ANTICIPATORY GUIDANCE FOR CHILDREN AND ADOLESCENTS WITH FETAL ALCOHOL SPECTRUM DISORDER (FASD): PRACTICE POINTS FOR PRIMARY HEALTH CARE PROVIDERS

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ABSTRACT

Background

Fetal Alcohol Spectrum Disorder (FASD) is an umbrella term that describes the range of effects that can occur in an individual who was prenatally exposed to alcohol and includes an array of complex neurodevelopmental and physical findings.

Objectives

To give primary healthcare providers (PHCP) evidence-based recommendations for supporting and managing the symptoms of FASD *after* patients have received a diagnosis.

Methods

Primary health recommendations for the management of children and adolescents with FASD were developed based on expert clinical judgment and supported by evidence-based research, where appropriate. The format was adapted from other health supervision practice guidelines as developed by the American Academy of Pediatrics. Clinical practice "Points" for the PHCP are highlighted. A reference table of anticipatory recommendations by age is presented.

Results

In most cases, the initial screening and referral for diagnosis will be made by the PHCP, and they will be responsible for ongoing management. It is anticipated that these recommendations will provide the PHCP with evidence to support the longitudinal health care of children and adolescents with FASD and their families as they transition throughout all developmental stages.

Conclusion

There is a pressing need for the involvement of PHCP in the active care of children and adolescents with FASD and their families over the lifespan. PHCP are trained in screening, prevention, and management of health needs, and are in the position to coordinate sub-specialty referrals as needed. Engaging PHCP will provide a truly integrated care system for individuals with FASD and their families.

Key Words: Fetal Alcohol Spectrum Disorder (FASD); children; adolescents; anticipatory guidance

What is FASD?

Fetal Alcohol Spectrum Disorder (FASD) is an umbrella term that describes the range of effects that can occur in an individual who was prenatally exposed to alcohol and includes an array of complex neurodevelopmental and physical

findings.¹ These effects can include lifelong physical, mental and behavioural difficulties, as well as learning disabilities. Alcohol exposure during pregnancy results in changes to the developing brain at both the neurochemical and structural levels. Often, these changes are not

detected until a child reaches early or middle school age when problems at school and at home become increasingly apparent. These challenges can include difficulties in adaptive functioning, social communication and attention, motor and sensory problems, memory deficits, and trouble learning from consequences. As the child grows and develops, they are at an increased risk for developing depression, anxiety and/or other mental health conditions, particularly if their symptoms and deficits are not appropriately identified and managed.

Scientific evidence has conclusively shown that alcohol consumption during pregnancy can cause fetal harm²⁻⁵; however, there is insufficient data to define any threshold for safe low-level drinking during pregnancy or when planning to become pregnant. Because the brain injury is the most common and serious consequence of prenatal exposure to alcohol and can occur at any time during a pregnancy, the safest choice for a woman who is pregnant or planning to become pregnant is **not to drink alcohol**.

It is possible for individuals with FASD to experience differences in their challenges due to the wide variation of alcohol-mediated effects on brain development. Some of the more common difficulties include:

- Cognitive functioning/learning disabilities.
- Executive functioning difficulty with judgment; planning; delaying gratification; inhibition; cognitive flexibility; learning from consequences; organization and impulsivity.
- Memory short and long term; verbal and visual information.
- Communication can be highly verbal, but lack comprehension skills for both the verbal and nonverbal cues; deficits in social communication (pragmatics) impacts daily adaptive and social function.
- Neuromotor Deficits impaired balance and coordination; and poor motor planning.
- Sensory Dysregulation hyper- or hyposensitivities or combinations of both to touch (e.g., low/high pain threshold, heat/cold); light; sound; taste and smell.

If these are not appropriately understood and addressed, children and adolescents with FASD are at increased risk for early school failure, involvement with the law, family disruption and homelessness.

In 2005, Loock and colleagues published "Identifying fetal alcohol spectrum disorder in primary care" as a guideline for diagnosis. These recommendations provided simple. straightforward approach to identifying patients at risk for FASD and outlined the appropriate screening tools for assessing problematic alcohol women. This reflected operationalization of the Canadian Diagnostic Guidelines for FASD¹ and aimed to improve diagnosis of FASD in the clinical setting. However, they did not include specific guidance for follow-up, monitoring and ongoing health support, in which the responsibility falls predominantly to the primary healthcare providers (PHCP). Thus, based on the expectation that PHCP can now screen women at-risk of having children with FASD and the availability of diagnostic services, the purpose of this paper is to provide PHCP with evidence-based anticipatory guidance for supporting and managing the symptoms of FASD after patients have received a diagnosis. These practice points support the nonpeer reviewed literature on medical management of FASD^{7,8}, and which can be reviewed for more in depth background.

In a survey conducted by the American Academy of Pediatrics, pediatricians reported knowledge about FASD, but did not feel adequately trained to integrate the management of the diagnosis in their everyday practice.⁹ Pediatricians and other PHCP are often required to coordinate appropriate mental health services, provide consultation to special education programs, and manage medication for comorbid health conditions. Thus, the goal of this document present best practice anticipatory recommendations that can be used by PHCP to better meet the challenges of ongoing management for patients with FASD after providing appropriate referrals for FASD diagnosis (please see Appendix I: Anticipatory Guidance for Children and Adolescents with FASD template).

Prevalence of FASD

Although exact prevalence rates for FASD remain largely unknown, from the estimates that do exist, it is clear that FASD is an important global health issue. In the United States, FASD has been estimated to occur at a rate of 9.1 per 1000 live births 10-13, and may be as high as 2-5% in some parts of the United States and in some Western European countries. 14 Currently, countries from Central and Eastern Europe, Africa, Asia and Canada are working with the World Health Organization to determine accurate prevalence rates for FASD. 15

It has been well-documented that alcohol consumption among women of child-bearing age is increasing ¹⁶ and that drinking patterns leading to adverse health outcomes are also increasing, including those associated with prenatal alcohol exposure. In the developed world, approximately 40% of women of childbearing age drink alcohol, and many are drinking at or around the time of conception. ^{17,18}

The economic and societal costs associated with FASD are significant. In Canada, the *total* adjusted annual cost associated with FASD at the individual level is an ~\$21,642¹⁹, with a lifetime cost from day of birth to 53 years old of ~\$5.3 billion. There are also the unmeasured indirect costs associated with stress and financial burden to the caregiver, as well as the loss of full potential for the individual with FASD. The costs associated with supporting individuals with FASD span many systems including education, child welfare, health, mental health, justice and social support (e.g., housing).

Speaking to Patients About Alcohol

PHCP are well positioned to counsel their patients about problematic alcohol use. They play an important role in providing education and information, as well as, communicating appropriate low-risk alcohol drinking guidelines. However, a recent US report revealed that health professionals do not routinely discuss problematic alcohol use (such as binge drinking) with their patients. PHCP may feel uncomfortable asking their patients about alcohol use; lack the time; receive inadequate remuneration; are influenced by issues of societal stigma; perceive professional

or systemic inadequacies; and/or lack information on how to manage a discussion on alcohol use and intervention strategies. 22,23 Recent meta-analytic evidence supports the effectiveness preconception interventions such as screening, education, motivational interviewing, and referral to reduce alcohol use in pregnancy.²⁴ These conversations need to be respectful, compassionate, and informed. PHCP thus have an important role in screening for problematic alcohol use, offering harm reduction, implementing brief intervention strategies, and/or providing referrals for diagnosis and treatment. 22,25

Screening for Problematic Alcohol Use

It is critical for PHCP to feel comfortable and confident screening for problematic alcohol use, especially during pregnancy. Currently, there are several screening tools available that are both sensitive and specific for both males and females. Screening tools for problematic alcohol use that are specific to women include the "CRAFFT" for adolescents, the "modified CAGE" (T-ACE and TWEAK) for pregnant and non-pregnant women 1,6 or the AUDIT (Alcohol Use Disorder Identification Test). Motivational interviewing is also an important skill set for conducting these types of investigations and is effective in reducing alcohol use in pregnancy.

FASD Screening Tool Kit

In partnership with the Public Health Agency of Canada (PHAC), the First Nations Inuit Health Branch (FNIHB) of Health Canada, and many FASD experts, practitioners and researchers, the Canadian Association of Pediatric Health Centres (CAPHC) facilitated the development of a National FASD Screening Tool Kit. 28,29 The tool kit is comprised of the Neurobehavioural Screening Tool (NST); Meconium Fatty Acid Ethyl Esters (FAEE) Testing; Maternal Drinking Guide – Fact Sheet and Tool: Medicine Wheel Student Index and Medicine Wheel Developmental History; and the FASD Screening & Referral Form for Youth Probation Officers. The use of these tools, such as the NST, continue to be evaluated across different clinical populations.³⁰ Ethical questions have been raised about biomarkers for prenatal alcohol exposure^{31,32}, which identify risk for developmental

disability in the child, but are not diagnostic. The child will need developmental monitoring and access to a supportive environment. These tests also extend a certain degree of culpability to the biological mother, and positive screens need to be carefully communicated³³ and the mother provided with supports for her own needs.^{33,34}

HOW IS THE DIAGNOSIS MADE?

Multidisciplinary Approach

FASD may be diagnosed in an individual with a history of prenatal alcohol exposure and a pattern of neurobehavioural characteristics with or without facial features. The Canadian diagnostic guidelines for FASD¹ describe a multidisciplinary process that consists of screening and referral, neurodevelopmental and physical assessment. diagnostic formulation, recommendations for management, and follow-up. The multidisciplinary team varies based on resources, but typically includes a coordinator, physician, psychologist, occupational therapist, speech therapist, and social worker. The team collects information from the individual and their family to determine the specific needs and goals of the assessment, to ascertain their readiness, and to identify community and cultural influences that may impact the assessment and subsequent resources available to the family. PHCP can provide important support, education, and advocacy to the family during the process of data gathering. Confirmation of prenatal alcohol exposure is a complex process that involves reliable sources, ideally the birth mother. The multidisciplinary assessment provides data on cognitive and adaptive functioning across multiple domains of brain function. A differential diagnosis is considered along with the physical examination and dysmorphology assessment to determine the functional strengths and needs that can inform the most appropriate management plan. The final report outlines the basis for diagnosis, details specific areas of strength and weakness, and provides concrete recommendations for follow-up and support. The PHCP receives a copy of the report and plays an important role in following-up on recommendations with the family, providing on-going education and support in the community, advocating for services and supports, and for a culture of acceptance and understanding.

METHODS

Primary health recommendations for the management of children with FASD were developed based on expert clinical judgment and supported by evidence-based research, where appropriate. The format was adapted from other health supervision practice guidelines as developed by the American Academy of Pediatrics (e.g., AAP Committee on Genetics. American Academy of Pediatrics: Health supervision for children with Down syndrome²⁶).

RESULTS & DISCUSSION

Key Points about FASD for PHCP Social and environmental factors

There are a number of potential factors that can influence the presentation and severity of FASD, including: maternal nutrition and stress in pregnancy, exposure to other drugs in utero, as well as, genetic and epigenetic influences. 5,35-39

Early postnatal adversity such as unstable home environment and exposure to neglect, abuse and domestic violence have a negative effect on early brain development. This is mediated largely through the stress response involving the hypothalamic pituitary axis leading to altered cortisol secretion and lifelong implications. 38,40 These effects have been well studied in the Adverse Childhood Experiences (ACE) study⁴¹ and in the Harvard Early Brain Development work. 42 Attachment disorders need also to be considered in children who have had chaotic early life experiences with multiple caregivers, as this can have a negative impact on development. 43-48 Many children with prenatal exposure to alcohol will come into the foster care system due to the systemic issues that contributed to the maternal drinking. These children are at risk for developmental disability from both the prenatal alcohol exposure and early life adversity. All children in foster care require developmental screening including the investigation of prenatal alcohol exposure. These should be evaluated by history and documented. 49,50 Ultimately, an early diagnosis, access to appropriate services, and stability of home environment are critical protective factors against negative outcomes in later years. 51,52

PRACTICE POINT #1: PHCP are in a position to screen for child maltreatment and to advocate for stability of placement for the child and education for their families to minimize or prevent further trauma. 53,54

Behaviour and Learning

Current research on the brain dysfunction associated with FASD supports a diffuse information processing, integration and selfregulation model that underlies many of the problems observed in this population. 55-58 Clinically, affected children present with complex difficulties in adaptive function that becomes increasingly evident with age as natural societal expectations increase. 59-61 The measure of intellectual ability alone often fails to reflect the extent of dysfunction associated with FASD and many children score in the low-average range.⁶² Executive functioning deficits describe significant areas of difficulty that include: cognitive flexibility, planning and organization, reasoning, inhibition, shifting, working memory and abstract reasoning. 63-65 Memory deficits in both verbal and visual information have also been documented. 55,66 Attention difficulties are common and studies support a comorbid attention deficit/hyperactivity disorder (ADHD) diagnosis in >65% of children with FASD. 67,68 Sustained attention appears to be more significantly impaired than immediate attention in children with FASD compared to children with ADHD.⁶⁹ Academic difficulties have been identified in the areas of mathematics and reading comprehension, both critical to functional numeracy of money, time management and learning through reading.^{70,71} In later school years, a shift from academics to basic life skills and employability options is more supportive and reduces the frustrations of being overwhelmed in a classroom setting.⁷²

Management of ADHD and Mood Disorders Comorbid with FASD

Educating caregivers on the complex impact of prenatal alcohol exposure on brain development is the initial step in the management of comorbid ADHD and mood dysregulation diagnoses.

Pharmacotherapeutics

The first line pharmacotherapeutics for ADHD are the stimulant medications (e.g., methylphenidate) in short or long acting formulations as described in the Multimodal Treatment of ADHD⁷³ and in the American Academy of Pediatrics guidelines for ADHD treatment.⁷⁴ Common side effects include loss of appetite and difficulty settling to sleep that requires ongoing monitoring of growth, nutrition planning and sleep management. Emotional lability can present as the medications wear off and is often a reason to discontinue the stimulant medication. Switching from one stimulant to another may result in fewer side effects. Consideration must be given if the adolescent or their family members are at risk of abuse, misuse or diverting the stimulant medication to a street drug when choosing to prescribe stimulants. If there is a family history of sudden death from cardiac arrhythmias or a known cardiac concern in the patient, a baseline electrocardiogram with measure of OT interval is recommended.

An alternative non-stimulant medication for ADHD management is Atomoxetine a selective norepinephrine transport inhibitor, which has been effective for the inattentive ADHD subtype and anxiety. This requires daily dosing and more than four weeks of administration to determine effectiveness. The most common side effects include nausea and abdominal discomfort.

Alpha 2 adrenergic agonists (e.g., short acting clonidine and long acting guanfacine, under the trade name of Intuniv), which facilitate dopamine and norepinephrine neurotransmission have been beneficial for the medical management of ADHD as an adjunct to stimulants or as a primary therapy. Cardiac, including electrocardiogram with QT interval and blood pressure monitoring is required for this family of

drugs. Side effects reported are initial sedation that improves over time.

Medication management for anxiety, depression and other mood disorders comorbid with FASD frequently involves the use of Selective Serotonin Reuptake Inhibitors or SSRIs. 78-80 However, there are no randomized control trials specific to FASD. Atypical antipsychotics are often prescribed for individuals with FASD for a variety of disruptive behaviours but there is no research evidence to support this. This raises an urgent call to researchers to do more in FASD intervention research, especially with the suspected prevalence of FASD in the population and the high incidence of comorbid mental health disorders. A recent panel of experts on therapeutic products for children in Canada led by Dr. Stuart MacLeod has raised these issues.

Dose guidelines are provided based on body weight with a "start low go slow" rule when changing doses based on clinical responses. It is essential to obtain information from caregivers and teachers on the responses in all environments by either a questionnaire or qualitative description. It is also important to ask the older child or youth how they feel the medication is working for them.

Interventions

Medications must be accompanied by classroom accommodations such as preferential seating, reducing over stimulation, reduced task length, extra time for tests, access to a scribe, reminders for organizational skills and modified learning if a learning disability is also present. Nonmedical approaches to ADHD management have been developed such as computer-based attention training and neurofeedback programs. However, these therapies are not readily available and can be very expensive and require a time commitment.

Parent coaching programs that teach how to provide positive discipline for negative behaviours and rewards for appropriate behaviours, such as the Triple P or Incredible Years programs have been beneficial for children and adolescents with ADHD and other disruptive behaviours. However, they have not been evaluated in the FASD population specifically but provide caregivers with basic parenting tools that are important. Anxiety and other internalizing

disorders can be managed with non-pharmaceutical interventions including behavioural parenting interventions or cognitive behavioural therapy for the affected child or youth.⁸⁴ These interventions require a skilled therapist who can modify the strategies according to the cognitive, communication and other challenges associated with FASD. However, it is unclear whether individuals with FASD can generalize from the therapy session to using the skills acquired in real life situations, as adaptive functioning is a fundamental deficit in FASD.

PRACTICE POINT #2: PHCP should provide ongoing monitoring of academic and social functioning with the family and education system. They are in a position to provide education and consultation to the schools and to interpret behaviour with an understanding of the neurological basis of FASD and the associated behaviours.

Self-Regulation

Difficulties with self-regulation have been considered a core deficit of FASD and describe problems with sustaining attention, difficulties with impulse control and inhibition, difficulty regulating appropriate responses to stress, mood and affect dysregulation, sleep disturbance and other mental health disorders. Disturbances in self-regulation can be noted as early as the period.85,86 neonatal and early infancy Presentations can include irritability, under-or over-responsiveness stimulation, to habituation, disorganized suck and swallow and disrupted sleep patterns. 87,88 Self-regulation problems can often be overlooked and further underscore the need for comprehensive history taking. During toddlerhood and into early school age, self-regulation difficulties present as hyperactivity, inattention, oppositional defiance and mood swings. ADHD may be diagnosed prior to identifying the history of prenatal alcohol exposure. This emphasizes the importance of asking about the possibility of alcohol use in the pregnancy when taking the history of any child presenting with developmental or behavioural disability. ⁶⁸ During adolescence and adulthood continued problems can lead to risk-taking and

impulsive behaviours that may lead to problems with the law. ^{60,61,72,89,90} When managing problems with self-regulation, environmental adaptation with an understanding of triggers is a good initial approach that may benefit from the involvement of an appropriate therapist. Medications that target the symptoms may be also considered along with a comprehensive behavioural management plan and appropriate supports. Medication management in FASD can be challenging and may require consultation with mental health services. ^{91,92}

PRACTICE POINT #3: PHCP should evaluate children with FASD for other mental health disorders (or refer for assessment). They should educate families on the behavioural and medication management when appropriate, in consultation with mental health specialists as needed.

Social Situations

While basic language may appear to be intact, higher order language deficits are common in the FASD population. 93 This includes communication functions such as predicting, referencing, inferencing, reasoning and narrative generation⁹⁴, which can impact social function, and the ability to understand instructions and information. 70,95-97 Children with FASD can be talkative, but tangential, making it difficult for the listener to follow what the individual is trying to say. This type of communication can present as "lying" or "telling stories", and must be considered when individuals are in different social situations (e.g., legal statements in court). Due to the fact that individuals with FASD often experience impairments in their understanding of social use of language and nonverbal social communication cues, they are victimization. 96,97,99 at an increased risk of

PRACTICE POINT #4: PHCP should screen communication skills of children with prenatal alcohol exposure and refer early to speech and language therapists.

Family Support

By framing FASD as a neurodevelopmental condition, the focus shifts from "changing" the

child to "providing" them with the most supportive environment. Stable home placement that includes positive stimulation leading to healthy attachments and learning opportunities provide the foundation for successful outcomes. It is critical that all caregivers receive training, support and respite to provide them with important tools for managing children and adolescents with FASD and to help burnout. 100-102 prevent reduce stress and Interventions for children with FASD that focus on a quality caregiving environment and integrate the role of family have been suggested 103, and caregiver training has been a repeated key to success across many research studies. 100,102,104 An online caregiver-training program has recently been developed and is currently undergoing larger field trials to provide caregivers with education, training and support to meet their needs, the needs of their child with FASD and their family based on caregiver and clinician input. 105 When children **FASD** encounter non-supportive with environments, their learning is impeded and this can lead to externalizing behaviours (e.g., aggression, acting out) or internalizing behaviours (e.g., shut down, withdrawal, anxiety). Training of foster care providers about FASD has been shown to improve stability of placement and reduce caregiver stress. 101,102

PRACTICE POINT #5: PHCP should be aware of FASD support services in their community and refer families to educational and family supports early.

Medical Review Including Medication Management

Children with FASD can have significant health concerns requiring coordination of care by a PHCP who understands their unique needs and provides continuity of care over the lifespan. A thorough medical history must be obtained from the individual or reliable caregiver, and a general physical and neurological exam are the gold standard. Screening for mental health disorders is also needed, as well as, screening for addictions for adolescents and young adults. Importantly, the majority of health issues are not specific to FASD and can present as other neurodevelopmental disabilities, which underscores the importance of

continuity of care. For example, if the child is in the foster care system, every effort should be made to maintain the same PHCP. For adolescents and young adults with FASD, having easy access to a PHCP with whom they have a trusting and nonjudgmental relationship can significantly improve quality of life and prevent many health complications.

The following list highlights the most frequent health issues associated with FASD requiring ongoing monitoring by their primary care provider:

Growth

Growth may be impaired in some children or may significant with normal, even neurodevelopmental impairment. Growth should be monitored at each visit using standardized growth curves. The growth curves produced by the Canadian Pediatric Endocrinology Group (CPEG) based on the 2010 WHO curves with some modifications are recommended. The Canadian Pediatric Society has endorsed these growth curves. 106 The detection of small stature (i.e., less that the 10%) is one criterion of full Fetal Alcohol Syndrome (FAS)¹, however it is important to rule out other causes of growth deficiency, such as malabsorption (celiac, short syndrome); pituitary growth hormone deficiency; nutritional deficiencies; other in utero drug exposures; gastroesophageal reflux; feeding disorders, including those related to sensory processing or emotional disturbances; other systemic chronic disease (e.g., cardiac, renal); genetic syndromes; or familial small stature (if biological parental stature is available).

Facial Dysmorphology

Elements of the characteristic face of FAS, with flat philtrum, thin upper lip and small palpebral fissures, may be observed in approximately 10% of patients who experienced prenatal exposure to alcohol. Facial analysis can be completed using a photographic software program from the University of Washington. However, there are a number of genetic syndromes that produce similar facial dysmorphology and must be investigated by appropriate genetic testing (e.g.,

Williams syndrome, Aarskog, syndrome 22q11.2 deletion syndrome). 1,109

Other Physical Effects of PAE

A number of birth defects have been described to recognize developing organ systems that can be damaged or disrupted by prenatal exposure to alcohol. While there is a long list of possibilities, the most important organ systems to screen include: vision (strabismus and myopia); hearing (malformations, sensorineural hearing loss that can impact language development); heart (septal defects needing surgery, arrhythmias that have major implications of side effects to certain medications); and palate (requiring plastic surgery to repair). It should be noted that these malformations are common in the general population and in other genetic syndromes, and therefore *not* specific to prenatal exposure to alcohol. 109 Additional minor dysmorphic features have also been documented such as a hockey stick palmar crease. 109 Thus, before an FASD diagnosis is made, it is critical to determine the etiology of these features (e.g., prenatal alcohol, genetic abnormality). Current genetic testing has revealed increasing number of chromosomal abnormalities that may be linked to the observed birth defects. Physicians can now request a microarray through genetic labs, if indicated, and refer for further genetic assessment counselling as appropriate. 109

Seizures

A retrospective chart review revealed that individuals with FASD are at increased risk of having a seizure disorder (~prevalence of 17.7%) compared to the general population. 110 Clinical presentation of the seizures can be subtle necessitating a thorough examination when conducting the child's history. An electroencephalogram (EEG) is recommended if there is a clinical suspicion. Atypical behaviours reported in sleep may require additional video EEG investigation and/or sleep deprived EEG. Epileptic abnormalities on the EEG need clinical correlation and may need treatment with an anticonvulsant. These medications need to be titrated with the same precautions as those used for other co-morbidities (e.g., ADHD).

Additionally, the knowledge of confirmed prenatal exposure to alcohol may also impact the efficacy of prescribed medications, rendering the patient more or less sensitive to the effects. Frequent monitoring is recommended. Any abnormalities on EEG also need to be considered in the context of the differential diagnosis of comorbid conditions, especially if medications such as stimulants for ADHD or selective serotonin reuptake inhibitors for mood have been prescribed.

Neuroimaging is not indicated in the diagnosis of FASD but can be considered as part of the differential diagnosis if there are abnormal hard neurological findings such as spasticity or microcephaly. Research level MRI studies are providing further information on the impact of alcohol on brain development.¹¹¹

Immune System

The effects of prenatal alcohol exposure on the developing neuroimmune system are complex and not fully understood. Longitudinal population based studies such as the Adverse Childhood Experiences Study, demonstrate that pre- and postnatal toxic stress can alter the immune system responses and the effects can be life-long. Prenatal alcohol exposure and/or exposure to traumatic stress is often linked to psychosocial determinants of health such as poverty; malnutrition and environmental exposure, which can further impact the developing immune system. With lower immune responses, the individual may be more prone to common infections.

Sleep

Disrupted sleep, both difficulties in falling asleep and staying asleep, is very common in FASD. 115-118 A description of the child's sleep patterns, the sleep environment and medications that are used throughout the day are essential pieces of the history that need to be collected by the PHCP. The causes of the sleep problems can be multiple and complex including the effects of prenatal alcohol exposure on the normal sleep cycle. Trauma and adverse life events, low iron levels (as found in restless leg syndrome), mental health issues such as anxiety or depression, school frustrations and bullying, chaotic home environment, technology in the bedroom, personal substance abuse in older

teens and young adults, and side effects of prescribed medications can all contribute to disrupted sleep patterns. Initial management steps include counselling regarding appropriate sleep hygiene and a stable emotional environment. Melatonin may be appropriate in conjunction with strict sleep hygiene and environmental modifications.

PRACTICE POINT #6: PHCP should screen children and adolescents with FASD for sleep disorders using a simple screen such as the 5-item pediatric sleep screening instrument, the BEARS (B=Bedtime Issues, E=Excessive Daytime Sleepiness, A=Night Awakenings, R=Regularity and Duration of Sleep, S=Snoring. 119 PHCP should provide early referrals to occupational therapists that can provide consultation on sensory processing, sleep and self-regulation. If Obstructive Sleep Apnea is suggested by clinical history, referral to a Sleep Clinic or an Ear, Nose and Throat Specialist is indicated. If nocturnal seizures are suspected, a sleep deprived EEG and/or referral to Neurologist is indicated.

Nutrition

Feeding difficulties often present during infancy where there has been prenatal exposure to drugs and alcohol. Symptoms may include poor coordination of suck and swallow, irritability associated with pain from gastroesophageal reflux, or coughing with intake of liquids. Feeding difficulties may also be related to environmental neglect and poor infant-maternal bonding. Early identification and management can prevent failure to thrive and other medical complications. Referral to services such as a Dietician, Occupational Therapist, Speech and Language Pathologist or a multidisciplinary feeding team may be indicated. In the preschool age group, sensitivities to different textures and tastes of foods can lead to food refusals. Nutritional deprivation in the early years can develop into lifelong unhealthy habits such as hoarding and stuffing of foods and not recognizing limits. Many medications prescribed for behaviour management can have side effects such as loss of appetite or excessive food cravings. The PHCP needs to use good judgment in choosing the appropriate

medications and frequently monitor weight, Body Mass Index and blood testing if indicated. 120,121

Elimination

Constipation is a frequent problem for all children and adolescents with disabilities including those with FASD. There are many factors to consider, including food choices, behavioural stool retention, gut motility, side effects of medications, and lack of home routines that must be contemplated in developing an appropriate management plan. Additional problems with urine control can be related to renal and bladder function. The PHCP needs to also be aware that trauma, especially sexual abuse, can trigger these problems and further investigations need to be approached with extreme sensitivity.

Mental Health Disorders

Over 90% of individuals with FASD have a coexisting mental health disorder^{52,122}, where ADHD is one of the most prevalent.⁶⁷ In teens and young adults there are often multiple conditions, such as depression, anxiety and substance abuse. These conditions may be primarily due to the impact of in utero alcohol exposure or secondary to adverse life experiences, or both. There is no evidence-based literature supporting unique management of mental health disorders in FASD. Anecdotal evidence suggests that the responses to medication are not predictably robust and may result in the prescribing of multiple medications. In recent years there has been an increase in the use of atypical antipsychotic drugs to manage aggression and "out of control" behavioural responses. General practice guidelines on the management of mental health conditions in the general population provides the PHCP with guidance in their management, recognizing that the neurobehavioural complexity of FASD may result in atypical responses to medications and thus close monitoring is recommended. 120,121 Cognitive therapy may be of limited efficacy when there are significant functional limitations in communication and cognitive function as is seen in FASD. Therapy that is trauma-informed and based attachment theory recommended.³³ Access to a mental health team that has received FASD training and who can help manage these complex cases is helpful in these situations.

PRACTICE POINT #7: The PHCP must evaluate the environment prior to prescribing medication for presenting behaviours. When doing so, they must follow patients closely for side effects as their responses may differ from children with no underlying FASD.

Dental Health

Although prenatal exposure of alcohol alone may not affect the development of dental enamel in utero, maternal malnutrition can play a role. A recent population based study suggests that children of mothers with an alcohol-related diagnosis have increased dental disease including dental admissions. 123 More importantly, the postnatal diet with inappropriate bottle feeding and high sugar intake compounded by poor oral hygiene can cause cavities. Individuals with FASD have difficulty remembering routines for daily especially personal hygiene without constant reminders. A picture list for daily routines on the bathroom mirror can be very helpful. Dental cavities and abscesses are painful and can contribute to irritable behaviours and disrupted sleep. Early recognition is very important. Regular six month routine dental monitoring is recommended. In the adolescent years, many teens with FASD have dental malocclusion related to midfacial under-development and may require orthodontic treatment.

PRACTICE POINT #8: The PHCP should be vigilant in assessing for health issues that may occur more frequently in individuals with FASD. Early identification and treatment of co-morbid health conditions will reduce the burden of care for individuals with FASD and the families caring for them. The PHCP has an important role in the early identification of individuals who may have experienced prenatal alcohol exposure by conducting a complete and sensitive medical and prenatal history.

COMMUNITY PARTNERSHIPS

What Partnerships Are Needed?

An integrated network of multidisciplinary, community-based care providers is fundamental to the management of children with FASD. Children with FASD and their families often have complex needs that span multiple systems (e.g., health, education, social services etc.) As FASD is a lifelong condition, supports and services will change over time, making continuity of care a major contributor to ongoing success. It is important to foster partnerships across all systems early in the diagnostic process, as they will be critical for supporting children and their families throughout the lifespan. The spectrum of services in a community may include screening; early intervention: access to a multidisciplinary diagnostic team; developmental, educational and behavioural interventions; primary health care; mental health and addiction services; rehabilitation services; tiered prevention services; and employment and housing support (for older individuals with FASD).

PRACTICE POINT #9: A coordinated team approach, with the PHCP as a key player, should be based on the principles of child-centered and family-centered care, inter-professional collaboration, evidence-based knowledge, flexible care provision, shared communication, effective resource coordination, and integration of care.

Role of the PHCP (e.g., Pediatrician, Family Physician)

The complexity of FASD requires a PHCP who is knowledgeable about FASD and has access to a collaborative multidisciplinary team. The PHCP may be a resource to help navigate the systems for the family and individual with FASD. A key element of providing care to families of children with FASD is providing education to caregivers including a discussion about the behavioural and cognitive effects of FASD, goals for intervention, and how to advocate for supports and services for their children. The supports will naturally need to change at different ages and life stages, and an anticipatory guidance approach is needed.

PRACTICE POINT #10: The PHCP and support team will play a key role in ensuring a successful transition from childhood to adulthood. This will include transition planning and helping families secure the services and resources their child will need in adulthood.

FASD AND Alcohol Resources in the Community

It is important to identify community-based resources related to FASD, alcohol, addictions, and prevention for each region. These resources should be readily available to families from their PHCP and/or community team and/or shared through community partnerships.

PRENATAL MANAGMENT

Education and Counselling from PHCP (e.g., Pediatrician, Family Physician)

All PHCP need to be aware of the risks associated with alcohol use during pregnancy and to provide prevention messaging to all women of childbearing age and their partners. There are many resources for primary prevention messaging such as posters, pamphlets and other media. Primary prevention messages displayed in a health care setting may not change behaviour, but can be used to start the conversation about "how to have a healthy pregnancy" by the PHCP.

Counselling

PHCP are in an ideal position to talk to all women about their alcohol use at multiple points in their lifespan. Alcohol use in pregnancy crosses all ethnic and economic strata. The question of alcohol use should be asked without judgment and with the offer of positive support. Women may be fearful about disclosing their alcohol use during pregnancy, as they may lose their child to Child Welfare authorities. Many women problematic alcohol use have also experienced multigenerational trauma, loss and undiagnosed mental health issues, lack of educational opportunities, stressful home unplanned environments, pregnancies intimate partner violence. It is incumbent that the PHCP recognizes these compounding factors. The Society Obstetricians and Gynecologists of

Canada have published *Alcohol Use and Pregnancy Consensus Clinical Guidelines* that describe best practices for counselling women about alcohol use and pregnancy, and provide evidence-based recommendations to help PHCP provide the best care in these sensitive situations. ¹²⁵

An emerging "high risk" group for alcohol use in pregnancy may be professional women, who may not be planning pregnancy, and who may use alcohol as a means to cope with stress and mental health issues. Binge drinking in adolescent girls is a concern, as this is an at-risk age group for unplanned and late recognition of pregnancy. The PHCP needs to feel comfortable asking questions about problematic alcohol and/or substance use to all women of childbearing age. Screening tools (e.g., TWEAK, T-ACE, CAGE) can be extremely useful for identifying "at-risk" populations, who would benefit from further counselling, referral, intervention, and/or for treatment problematic alcohol and/or substance use. The PHCP is in a position to provide prevention messaging to male partners to help them understand the male's role in having a healthy pregnancy, including information on alcohol and stress in pregnancy.

Harm Reduction

There are many women who are unable to change behaviour even after they receive information about problematic alcohol and/or substance use and may require a more personalized approach. This approach needs to be framed within a social determinants of health context and based on harm reduction. Brief interventions, such as motivational interviewing, have been effective. The PHCP needs to know the next level of tertiary prevention and how these patients can access more intensive supports and programs that provide access to contraception, physical and mental health care and targeted educational resources for women. Although the primary goal is less prenatal alcohol exposedpregnancies, if this cannot be achieved, harm reduction is supported without blaming or shaming. 33,34,86

Prenatal Investigations

Prenatal fetal ultrasound may identify differences in limb length, congenital cardiac defects, and other alcohol-related birth defects, if present. However, most cases of FASD show no specific features on prenatal imaging that would identify FASD. At this time, there are no specific genetic tests that would identify FASD.

Maternal Nutrition

FASD has been associated with the "poverty trap¹²⁶", which is associated with a lack of resources leading to poor nutrition during pregnancy, including decreased folate, vitamin, iron and protein intake. Additionally, the calories from alcohol reduce food intake and monies may be directed to the purchase of alcohol rather than nutritious foods. These factors, along with lack of antenatal care, can also adversely impact the management of gestational diabetes.

Nutrient deficiencies compound effects of prenatal alcohol exposure 128, and the potential of using nutrient supplementation as a factor, protective currently is investigation. 129-131 Maternal iron deficiency contributes to poor iron stores in the newborn, which can impact cognitive development. 132,133 Micronutrient supplements, such as choline 134, may also improve cognitive function following alcohol exposure. prenatal However, enrichment does not replace the need to provide the woman and her partner with advice on avoiding alcohol in pregnancy.

CONCLUSION

There is a pressing need for the engagement of PHCP in the active care of individuals with FASD and their families over the lifespan. PHCP are trained in screening, prevention, and management of health needs, and are in the position to coordinate sub-specialty referrals as needed. They can also provide families and their communities with education and interdisciplinary coordinated care. Individuals with FASD are often involved with multiple systems such as health, education, social services, and justice. The role of a PHCP can include critical consultation and advocacy for affected individuals. Finally, subspecialty care

may not always be easily accessible to individuals and families in remote areas, and the PHCP can provide an important link to expert care. The purpose of these guidelines is to provide PHCP with the best evidence for the medical, behavioural, and mental health recommendations for the care of individuals with FASD in their communities (see worksheet template in Appendix I). Engaging PHCP will provide a truly integrated care system for individuals affected by prenatal alcohol exposure and their families.

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REFERENCES

- 1. Chudley AE, Conry J, Cook JL, Loock C, Rosales T, LeBlanc N. Fetal alcohol spectrum disorder: Canadian guidelines for diagnosis. CMAJ 2005;172(5 Suppl):S1-S21.
- 2. Clarren SK, Smith DW. The fetal alcohol syndrome. N Engl J Med May 11 1978;298(19):1063-1067.
- 3. Mattson SN, Schoenfeld AM, Riley EP. Teratogenic effects of alcohol on brain and behavior. Alcohol research & health: The Journal of the National Institute on Alcohol Abuse and Alcoholism 2001;25(3):185.
- 4. Abel EL, Sokol RJ. Maternal and fetal characteristics affecting alcohol's teratogenicity. Neurobehav Toxicol Teratol Jul-Aug 1986;8(4):329-334.
- 5. Uban KA, Bodnar T, Butts K, Sliwowska JH, Comeau W, Weinberg J. Direct and indirect

- mechanisms of alcohol teratogenicity: Implications for understanding alterations in brain and behavior in FASD. Fetal Alcohol Spectrum Disorder: Management and policy perspectives of FASD, First Edition: Wiley; 2011.
- 6. Loock C, Conry J, Cook JL, Chudley AE, Rosales T. Identifying fetal alcohol spectrum disorder in primary care. CMAJ Mar 1 2005;172(5):628-630.
- 7. Chudley AE, Longstaffe SE. Fetal Alcohol Syndrome and Fetal Alcohol Spectrum Diosrder. In: Cassidy S, Allanson J, eds. Management of Genetic Syndromes. Third ed. New York, NY: John Wiley and Sons, Inc.; 2010:363-380.
- 8. Riley EP, Clarren SK, Weinberg J, Jonsson E. Fetal Alcohol Spectrum Disorder: Management and Policy Perspectives of FASD. Wiley-Blackwell; 2010.
- 9. Gahagan S, Sharpe TT, Brimacombe M, et al. Pediatricians' knowledge, training, and experience in the care of children with fetal alcohol syndrome. Pediatrics Sep 2006;118(3):e657-668.
- 10. Chavez GF, Cordero JF, Becerra JE. Leading major congenital malformations among minority groups in the United States, 1981-1986. MMWR CDC Surveill Summ 1988;37(3):17-24.
- 11. Sokol RJ, Clarren SK. Guidelines for use of terminology describing the impact of prenatal alcohol on the offspring. Alcohol Clin Exp Res 1989;13(4):597-598.
- 12. Sampson PD, Streissguth AP, Bookstein FL, et al. Incidence of fetal alcohol syndrome and prevalence of alcohol-related neurodevelopmental disorder. Teratology 1997;56(5):317-326.
- 13. Sampson PD, Bookstein FL, Barr HM, Streissguth AP. Prenatal alcohol exposure, birthweight, and measures of child size from birth to age 14 years. Am J Public Health 1994;84(9):1421-1428.
- 14. May PA, Gossage JP, Kalberg WO, et al. Prevalence and epidemiologic characteristics of FASD from various research methods with an emphasis on recent in-school studies. Dev Disabil Res Rev 2009;15(3):176-192.
- 15. Health CfAaM. Fetal alcohol spectrum disorders: How widespread are they in Canada? 2013; http://www.camh.ca/en/hospital/about_camh/n ewsroom/news_releases_media_advisories_an

- d_backgrounders/current_year/Pages/Fetal-Alcohol-Spectrum-Disorders-How-Widespread-Are-They-in-Canada.aspx.
 Accessed March 21, 2014.
- 16. de SL, Memo L, Pichini S, Tarani L, Vagnarelli F. Fetal alcohol syndrome: new perspectives for an ancient and underestimated problem. J Matern Fetal Neonatal Med 2011;24 Suppl 1:34-37.
- 17. Paintner A, Williams AD, Burd L. Fetal alcohol spectrum disorders--implications for child neurology, part 1: prenatal exposure and dosimetry. J Child Neurol 2012;27(2):258-263
- 18. Maier SE, West JR. Drinking patterns and alcohol-related birth defects. Alcohol research & health: The Journal of the National Institute on Alcohol Abuse and Alcoholism 2001;25(3):168-174.
- 19. Stade BC, Ali A, Bennett D, et al. The burden of prenatal exposure to alcohol: revised measurement of cost. The Canadian Journal of Clinical Pharmacology 2009;16(1):e91-102.
- 20. Popova S, Lange S, Burd L, Rehm J. Health care burden and cost associated with fetal alcohol syndrome: based on official Canadian data. PLoS.One 2012;7(8):e43024.
- 21. CCSA CCoSA. Drinking Guidelines. 2013; http://www.ccsa.ca/Eng/topics/alcohol/drinking-guidelines/Pages/default.aspx. Accessed April 3, 2014.
- 22. McKnight-Eily LR, Liu Y, Brewer RD, et al. Vital signs: communication between health professionals and their patients about alcohol use--44 states and the District of Columbia, 2011. MMWR Morb Mortal Wkly Rep Jan 10 2014;63(1):16-22.
- 23. McCormick KA, Cochran NE, Back AL, Merrill JO, Williams EC, Bradley KA. How primary care providers talk to patients about alcohol: a qualitative study. J Gen Intern Med Sep 2006;21(9):966-972.
- 24. Shannon GD, Alberg C, Nacul L, Pashayan N. Preconception Healthcare and Congenital Disorders: Systematic Review of the Effectiveness of Preconception Care Programs in the Prevention of Congenital Disorders. Matern Child Health J Oct 4 2013.
- 25. Rendall-Mkosi K, Morojele N, London L, Moodley S, Singh C, Girdler-Brown B. A randomized controlled trial of motivational interviewing to prevent risk for an alcoholexposed pregnancy in the Western Cape,

- South Africa. Addiction Apr 2013;108(4):725-732.
- 26. National Institute of Alcohol Abuse and Alcoholism N. Assessing alcohol problems: A guide for clinicians and researchers, Second edition. Washington, DC: Dept. of Health and Human Services, Public Health Service;2003.
- 27. Babor TF, Higgins-Biddle J, Saunders JB, Monteiro MG. AUDIT. The alcohol use disorders identification test: Guidelines for use in primary care. 2nd Edition. Geneva, Switzerland: Department of Mental Health and Substance Dependence, World Health Organization;2001.
- 28. CAPHC. National screening tool kit for children and youth identified and potentially affected by FASD. 2010; http://ken.caphc.org/xwiki/bin/view/FASDScreeningToolkit/National+Screening+Tool+Kit+for+Children+and+Youth+Identified+and+Potentially+Affected+by+FASD. Accessed March 17, 2014.
- 29. Goh YI, Chudley AE, Clarren SK, et al. Development of Canadian screening tools for fetal alcohol spectrum disorder. Can J Clin Pharmacol Summer 2008;15(2):e344-366.
- 30. LaFrance MA, McLachlan K, Nash K, et al. Evaluation of the Neurobehavioral Screening Tool in Children with Fetal Alcohol Spectrum Disorders (FASD). J Popul Ther Clin Pharmacol 2014;21(2):e197-210.
- 31. Zizzo N, DiPietro N, Green CR, Reynolds JN, Bell E, Racine E. Review and reflections on ethics in screening for biomarkers of prenatal alcohol exposure. Alcohol Clin Exp Res 2013;(in press).
- 32. Yan A, Bell E, Racine E. Ethical and social challenges in newborn screening for prenatal alcohol exposure. Can J Neurol Sci Jan 2014;41(1):115-118.
- 33. Poole N. Bringing a women's health perspective to FASD prevention. In: Riley EP, Clarren SK, Weinberg J, Jonsson E, eds. Fetal Alcohol Spectrum Disorder: Management and policy perspectives: Wiley-Blackwell; 2010:161-174.
- 34. Grant TM. Maternal alcohol and drug abuse: Effective case management with high-risk mothers and their children. In: Rubin A, ed. Programs and interventions for maltreated children and families at risk. The clinician's guide to evidence-based practice series: Wiley Blackwell Publishers; 2011:207-221.

- 35. Kobor MS, Weinberg J. Focus on: Epigenetics and fetal alcohol spectrum disorders. Alcohol Research & Health 2011;34(1):29-37.
- 36. Hellemans KG, Sliwowska JH, Verma P, Weinberg J. Prenatal alcohol exposure: fetal programming and later life vulnerability to stress, depression and anxiety disorders. Neurosci Biobehav Rev May 2010;34(6):791-807.
- 37. Chudley AE. Genetic factors in Fetal Alcohol Spectrum Disorder. In: Riley EP, Clarren SK, Weinberg J, Jonsson E, eds. Fetal Alcohol Spectrum Disorder. Management and Policy Perspectives of FASD. New York: Wiley/Blackwell; 2011:109-126.
- 38. Zhang X, Sliwowska JH, Weinberg J. Prenatal alcohol exposure and fetal programming: effects on neuroendocrine and immune function. Exp Biol Med (Maywood.). 2005;230(6):376-388.
- 39. Hall JG. Epigenetics: What does it mean for paediatric practice? Paediatr Child Health Jan 2014;19(1):27-30.
- 40. Hellemans KG, Verma P, Yoon E, Yu WK, Young AH, Weinberg J. Prenatal alcohol exposure and chronic mild stress differentially alter depressive- and anxiety-like behaviors in male and female offspring. Alcohol Clin Exp Res Apr 2010;34(4):633-645.
- 41. CDC CfDC. Adverse Childhood Experiences (ACE) Study. 2013; http://www.cdc.gov/ace/about.htm. Accessed March 28, 2014.
- 42. University CotDCaH. http://developingchild.harvard.edu/.
- 43. O'Connor MJ, Kogan N, Findlay R. Prenatal alcohol exposure and attachment behavior in children. Alcohol Clin Exp Res Oct 2002;26(10):1592-1602.
- 44. Garner AS, Shonkoff JP, Committee on Psychosocial Aspects of C, et al. Early childhood adversity, toxic stress, and the role of the pediatrician: translating developmental science into lifelong health. Pediatrics Jan 2012;129(1):e224-231.
- 45. Shonkoff JP. Protecting brains, not simply stimulating minds. Science Aug 19 2011;333(6045):982-983.
- 46. Shonkoff JP, Boyce WT, McEwen BS. Neuroscience, molecular biology, and the childhood roots of health disparities: building a new framework for health promotion and disease prevention. JAMA Jun 3 2009;301(21):2252-2259.

- 47. Shonkoff JP, Garner AS, Committee on Psychosocial Aspects of C, et al. The lifelong effects of early childhood adversity and toxic stress. Pediatrics Jan 2012;129(1):e232-246.
- 48. Ahmed-Landeryou MJ. Fetal central nervous system development and alcohol--the evidence so far. Fetal Pediatr Pathol Dec 2012;31(6):349-359.
- 49. Fuchs D, Burnside L, Marchenski S, Mudry A. Children with FASD-related disabilities receiving services from child welfare agencies in Manitoba. Int J Ment Health Addiction 2010;8:232-244.
- 50. Zeanah CH, Shauffer C, Dozier M. Foster care for young children: why it must be developmentally informed. J Am Acad Child Adolesc Psychiatry Dec 2011;50(12):1199-1201.
- 51. Streissguth AP, Barr HM, Kogan J, Bookstein FL. Understanding the occurrence of secondary disabilities in clients with fetal alcohol syndreome (FAS) and fetal alcohol effects (FAE). Final report to the Centers for Disease Control and Prevention. Seattle: University of Washington, Fetal Alcohol and Drug Unit;1996.
- 52. Streissguth AP, Bookstein FL, Barr HM, Sampson PD, O'Malley K, Young JK. Risk factors for adverse life outcomes in fetal alcohol syndrome and fetal alcohol effects. J Dev Behav Pediatr 2004;25(4):228-238.
- 53. Macmillan HL. Protecting children from maltreatment: A Canadian call to action. Paediatr Child Health Oct 2013;18(8):409-410.
- 54. Tonmyr L, Hovdestad WE, Draca J. Commentary on Canadian child maltreatment data. Journal of Interpersonal Violence Jan 2014;29(1):186-197.
- 55. Mattson SN, Crocker N, Nguyen TT. Fetal alcohol spectrum disorders: neuropsychological and behavioral features. Neuropsychol Rev 2011;21(2):81-101.
- 56. Rasmussen C, Andrew G, Zwaigenbaum L, Tough S. Neurobehavioural outcomes of children with fetal alcohol spectrum disorders: A Canadian perspective. Paediatr Child Health 2008;13(3):185-191.
- 57. Quattlebaum JL, O'Connor MJ. Higher functioning children with prenatal alcohol exposure: Is there a specific neurocognitive profile? Child Neuropsychol Aug 21 2012.
- 58. Paley B, O'Connor MJ. Neurocognitive and neurobehavioural impairments in individuals

- with fetal alcohol spectrum disorders: Recognition and assessment. Int J Disabil Hum Dev 2007;6:127-142.
- 59. Ware AL, Crocker N, O'Brien JW, et al. Executive function predicts adaptive behavior in children with histories of heavy prenatal alcohol exposure and attention-deficit/hyperactivity disorder. Alcohol Clin Exp Res 2012;36(8):1431-1441.
- 60. Fagerlund A, Autti-Ramo I, Hoyme HE, Mattson SN, Korkman M. Risk factors for behavioural problems in foetal alcohol spectrum disorders. Acta Paediatr 2011;100(11):1481-1488.
- 61. Fagerlund A, Autti-Ramo I, Kalland M, et al. Adaptive behaviour in children and adolescents with foetal alcohol spectrum disorders: a comparison with specific learning disability and typical development. Eur Child Adolesc Psychiatry Apr 2012;21(4):221-231.
- 62. Astley SJ. Profile of the first 1,400 patients receiving diagnostic evaluations for fetal alcohol spectrum disorder at the Washington State Fetal Alcohol Syndrome Diagnostic & Prevention Network. Can J Clin Pharmacol 2010;17(1):e132-e164.
- 63. Rasmussen C. Executive functioning and working memory in fetal alcohol spectrum disorder. Alcohol Clin Exp Res 2005;29(8):1359-1367.
- 64. Pei J, Job J, Kully-Martens K, Rasmussen C. Executive function and memory in children with Fetal Alcohol Spectrum Disorder. Child Neuropsychol 2011;17(3):290-309.
- 65. Quattlebaum JL, O'Connor MJ. Higher functioning children with prenatal alcohol exposure: is there a specific neurocognitive profile? Child Neuropsychol 2013;19(6):561-578.
- 66. Manji S, Pei J, Loomes C, Rasmussen C. A review of the verbal and visual memory impairments in children with foetal alcohol spectrum disorders. Dev Neurorehabil Aug 2009;12(4):239-247.
- 67. Pei J, Denys K, Hughes J, Rasmussen C. Mental health issues in fetal alcohol spectrum disorder. J Ment Health 2011;20(5):438-448.
- 68. Kooistra L, Crawford S, Gibbard B, Ramage B, Kaplan BJ. Differentiating attention deficits in children with fetal alcohol spectrum disorder or attention-deficit-hyperactivity disorder. Dev Med Child Neurol Feb 2010;52(2):205-211.

- 69. Mattson SN, Calarco KE, Lang AR. Focused and shifting attention in children with heavy prenatal alcohol exposure. Neuropsychology May 2006;20(3):361-369.
- 70. Rasmussen C, Wyper K, Talwar V. The relation between theory of mind and executive functions in children with fetal alcohol spectrum disorders. Can J Clin Pharmacol Summer 2009;16(2):e370-380.
- 71. Coles CD, Kable JA, Taddeo E. Math performance and behavior problems in children affected by prenatal alcohol exposure: intervention and follow-up. J Dev Behav Pediatr Feb 2009;30(1):7-15.
- 72. Paley B, O'Connor MJ. Behavioral interventions for children and adolescents with fetal alcohol spectrum disorders. Alcohol research & health: The Journal of the National Institute on Alcohol Abuse and Alcoholism 2011;34(1):64-75.
- 73. A 14-month randomized clinical trial of treatment strategies for attention-deficit/hyperactivity disorder. The MTA Cooperative Group. Multimodal Treatment Study of Children with ADHD. Arch Gen Psychiatry Dec 1999;56(12):1073-1086.
- 74. Subcommittee on Attention-Deficit/Hyperactivity D, Steering Committee on Quality I, Management, et al. ADHD: clinical practice guideline for the diagnosis, evaluation, and treatment of attention-deficit/hyperactivity disorder in children and adolescents. Pediatrics Nov 2011;128(5):1007-1022.
- 75. Sumner CR, Gathercole S, Greenbaum M, et al. Atomoxetine for the treatment of attention-deficit/hyperactivity disorder (ADHD) in children with ADHD and dyslexia. Child and Adolescent Psychiatry and Mental Health 2009;3:40.
- Michelson D, Allen AJ, Busner J, et al. Oncedaily atomoxetine treatment for children and adolescents with attention deficit hyperactivity disorder: a randomized, placebo-controlled study. Am J Psychiatry Nov 2002;159(11):1896-1901.
- 77. Cinnamon Bidwell L, Dew RE, Kollins SH. Alpha-2 adrenergic receptors and attention-deficit/hyperactivity disorder. Current Psychiatry Reports Oct 2010;12(5):366-373.
- 78. Olatunji BO, Cisler JM, Deacon BJ. Efficacy of cognitive behavioral therapy for anxiety disorders: a review of meta-analytic findings.

- Psychiatr Clin North Am Sep 2010;33(3):557-577.
- 79. Walkup JT, Albano AM, Piacentini J, et al. Cognitive behavioral therapy, sertraline, or a combination in childhood anxiety. N Engl J Med Dec 25 2008;359(26):2753-2766.
- 80. Silverman WK, Ollendick TH. Evidence-based assessment of anxiety and its disorders in children and adolescents. J Clin Child Adolesc Psychol. Sep 2005;34(3):380-411.
- 81. Alliance TCA-DHDR. http://www.caddra.ca.
- 82. Steiner NJ, Sheldrick RC, Gotthelf D, Perrin EC. Computer-based attention training in the schools for children with attention deficit/hyperactivity disorder: a preliminary trial. Clin Pediatr (Phila.) Jul 2011;50(7):615-622.
- 83. Arns M, de Ridder S, Strehl U, Breteler M, Coenen A. Efficacy of neurofeedback treatment in ADHD: the effects on inattention, impulsivity and hyperactivity: a meta-analysis. Clin EEG Neurosci Jul 2009;40(3):180-189.
- 84. Forehand R, Jones DJ, Parent J. Behavioral parenting interventions for child disruptive behaviors and anxiety: what's different and what's the same. Clin Psychol Rev Feb 2013;33(1):133-145.
- 85. Haley DW, Handmaker NS, Lowe J. Infant stress reactivity and prenatal alcohol exposure. Alcohol Clin Exp Res 2006;30(12):2055-2064.
- 86. Motz M, Espinet SD, Jeong JJ, et al. The role of the mother-child relationship in developmental outcomes of infants and young children with and without prenatal alcohol exposure. J Popul Ther Clin Pharmacol 2011;18(3):e544-563.
- 87. Molteno CD, Jacobson JL, Carter RC, Dodge NC, Jacobson SW. Infant emotional withdrawal: a precursor of affective and cognitive disturbance in fetal alcohol spectrum disorders. Alcohol Clin Exp Res Feb 2014;38(2):479-488.
- 88. Jacobson SW, Jacobson JL, Stanton ME, Meintjes EM, Molteno CD. Biobehavioral markers of adverse effect in fetal alcohol spectrum disorders. Neuropsychol Rev Jun 2011;21(2):148-166.
- 89. Lambert BL, Bann CM, Bauer CR, et al. Risktaking behavior among adolescents with prenatal drug exposure and extrauterine environmental adversity. J Dev Behav Pediatr. Nov-Dec 2013;34(9):669-679.

- 90. Dej E. What was once sick is now bad: the shift from victim to deviant identity for those diagnosed with FASD. Can J Sociol 2011;36(2):137-160.
- 91. Frankel F, Paley B, Marquardt R, O'Connor M. Stimulants, neuroleptics, and children's friendship training for children with fetal alcohol spectrum disorders. J Child Adolesc Psychopharmacol Dec 2006;16(6):777-789.
- 92. O'Malley KD, Koplin B, Dohner VA. Psychostimulant clinical response in fetal alcohol syndrome. Can J Psychiatry Feb 2000;45(1):90-91.
- 93. O'Leary C, Zubrick SR, Taylor CL, Dixon G, Bower C. Prenatal alcohol exposure and language delay in 2-year-old children: the importance of dose and timing on risk. Pediatrics Feb 2009;123(2):547-554.
- 94. Coggins T, Friet T, Morgan T. Analysing narrative productions in older school age children and adolescents with fetal alcohol syndrome: An experimental toll for clinical applications. Clin Linguist Phon 1998;12(3):221-236.
- 95. Rasmussen C, Becker M, McLennan J, Urichuk L, Andrew G. An evaluation of social skills in children with and without prenatal alcohol exposure. Child Care Health Dev Sep 2011;37(5):711-718.
- 96. McGee CL, Bjorkquist OA, Price JM, Mattson SN, Riley EP. Social information processing skills in children with histories of heavy prenatal alcohol exposure. J Abnorm Child Psychol Aug 2009;37(6):817-830.
- 97. McGee CL, Fryer SL, Bjorkquist OA, Mattson SN, Riley EP. Deficits in social problem solving in adolescents with prenatal exposure to alcohol. Am J Drug Alcohol Abuse 2008;34(4):423-431.
- 98. Rasmussen C, Talwar V, Loomes C, Andrew G. Brief report: lie-telling in children with fetal alcohol spectrum disorder. J Pediatr Psychol Mar 2008;33(2):220-225.
- 99. Bishop S, Gahagan S, Lord C. Re-examining the core features of autism: a comparison of autism spectrum disorder and fetal alcohol spectrum disorder. J Child Psychol Psychiatry Nov 2007;48(11):1111-1121.
- 100. Petrenko CL, Tahir N, Mahoney EC, Chin NP. Prevention of secondary conditions in Fetal Alcohol Spectrum Disorders: Identification of systems-level barriers. Matern Child Health J Nov 1 2013.

- 101. Brown JD, Bednar LM. Challenges of parenting children with Fetal Alcohol Spectrum Disorder: A concept map. J Fam Soc Work 2004;8(3):1-17.
- 102. Pelech W, Badry D, Daoust G. It takes a team: Improving placement stability among children and youth with FASD in care in Canada. Child Youth Serv Rev 2013;35:120-127.
- 103. Olson HC, Oti R, Gelo J, Beck S. "Family matters:" fetal alcohol spectrum disorders and the family. Dev Disabil Res Rev 2009;15(3):235-249.
- 104. Peadon E, Rhys-Jones B, Bower C, Elliott EJ. Systematic review of interventions for children with Fetal Alcohol Spectrum Disorders. BMC Pediatr 2009;9:35.
- 105. Green CR, Roane J, Hewitt A, et al. Frequent behavoural challenges in children with Fetal Alcohol Spectrum Disorder: A needs-based assessment reported by caregivers and clinicians. J Popul Ther Clin Pharmacol 2014;21(3):e405-e420.
- 106. Lawrence S, Cummings E, Chanoine JP, et al.
 Canadian Pediatric Endocrine Group
 extension to WHO growth charts: Why
 bother? Paediatr Child Health Jun
 2013;18(6):295-297.
- 107. Astley SJ, Clarren SK. Measuring the facial phenotype of individuals with prenatal alcohol exposure: correlations with brain dysfunction. Alcohol Alcohol Mar-Apr 2001;36(2):147-159.
- 108. Andrew G. Diagnosis of FASD: An overview. In: Riley EP, Clarren SK, Weinberg J, Jonsson E, eds. Fetal Alcohol Spectrum Disorder: Management and policy perspectives: Wiley-Blackwell; 2011:127-148.
- 109. Leibson T, Neuman G, Chudley AE, Koren G. The differential diagnosis of fetal alcohol spectrum disorder. J Popul Ther Clin Pharmacol 2014;21(1):e1-e30.
- 110. Bell SH, Stade B, Reynolds JN, et al. The remarkably high prevalence of epilepsy and seizure history in fetal alcohol spectrum disorders. Alcohol Clin Exp Res 2010;34(6):1084-1089.
- 111. Treit S, Lebel C, Baugh L, Rasmussen C, Andrew G, Beaulieu C. Longitudinal MRI reveals altered trajectory of brain development during childhood and adolescence in fetal alcohol spectrum disorders. J Neurosci Jun 12 2013;33(24):10098-10109.
- 112. Anda RF, Centers for Disease C, Felitti VJ, Permanente K. The adverse childhood

- experiences study. 2008; http://acestudy.org/. Accessed March 27, 2014.
- 113. Anda RF, Felitti VJ, Bremner JD, et al. The enduring effects of abuse and related adverse experiences in childhood. A convergence of evidence from neurobiology and epidemiology. Eur Arch Psychiatry Clin Neurosci. Apr 2006;256(3):174-186.
- 114. Dube SR, Felitti VJ, Dong M, Giles WH, Anda RF. The impact of adverse childhood experiences on health problems: evidence from four birth cohorts dating back to 1900. Prev Med Sep 2003;37(3):268-277.
- 115. Jan JE, Asante KO, Conry JL, et al. Sleep health issues for children with FASD: Clinical considerations. Int J Pediatr 2010;2010.
- 116. Ipsiroglu OS, McKellin WH, Carey N, Loock C. "They silently live in terror..." why sleep problems and night-time related quality-of-life are missed in children with a fetal alcohol spectrum disorder. Soc Sci Med Feb 2013;79:76-83.
- 117. Chen ML, Olson HC, Picciano JF, Starr JR, Owens J. Sleep problems in children with fetal alcohol spectrum disorders. J Clin Sleep Med 2012;8(4):421-429.
- 118. Wengel T, Hanlon-Dearman AC, Fjeldsted B. Sleep and sensory characteristics in young children with fetal alcohol spectrum disorder. J Dev Behav Pediatr Jun 2011;32(5):384-392.
- 119. Mindell JA, Owens JA. A clinical guide to pediatric sleep: Diagnosis and management of sleep problems, volume 2. Lippincott Williams & Wilkins; 2003.
- 120. Ho J, Panagiotopoulos C, McCrindle B, et al. Management recommendations for metabolic complications associated with second-generation antipsychotic use in children and youth. Paediatr Child Health Nov 2011;16(9):575-580.
- 121. Ho J, Panagiotopoulos C, McCrindle B, Grisaru S, Pringsheim T, group Cg. Management recommendations for metabolic complications associated with second generation antipsychotic use in children and youth. J Can Acad Child Adolesc Psychiatry Aug 2011;20(3):234-241.
- 122. Streissguth AP, Barr HM, Bookstein FL, Sampson PD, Olson HC. The long-term neurocognitive consequences of prenatal alcohol exposure: A 14-year study. Psychological Sicence 1999;10:186-190.
- 123. O'Leary CM, Slack-Smith LM. Dental hospital admissions in the children of mothers

- with an alcohol-related diagnosis: a population-based, data-linkage study. J Pediatr Aug 2013;163(2):515-520 e511.
- 124. Paley B, O'Connor MJ. Intervention for individuals with fetal alcohol spectrum disorders: treatment approaches and case management. Dev Disabil Res Rev 2009;15(3):258-267.
- 125. Carson G, Cox LV, Crane J, et al. Alcohol use and pregnancy consensus clinical guidelines. J Obstet Gynaecol Can Aug 2010;32(8 Suppl 3):S1-31.
- 126. Thanh NX, Jonsson E, Moffatt J, Dennett L. Fetal alcohol spectrum disorder--poverty trap? J Popul Ther Clin Pharmacol 2013;20(1):e63-66.
- 127. Fuglestad AJ, Fink BA, Eckerle JK, et al. Inadequate intake of nutrients essential for neurodevelopment in children with fetal alcohol spectrum disorders (FASD). Neurotoxicol Teratol Sep-Oct 2013;39:128-132.
- 128. Keen CL, Uriu-Adams JY, Skalny A, et al. The plausibility of maternal nutritional status being a contributing factor to the risk for fetal alcohol spectrum disorders: the potential influence of zinc status as an example. Biofactors Mar-Apr 2010;36(2):125-135.
- 129. Patten AR, Sickmann HM, Dyer RA, Innis SM, Christie BR. Omega-3 fatty acids can reverse the long-term deficits in hippocampal synaptic plasticity caused by prenatal ethanol exposure. Neurosci Lett Sep 13 2013;551:7-
- 130. Thomas JD, Abou EJ, Dominguez HD. Prenatal choline supplementation mitigates the adverse effects of prenatal alcohol exposure on development in rats. Neurotoxicol Teratol Sep-Oct 2009;31(5):303-311.
- 131. Thomas JD, Idrus NM, Monk BR, Dominguez HD. Prenatal choline supplementation mitigates behavioral alterations associated with prenatal alcohol exposure in rats. Birth Defects Res A Clin Mol Teratol Oct 2010;88(10):827-837.
- 132. Carter RC, Jacobson SW, Molteno CD, Jacobson JL. Fetal alcohol exposure, irondeficiency anemia, and infant growth. Pediatrics 2007;120(3):559-567.
- 133. Carter RC, Jacobson JL, Burden MJ, et al. Iron deficiency anemia and cognitive function in infancy. Pediatrics Aug 2010;126(2):e427-434.

- 134. Wozniak JR, Fuglestad AJ, Eckerle JK, et al. Choline supplementation in children with fetal alcohol spectrum disorders has high feasibility and tolerability. Nutrition Research Nov 2013;33(11):897-904.
- 135. American Academy of Pediatrics. Committee on G. American Academy of Pediatrics: Health supervision for children with Down syndrome. Pediatrics Feb 2001;107(2):442-449.
- 136. Peadon E, O'Leary C, Bower C, Elliott E. Impacts of alcohol use in pregnancy--the role of the GP. Aust Fam Physician Nov 2007;36(11):935-939.
- 137. Brennan D, Giles S. Ocular Involvement in Fetal Alcohol Spectrum Disorder: a review. Curr Pharm Des Feb 5 2014.
- 138. Davies L, Dunn M, Chersich M, et al. Developmental delay of infants and young children with and without fetal alcohol spectrum disorder in the Northern Cape Province, South Africa. Afr J Psychiatry (Johannesbg) Sep 2011;14(4):298-305.
- 139. Stephen JM, Kodituwakku PW, Kodituwakku EL, et al. Delays in auditory processing identified in preschool children with FASD. Alcohol Clin Exp Res Oct 2012;36(10):1720-1727.
- 140. de Beer M, Kritzinger A, Zsilavecz U. Young children with fetal alcohol spectrum disorder-communication profiles. S Afr J Commun Disord Dec 2010;57:33-42.
- 141. Kalberg WO, Provost B, Tollison SJ, et al. Comparison of motor delays in young children with fetal alcohol syndrome to those with prenatal alcohol exposure and with no prenatal alcohol exposure. Alcohol Clin Exp Res Dec 2006;30(12):2037-2045.
- 142. Whaley SE, O'Connor, Mj, Gunderson B. Comparison of the adaptive functioning of children prenatally exposed to alcohol to a nonexposed clinical sample. Alcohol Clin Exp Res Jul 2001;25(7):1018-1024.
- 143. Sood B, Delaney-Black V, Covington C, et al. Prenatal alcohol exposure and childhood behavior at age 6 to 7 years: I. dose-response effect. Pediatrics Aug 2001;108(2):E34.
- 144. Jirikowic T, Olson HC, Kartin D. Sensory processing, school performance, and adaptive behavior of young school-age children with fetal alcohol spectrum disorders. Phys Occup Ther Pediatr May 2008;28(2):117-136.
- 145. Alex K, Feldmann R. Children and adolescents with fetal alcohol syndrome

- (FAS): better social and emotional integration after early diagnosis. Klin Padiatr Mar 2012;224(2):66-71.
- 146. Stevens SA, Nash K, Fantus E, Nulman I, Rovet J, Koren G. Towards identifying a characteristic neuropsychological profile for fetal alcohol spectrum disorders. 2. Specific caregiver-and teacher-rating. J Popul Ther Clin. Pharmacol 2013;20(1):e53-62.
- 147. Pruett D, Waterman EH, Caughey AB. Fetal alcohol exposure: consequences, diagnosis, and treatment. Obstet Gynecol Surv Jan 2013;68(1):62-69.
- 148. Mattson SN, Roesch SC, Glass L, et al. Further development of a neurobehavioral profile of fetal alcohol spectrum disorders. Alcohol Clin Exp Res Mar 2013;37(3):517-528.

APPENDIX I: Anticipatory Guidance for Children and Adolescents with FASD (modeled after Appendix I from 135)

													Late	Adolescence
	Prenatal	Infancy Birth to 1 y				Early Childhood, 1 to 5 y				Childhood	(14 to 21 y)			
													(6 to 13 y)	
m = months; y =		Neo-	2 m	4 m	6 m	9 m	12 m	15 m	18 m	24 m	3 y	4 y		
years		natal												
Maternal Alcohol	S		S - sc	reen as	require	d for re	ferral an	d/or trea	tment re	equired;	intervi	ew/infor	mation gather	ring for
Use Screening		mu	ltidiscij	olinary	FASD a	assessm	ent shou	ld be do	ne by ap	propria	tely train	ned indi	vidual workin	g with team
Diagnosis		•	•	•	•	•	•	•	•	•	•	•	•	•
Medical Evaluation														
Growth (Length, weight, head circumference)	0	0	0	0	0	0	0	0	0	0	0	0	0	0
Hearing Screening		S/o	S/o	S/o	S/o	S/o	S/o	S/o	S/o	S/o	S/o	S/o	S/o	S/o
Vision Screening		S/o	S/o	S/o	S/o	S/o	S/o	S/o	S/o	S/o	S/o	S/o	S/o	S/o
ARBD (complete review of systems, e.g., cardiac, musculoskeletal, spina bifida etc.)		0	0	0	0	0	0	0	0	0	0	0	0	0
Seizures		S/o	S/o	S/o	S/o	S/o	S/o	S/o	S/o	S/o	S/o	S/o	S/o	S/o

DCD										S	S/o	S/o	S/o	S/o
Toileting/Enuresis/									S	S	S	S	S	
Encopresis														
Eating/Feeding		S/o	S/o	S/o	S/o	S/o	S	S	S	S	S	S		
Sleeping		S	S	S	S	S	S	S	S	S	S	S	S	S
Sexuality													S	S
Psychiatric/														
Behavioural														
ADHD											S/o	S/o•	•	•
Mood disorder	S/o										S/o	S/o	S/o	•
(anxiety/depression)														
ASD									S/o •					
Substance Use	S/o												S/o	S/o
Psychosocial														
Development		S/o	S/o	S/o	S/o	S/o	S/o							
Behaviour		S/o	S/o	S/o	S/o	S/o	S/o							
Preschool/School										0	0	0	0	0
Performance														
Socialization/Peers										0	0	0	0	О

Referrals (at any appointment as													
needed)													
SLP						•	•	•	•	•	•	•	•
OT	•	•	•	•	•	•	•	•	•	•	•	•	•
PT	•	•	•	•	•	•	•	•	•	•	•	•	•
Early Intervention	•	•	•	•	•	•	•	•	•	•	•	•	•
FASD Family	•	•	•	•	•	•	•	•	•	•	•	•	•
Support													
FASD Community	•	•	•	•	•	•	•	•	•	•	•	•	•
Support													
Justice												•	•

tion Disorder; OT = Occupational Therapist; PT = Physiotherapist; SLP = Speech & Language Pathologist.

TABLE 1 Anticipatory Recommendations by Age

	Physical Examination	Discuss and Review	Evaluate for	Anticipatory Guidance
Birth to 1 month	- Measure and plot growth	- Prenatal alcohol use	- Alcohol Related Birth Defects	- Self-regulation: Infants
	parameters (weight, length, head		(ARBD): cardiac defects,	with prenatal alcohol
	circumference), including growth	- Maternal substance use	gastrointestinal defects, cleft lip and	exposure and/or other
	at birth using appropriate growth	patterns including screening	palate, orthopedic anomalies (e.g.,	prenatal substance exposure
	curves	for current substance misuse	radioulnar synostosis), spina bifida -	may have difficulty
		using screening tool such as	see Table 2 from ¹³⁶)	regulating their body
	- Complete dysmorphology	CRAFFT or modified CAGE		temperature, movements, and
	examination. This may be	(T-ACE and TWEAK) ⁶	- Growth retardation, especially	reactions to their
	conducted by geneticist/		symmetric growth retardation	environment. They may be
	dysmorphologist experienced in	- Mental health screening for		jittery, irritable, and show
	FASD. Palpebral fissure lengths	depression, post-partum	- Vision (e.g., optic nerve	tone abnormalities. Infants
	(PFL) should be plotted on PFL	depression or other mental	hypoplasia, abnormal retinal vessels,	may respond to swaddling,
	curves. An assessment of lip	health disorders	coloboma, microphthalmia ¹³⁷)	calm quiet environments, and
	fullness and development of		,	patient well-regulated
	philtrum should be conducted	- Social screening for risk	- Hearing: universal newborn hearing	caregiving. Parents may
	using the Washington lip-	factors such as intimate	recommended	benefit from counselling and
	philtrum guide.	partner violence, mental		respite care.
		health disorder such as	- Newborn metabolic screening	
	- Complete systems exam	depression, and poverty		- Sleep: Further evidence of
	including but not limited to		- Iron deficiency ¹³²	abnormal self-regulation may
	intraoral/palate, vision, hearing,			be poorly regulated sleep,
	skin (e.g., hemangiomas, other		- Microcephaly: if present,	reduced sleep, or disrupted
	birthmarks, neurocutaneous		recommend head ultrasound and	sleep patterns. Parents may
	abnormalities), cardiac (septal		consider MRI	benefit from sleep
	defects, patent ductus arteriosus,			consultation and respite.
	other), orthopedic/skeletal (e.g.,		- Seizure disorder	•
	congenital hip dislocation,			- Feeding: Infants may have
	scoliosis, limb/digit anomalies),			poorly coordinated suck and
	genitourinary (e.g.,			swallow, regurgitate more
	hydronephrosis, labial hypoplasia,			frequently. They benefit from
	sphincter tone, tufts), neurological			patient responsive feeding

I	(e.g., head shape, microcephaly, focal abnormalities, tone abnormalities)			practices. Parents may need consultation from feeding specialists and occupational therapists.
Infancy	- Measure and plot growth parameters (weight, length, head circumference), including growth parameters at birth using appropriate growth curves. - Complete dysmorphology examination. This may be conducted by geneticist/ dysmorphologist experienced in FASD. PFL should be plotted on PFL curves. An assessment of lip fullness and development of philtrum should be conducted using the Washington lip-philtrum guide. - Complete systems exam, including oral/dental/palate, vision, hearing, cardiac, hips, orthopedic/skeletal abnormalities, genitourinary, neurologic (e.g., abnormalities of tone, persistent primitive reflexes, motor deficits) - Complete developmental assessment looking for delays in expressive and receptive language, problem solving delays, fine and gross motor delays, social and emotional delays, self-regulatory delays	- Developmental milestones and behaviour 138 - Parent child relationship and stressors - Social behaviour and development - Parenting stress and sources of support	- ARBD as above - Vision – refer for formal ophthalmologic evaluation if any visual concerns ¹³⁷ - Hearing – refer for formal audiologic evaluation if newborn screen has not been done, if there are any speech or language delays, or if there are any concerns regarding hearing or speech development ¹³⁹ - Developmental delay ¹³⁸ – refer for formal developmental assessment - Iron deficiency, other nutritional deficiency	- Behavioural disturbance including difficulties with self-regulation, disorganization, emerging hyperactivity, emotional dysregulation - Discuss attachment behaviours and healthy parent child interactions; refer for supports in high risk groups and as needed - Feeding disturbances including food selectivity and atypical feeding behaviours - Screen for sleep disturbances - Daycare: issues related to sensitivity to environmental change, behavioural management - Discuss early intervention supports in the community including developmental supports, mental health services, and therapy supports including physiotherapy, occupational therapy, and speech and language therapy

				- Caregiver support and respite
1 to 5 years: Early Childhood	- Continue to monitor growth parameters (weight, length, head circumference) using appropriate growth curves. - Complete systems exam, including oral/dental/palate, vision, hearing, cardiac, hips, orthopedic/skeletal abnormalities, genitourinary, neurologic (e.g., abnormalities of tone, persistent primitive reflexes, motor deficits), and minor dysmorphic features particularly in the child who has previously been assessed as non-dysmorphic as minor findings may still be found - Complete developmental assessment looking for delays in expressive and receptive language, problem solving delays, fine and gross motor delays, social and emotional delays, self-regulatory delays – follow-up should include a pre-kindergarten developmental assessment for the purpose of informing school supports and programming	- Developmental milestones and behaviour ^{138,140-142} - Parent child relationship and stressors - Social behaviour and development ^{142,143} – risk for internalizing and externalizing behaviours - Sensory differences and influence on self-regulation/behaviour, sleep, eating, toileting ^{118,144} - Parenting stress and support	- Vision – refer for preschool vision screening - Hearing – refer for formal audiologic screening/ evaluation if newborn screen has not been done, if there are any speech or language delays, or if there are any concerns regarding hearing or speech development 139 - Developmental delay 138,140 – refer for formal developmental assessment - Iron deficiency, other nutritional deficiency	- Discuss school readiness and support early intervention programs aimed at academic and social emotional health - Discuss neurobehavioural differences in learning and social behaviour and need for continued and increased supports with age as adaptive gaps widen 142,145 - Education re: behaviour in context of prenatal alcohol exposure "reframing behaviour" 146-148 - Education and support re: sleep disorder – refer for consultation as needed - Discuss need for caregiver support and respite - Advocate for educational and community based supports for child
6 to 13 years: Late Childhood	- Continue to monitor growth parameters (e.g., weight, length, head circumference) using appropriate growth curves.	- Evolution of comorbid diagnoses such as ADHD, ODD and Conduct Disorder Pharmacological interventions	 Vision – refer for preschool vision screening Hearing – refer for formal audiologic screening/ evaluation if 	- Collaborate with education system, social services, and other service providers that have contact with the patient

	- Complete systems exam, including oral/dental/palate, vision, hearing, cardiac, hips, orthopedic/skeletal abnormalities, genitourinary, neurologic (e.g., abnormalities of tone, persistent primitive reflexes, motor deficits), and minor dysmorphic features particularly in the child who has previously been assessed as non-dysmorphic as minor findings may still be found - Assess Tanner Stage and evolving pubertal development - Comprehensive neuropsychological assessment looking for delays and/or deficits in the following domains: Motor; Neurophysiology/Neuroanatomy; Cognition; Communication; Academic Achievement; Memory; Attention; Executive Function; Anxiety and Depression; Adaptive Behaviour, Social Skills and Social Communication	- Early presentations of mental health issues such as depression, low self-esteem, anxiety and mood regulation Pharmacological interventions - Determinants of health (housing, food security) - Parent child relationship and stressors - Social behaviour and development 142,143 - risk for internalizing and externalizing behaviours - Sensory differences and influence on self-regulation, sleep, eating 118,144 - Parenting stress and support - New contact with the justice system - Understanding of puberty, sexual activity, contracention	newborn screen has not been done, if there are any speech or language delays, or if there are any concerns regarding hearing or speech development 139 - Cognitive disability or learning difficulties/disability – refer for formal psychoeducational assessment - Iron deficiency, other nutritional deficiency Evidence of abuse (physical, emotional, sexual) - Refer for follow-up based on physical findings, when indicated - Existence of trusting and stable relationships and/or primary advocate for the individuals (caregiver) - Drug and/or alcohol abuse and subsequent referral to appropriate treatment centres and programs	- Identify potential advocates and/or mentors, if needed - Discuss neurobehavioural differences in learning and social behaviour and need for continued and increased supports with age as adaptive gaps widen 142,145 - Education re: behaviour in context of prenatal alcohol exposure "reframing behaviour" 146-148 - Advocate for educational and community based supports for child - Drug and/or alcohol education - Sex education, personal safety/abuse, and contraception when needed
		sexual activity, contraception		
13-21 years or older: Adolescence to Early Adulthood	- Depending on the age and cognitive capacity of the teen, choose to do the exam with or without the caregiver present, include teen in the decision.	- Transition to adulthood -School transitions to higher grades and need for assessment and supports that may not have been	Vision and hearing Oral health as many teens have significant malocclusions needing orthodontic work	- Discuss physical health needs, and pubertal changes - Discuss medication including transitioning to independence, medication
	- Corroborate information from the teen in a separate interview	previously identified	- Blood work for iron deficiency, other nutritional deficiencies based	preferences (e.g., stimulants), side effects of medications.

with the caregiver, keeping in
mind how long they have had a
relationship.

- Continue to monitor growth parameters (weight, height, head circumference) using appropriate growth curves. Note growth impact of prescribed medications: weight loss with stimulants, excessive weight gain with atypical antipsychotics and antidepressants
- Complete systems exam, especially if assessment for birth defects or acquired health conditions have not been previously conducted. Include oral/dental/palate, vision, hearing, cardiac, orthopedic/skeletal abnormalities, genitourinary, neurologic (e.g., abnormalities of tone, balance and coordination with eyes open and closed, motor planning, soft signs such as past pointing on finger nose testing), and minor dysmorphic features, Tanner staging of pubertal development with note of gynecomastia in males treated with atypical antipsychotics.
- Complete developmental/mental status assessment by history and current report: Previous mental health diagnosis and interventions (ADHD is the

- Current behavioural and mental health concerns and management
- Sleep hygiene
- Nutrition and self care
- Caregiver child relationship and stability of placement
- Caregiver stress and supports, knowledge about FASD and role as advocate in transition years to adulthood

on diet history

- Monitor side effects of medications, if applicable
- Blood work for fasting insulin and glucose, prolactin, thyroid function with atypical antipsychotics; electrocardiograms for arrhythmia in teens with positive family history or taking stimulants
- History to suggest seizures and need for electroencephalography
- Sexual activity and knowledge about pregnancy
- Substance use
- Current and merging concerns about mental health

- Discuss sexual health, especially around contraception for both male and female teens to prevent unplanned pregnancy and sexually transmitted diseases.
- The harm of alcohol use in pregnancy needs to be emphasized repeatedly and at a level understood by the teen.
- Discuss mental health concerns and refer to an Adolescent Psychiatrist or mental health therapist as needed
- Trauma-informed therapy might be indicated.
- Advocate for education supports and, if needed, an updated assessment of function
- Advocate for community supports and high level of supervision to ensure safe and meaningful participation and prevention of victimization, as appropriate
- Discuss with caregiver(s) the process of transition to adult services. Current guidelines recommend that

most common comorbidity in	this start at the age of 12
FASD).	years with ongoing
Past history of maltreatment or	assessment of the teens'
trauma, number of placements,	ability to become independent
relationship to current	in decision making, shared
caregiver(s).	decision-making or the need
Impairments in communication	for legal guardianship as an
(not understanding what is	adult. Documentation and
requested, responses that are	paperwork will be different in
vague, not connected, tangential	different jurisdictions. This
and immature)	may require navigation of the
Problems in the area of executive	system by an informed person
function (e.g., problem solving,	or trustee to ensure safe
inhibition, flexible thinking,	living, financial management,
predicting, using judgment)	employment options and
Social and emotional difficulties	ongoing maintain of health
(including bullying or being	care. Special time allocations
victimized by others, positive	needed to be considered for
participation with peers in	these visits.
community-based and sports	
activities)	- Teens living with FASD
Self-regulatory difficulties with	need to be informed about
externalizing behaviours of	their diagnosis and need for
aggression or internalizing	life long mentors and
behaviours of low self- esteem	supports.
and anxiety	
Sexual activity including past or	- Timing and process to be
present history of abuse	determined in consultation
Substance use	with primary caregiver. Care
Suicidal ideation or attempts	is needed to prevent blaming.
Sleep pattern and diet	
School experiences (i.e., extra	- Discuss need for caregiver
supports in the classroom,	support and respite,
cognitive and academic	especially if there are
assessments)	increasing mental health
Ability to perform daily living	issues for the teen and stress
and self-care activities	within the family
independently.	

			- Advocate for the teen to enter foster or group home
			care, where they can develop
			positive and trusting relationships and set realistic
			goals for the future. Care
			agreements can be extended
			to age 21 years in some jurisdictions to plan the
			transition process to adult
			services.
			- Transfer of medical and
			mental health care to the adult
			system needs to be planned
			with appropriate care
			summaries and training of the adult health care providers on
			FASD.
			1 1 1 1 EAGE
			 Young adults with FASD may become parents
			themselves and there needs to
			be consideration of their
			ability to parent and what
			supports they will need to be able to care for their children.
ADHD = Attention defi	cit/Hyperactivity Disorder: ODD = Oppo	sitional Defiant Disorder	able to care for their efficient.