlier. Self-help skills, adaptive behaviour and curiosity about the environment during the first three years should be at a level consistent with normal intellectual development. However, motor milestones may be somewhat delayed and motor clumsiness is

usual (although not a necessary diagnostic feature). Isolated special skills, often related to abnormal preoccupations, are common, but are not required for diagnosis.

- B. Qualitative abnormalities in reciprocal social interaction (criteria as for autism).
- C. An unusually intense circumscribed interest or restricted, repetitive, and stereotyped patterns of behaviour, interests and activities (criteria as for autism; however it would be less usual for these to include either motor mannerisms or preoccupations with part- objects or non-functional elements of play materials).
- D. The disorder is not attributable to the other varieties of pervasive developmental disorder; schizotypal disorder (F21); simple schizophrenia (F20.6); reactive and disinhibited attachment disorder of childhood (F94.1 and .2); obsessional personality disorder (F60.5); obsessive-compulsive disorder (F42).

(Adapted from the ICD-10 Classification of Mental and Behavioural Disorders: Diagnostic Criteria for Research. Geneva: World Health Organization; 1993)

## Signs and symptoms of possible autism in preschool children (or equivalent mental age)

<b>Social interaction and reciprocal communication behaviours</b>
<p><i>Spoken language</i></p> <ul style="list-style-type: none"> <li>• Language delay (in babble or words, for example less than ten words by the age of 2 years)</li> <li>• Regression in or loss of use of speech</li> <li>• Spoken language (if present) may include unusual:</li> <li>• non-speech like vocalisations               <ul style="list-style-type: none"> <li>• odd or flat intonation</li> <li>• frequent repetition of set words and phrases ('echolalia')</li> <li>• reference to self by name* or "you" or "she / he" beyond 3 years</li> <li>• Reduced and/or infrequent use of language for communication, for example use of single words although able to speak in sentences</li> </ul> </li> </ul> <p>*Part of the spoken language section may not be applicable to local setting</p>
<p><i>Responding to others</i></p> <ul style="list-style-type: none"> <li>• Absent or delayed response to name being called, despite normal hearing</li> <li>• Reduced or absent responsive social smiling</li> <li>• Reduced or absent responsiveness to other people's facial expressions or feelings</li> <li>• Unusually negative response to the requests of others (demand avoidant behaviour)</li> <li>• Rejection of cuddles initiated by parent or carer, although may initiate cuddles themselves</li> </ul>
<p><i>Interacting with others</i></p> <ul style="list-style-type: none"> <li>• Reduced or absent awareness of personal space, or unusually intolerant of people entering their personal space</li> <li>• Reduced or absent social interest in others, including children of his/her own age – may reject others; if interested in others, may approach others inappropriately, seeming to be aggressive or disruptive</li> <li>• Reduced or absent imitation of others' actions</li> <li>• Reduced or absent initiation of social play with others, plays alone</li> <li>• Reduced or absent enjoyment of situations that most children like, for example, birthday parties</li> <li>• Reduced or absent sharing of enjoyment</li> </ul>

*Eye contact, pointing and other gestures*

- Reduced or absent use of gestures and facial expressions to communicate (although may place adult's hand on objects)
- Reduced and poorly integrated gestures, facial expressions, body orientation, eye contact (looking at people's eyes when speaking) and speech used in social communication
- Reduced or absent social use of eye contact assuming adequate vision
- Reduced or absent joint attention shown by lack of:
  - gaze switching
  - following a point (looking where the other person points to – may look at hand)
  - using pointing at or showing objects to share interest

*Ideas and imagination*

- Reduced or absent imagination and variety of pretend play

**Unusual or restricted interests and/or rigid and repetitive behaviours**

- Repetitive 'stereotypical' movements such as hand flapping, body rocking while standing, spinning, finger-flicking
- Repetitive or stereotyped play, for example opening and closing doors
- Over-focused or unusual interests
- Excessive insistence on following own agenda
- Extremes of emotional reactivity to change or new situations, insistence on things being 'the same'
- Over or under reaction to sensory stimuli, for example textures, sounds, smells
- Excessive reaction to taste, smell, texture or appearance of food or extreme food fads

## Signs and symptoms of possible autism in primary school children (aged 5 – 11 years or equivalent mental age)<sup>23</sup>

<p><b>Social interaction and reciprocal communication behaviours</b></p> <p><i>Spoken language</i></p> <ul style="list-style-type: none"> <li>● Spoken language may be unusual in several ways:             <ul style="list-style-type: none"> <li>● very limited use</li> <li>● monotonous tone</li> <li>● repetitive speech, frequent use of stereotyped (learnt) phrases, content dominated by excessive</li> <li>● information on topics of own interest</li> <li>● talking 'at' others rather than sharing a two-way conversation</li> <li>● responses to others can seem rude or inappropriate</li> </ul> </li> </ul>
<p><i>Responding to others</i></p> <ul style="list-style-type: none"> <li>● Reduced or absent response to other people's facial expressions or feelings</li> <li>● Reduced or delayed response to name being called, despite normal hearing</li> <li>● Subtle difficulties in understanding other's intentions; may take things literally and misunderstand sarcasm or metaphor</li> <li>● Unusually negative response to the requests of others (demand avoidant behaviour)</li> </ul>
<p><i>Interacting with others</i></p> <ul style="list-style-type: none"> <li>● Reduced or absent awareness of personal space, or unusually intolerant of people entering their personal space</li> <li>● Reduced or absent social interest in people, including children of his/her own age – may reject others; if interested in others, may approach others inappropriately, seeming to be aggressive or disruptive</li> <li>● Reduced or absent greeting and farewell behaviours</li> <li>● Reduced or absent awareness of socially expected behaviour</li> <li>● Reduced or absent ability to share in the social play or ideas of others, plays alone</li> <li>● Unable to adapt style of communication to social situations, for example may be overly formal or inappropriately familiar</li> <li>● Reduced or absent enjoyment of situations that most children like</li> </ul>
<p><i>Eye contact, pointing and other gestures</i></p> <ul style="list-style-type: none"> <li>● Reduced and poorly integrated gestures, facial expressions and body orientation, eye contact (looking at people's eyes when speaking), and speech used in social communication</li> <li>● Reduced or absent social use of eye contact assuming adequate vision</li> <li>● Reduced or absent joint attention shown by lack of:             <ul style="list-style-type: none"> <li>● gaze switching</li> <li>● following a point (looking where the other person points to – may look at hand)</li> <li>● using pointing at or showing objects to share interest</li> </ul> </li> </ul>

*Ideas and imagination*

- Reduced or absent flexible imaginative play or creativity, although scenes seen on visual media (for example, television) may be re-enacted
- Makes comments without awareness of social niceties or hierarch

**Unusual or restricted interests and/or rigid and repetitive behaviours**

- Repetitive “stereotypical” movements such as hand flapping, body rocking while standing, spinning, finger-flicking
- Play repetitive and oriented towards objects rather than people
- Over-focused or unusual interests
- Rigid expectation that other children should adhere to rules of play
- Excessive insistence on following own agenda
- Extremes of emotional reactivity that are excessive for the circumstances
- Strong preferences for familiar routines and things being ‘just right’
- Dislike of change, which often leads to anxiety or other forms of distress (including aggression)
- Over or under reaction to sensory stimuli, for example textures, sounds, smells
- Excessive reaction to taste, smell, texture or appearance of food or extreme food fads

**Other factors that may support a concern about autism**

- Unusual profile of skills or deficits (for example, social or motor coordination skills poorly developed, while particular areas of knowledge, reading or vocabulary skills are advanced for chronological or mental age)
- Social and emotional development more immature than other areas of development, excessive trusting (naivety), lack of common sense, less independent than peers

## Signs and symptoms of possible autism in secondary school children (older than 11 years or equivalent mental age)<sup>23</sup>

<b>Social interaction and reciprocal communication behaviours</b>
<p><i>Spoken language</i></p> <ul style="list-style-type: none"> <li>● Spoken language may be unusual in several ways:               <ul style="list-style-type: none"> <li>● very limited use</li> <li>● monotonous tone</li> <li>● repetitive speech, frequent use of stereotyped (learnt) phrases, content dominated by excessive</li> <li>● information on topics of own interest</li> <li>● talking “at” others rather than sharing a two-way conversation</li> <li>● responses to others can seem rude or inappropriate</li> </ul> </li> </ul>
<p><i>Interacting with others</i></p> <ul style="list-style-type: none"> <li>● Reduced or absent awareness of personal space, or unusually intolerant of people entering their personal space</li> <li>● Long-standing difficulties in reciprocal social communication and interaction: few close friends or reciprocal relationships</li> <li>● Reduced or absent understanding of friendship; often an unsuccessful desire to have friends (although may find it easier with adults or younger children),</li> <li>● Social isolation and apparent preference for aloneness</li> <li>● Reduced or absent greeting and farewell behaviours</li> <li>● Lack of awareness and understanding of socially expected behaviour</li> <li>● Problems losing at games, turn-taking and understanding ‘changing the “rules”</li> <li>● May appear unaware or uninterested in what other young people his or her age are interested in</li> <li>● Unable to adapt style of communication to social situations, for example may be overly formal or inappropriately familiar</li> <li>● Subtle difficulties in understanding other’s intentions; may take things literally and misunderstand sarcasm or metaphor</li> <li>● Makes comments without awareness of social niceties or hierarchies</li> <li>● Unusually negative response to the requests of others (demand avoidant behaviour)</li> </ul>
<p><i>Eye contact, pointing and other gestures</i></p> <ul style="list-style-type: none"> <li>● Poorly integrated gestures, facial expressions, body orientation, eye contact (looking at people’s eyes when speaking) assuming adequate vision, and spoken language used in social communication</li> </ul>
<p><i>Ideas and imagination</i></p> <ul style="list-style-type: none"> <li>● History of a lack of flexible social imaginative play and creativity, although scenes seen on visual media(for example, television) may be re-enacted</li> </ul>

- Unusual or restricted interests and/or rigid and repetitive behaviours
- Repetitive “stereotypical” movements such as hand flapping, body rocking while standing, spinning, finger-flicking
- Preference for highly specific interests or hobbies
- A strong adherence to rules or fairness that leads to argument
- Highly repetitive behaviours or rituals that negatively affect the young person’s daily activities
- Excessive emotional distress at what seems trivial to others, for example change in routine
- Dislike of change, which often leads to anxiety or other forms of distress including aggression
- Over or under reaction to sensory stimuli, for example textures, sounds, smells
- Excessive reaction to taste, smell, texture or appearance of food and/or extreme food fad

**Other factors that may support a concern about autism**

- Unusual profile of skills and deficits (for example, social or motor coordination skills poorly developed, while particular areas of knowledge, reading or vocabulary skills are advanced for chronological or mental age)
- Social and emotional development more immature than other areas of development, excessive trusting (naivety), lack of common sense, less independent than peers

## Diagnostic Tools

Diagnostic Tool	Description	Sensitivity	Specificity	Comments
<b>Autism Diagnostic Interview-Revised (ADI-R)</b> <small>122, level III</small>	<ul style="list-style-type: none"> <li>● Structured interview conducted with the parents/ carers of individuals for assessment of ASD.</li> <li>● It is used for diagnostic purposes for anyone with a mental age of at least 18 months.</li> <li>● The interview measures behaviour in the areas of reciprocal social interaction, communication and language, and patterns of behaviour</li> </ul>	1.0	> 0.97	<ul style="list-style-type: none"> <li>● The levels of agreement between the ADI-R and ADOS is moderate in social (k=0.56) and communication (k=0.48) algorithm cut-offs scores.</li> <li>● The level of agreement between the ADI-R and ADOS is substantial in diagnosing autism (k=0.62) and moderate for spectrum diagnosis (k=0.54).</li> </ul>
<b>Diagnostic Interview for Social and Communication Disorders (DISCO)</b> <small>123, level II-3</small>	<ul style="list-style-type: none"> <li>● Semi-structured interview used by the parents/ carers to identify the impairments of social interaction, social communication and social imagination together with the associated repetitive behaviours</li> </ul>	0.80	0.79	<ul style="list-style-type: none"> <li>● DISCO accurately identify ASD in young children with an average intelligence or mild intellectual disability</li> <li>● The agreement between DISCO algorithm and ADOS was substantial (k=0.69, p &lt; 0.001).</li> </ul>

Diagnostic Tool	Description	Sensitivity	Specificity	Comments
<p><b>Childhood Autism Rating Scale (CARS)</b> 124, level II-2</p>	<ul style="list-style-type: none"> <li>● Consists of 14 domains assessing behaviours associated with autism, with the 15th domain rating general impressions of autism.</li> <li>● Each domain is scored on a scale ranging from one to four; higher scores are associated with a higher level of impairment.</li> <li>● Scores below 30 indicate that the individual is in non-autistic range.</li> </ul>	<p>2 years old: 0.79</p> <p>4 years old: 0.86</p>	<p>2 years old: 0.81</p> <p>4 years old: 0.80</p>	<ul style="list-style-type: none"> <li>● In younger children, CARS is useful to distinguish ASD from pervasive developmental disorder not otherwise specified (PDD-NOS).</li> <li>● Using an ASD cut-off score of 25.5, CARS produced the highest level of agreement among diagnostic instruments and clinical judgement:</li> <li>● <b>2 years old:</b> <ul style="list-style-type: none"> <li>● CARS and DSM-IV, <math>k=0.75</math>, <math>p &lt; 0.001</math></li> <li>● CARS and ADOS, <math>k=0.70</math>, <math>p &lt; 0.001</math></li> </ul> </li> <li>● <b>4 years old:</b> <ul style="list-style-type: none"> <li>● CARS and DSM IV, <math>k=0.74</math>, <math>p &lt; 0.001</math></li> <li>● CARS and ADOS, <math>k=0.73</math>, <math>p &lt; 0.001</math></li> </ul> </li> </ul>

Diagnostic Tool	Description	Sensitivity	Specificity	Comments
<p><b>Autism Diagnostic Observation Schedule (ADOS)</b>  <small>125, level III, 126, level II-2</small></p>	<ul style="list-style-type: none"> <li>● Semi-structured assessment of communication, social interaction and play (or imaginative use of materials) for individuals suspected of having ASD.</li> <li>● Consists of four modules, each of which is appropriate for children and adults of differing developmental and language levels, ranging from nonverbal to verbally-fluent.</li> </ul>	0.82	0.60	<ul style="list-style-type: none"> <li>● The prevalence of best-estimate clinical diagnosis at nine years is 43% for ADOS alone, 82% when combined with clinical judgement and 90% when combined with both clinical judgement and ADI-R.</li> </ul>

## LIST OF ABBREVIATIONS

3Di	Developmental, Dimensional and Diagnostic Interview
ABA	Applied Behaviour Analysis
AAC	Augmentative and Alternative Communication
AAP	Atypical antipsychotics
ABR	Auditory Brainstem Evoked Response
ADHD	Attention Deficit Hyperactivity Disorder
ADI-R	Autism Diagnostic Interview- Revised
ADOS	The Autism Diagnostic Observation Schedule
AGREE	Appraisal of Guidelines for Research and Evaluation
AR	Acoustic Reflexes
ASD	Autism Spectrum Disorder
ASSQ	Autism Spectrum Screening Questionnaire
ASDS	Asperger Syndrome Diagnostic Scale
CARS	Childhood Autism Rating Scale
CBCL	Child Behaviour Checklist
CDSR	Cochrane Database of Systematic Reviews
CI	Confidence Interval
CGI	Clinical Global Impression
CHAT	Checklist for Autism in Toddlers
CHAT-23	Checklist for Autism in Toddlers for Chinese Children
CPG	Clinical Practice Guidelines
Db	Decibel
DCD	Developmental Coordination Disorder
DIR	Developmental, Individual-Difference, Relationship-Based
DISCO	the Diagnostic Interview for Social and Communication Disorders
DG	Development Group
DSM-IV-TR	Diagnostic and Statistical Manual of Mental Disorders, Fourth Edition, Text Revision
DSM-5	Diagnostic and Statistical Manual of Mental Disorders, Fifth Edition
EEG	Electroencephalography
ES	Effect Size
GARS	Gilliam Autism Rating Scale
GARS-2	Gilliam Autism Rating Scale Second Edition
G-I-N	Guidelines International Network
ICD-10	International Classification of Diseases
IQ	Intelligence Quotient
IRD	Improvement Rate Difference
MaHTAS	Malaysia Health Technology Assessment Section
M-CHAT	Modified Checklist for Autism in Toddlers
MD	Mean Difference

MDT	Multidisciplinary Team
MMR	Measles, Mumps, Rubella
MOH	Ministry of Health
OCD	Obsessive Compulsive Disorder
ODD	Oppositional Defiant Disorder
OT	Occupational Therapy
OR	Odd Ratio
P	p value
PDD	Pervasive Developmental Disorder
PDD-NOS	Pervasive Developmental Disorder Not Otherwise Specified
PECS	Picture Exchange Communication System
PND	Percentage of Non-overlapping Data
PPV	Positive Predictive Value
RC	Review Committee
RCT	Randomised Controlled Trial
RIT	Reciprocal Imitation Training
RPMT	Responsive Education and Prelinguistic Milieu Teaching
RR	Risk Ratio
SCI	Social Communication Intervention
SCQ	Social Communication Questionnaire
SMD	Standard Mean Difference
SIT	Sensory Integration Therapy
SR	Systematic Review
SRS	Social Responsiveness Scale
SSRI	Selective Serotonin Reuptake Inhibitors
SGD	Speech Generating Device
TEOAE	Transient Evoked Otoacoustic Emissions
TEACCH	Treatment and Education of Autistic and Related Communication Handicapped Children
TCA	Tricyclic Antidepressants
WMD	Weighted Mean Difference

## Acknowledgement

The members of the Development Group of the CPG would like to express their gratitude and appreciation to the following for their contributions:

- Panel of external reviewers who reviewed the draft
- Ms. Loong Ah Moi – Nursing officer / Information Specialist, MaHTAS, Medical Development Division, Ministry of Health
- Ms. Cheoh Siew Tin – Public Health Matron, who was involved in the early stage of development of the CPG
- Technical Advisory Committee for CPG for their valuable input and feedback
- All those who have contributed directly or indirectly to the development of the CPG
- Mr. Wong Siew Tung for proof reading

## Disclosure Statement

The panel members of both Development Group and Review Committee had completed disclosure forms. None held shares in pharmaceutical firms or acts as consultants to such firms. (Details are available upon request from the CPG Secretariat)

## Source of Funding

The development of the CPG on Management of Autism Spectrum Disorder was supported financially in its entirety by the Ministry of Health Malaysia.

Malaysian Health Technology Assessment Section (MaHTAS)  
Medical Development Division, Ministry of Health Malaysia  
Level 4, Block E1, Precinct 1  
Federal Government Administrative Centre  
62590, Putrajaya, Malaysia

ISBN 978-967-0769-01-1



9 789670 769011