Section A: Assessing maternal alcohol use

The timing, frequency and quantity of prenatal alcohol exposure (PAE) are linked to the pattern and severity of fetal outcomes, but may not be available or reliable. (4, 18-21) In addition, both maternal and fetal characteristics are associated with variability in alcohol-related outcomes. Brain growth and development occur throughout pregnancy hence adverse cognitive, behavioural and neurodevelopmental outcomes may result from exposure at any time during pregnancy and may occur in the absence of facial anomalies or structural central nervous system abnormalities. (22)

It is likely that multiple mechanisms are involved in damage to the brain from PAE and **no** 'safe' threshold for alcohol consumption during pregnancy has been established. (23) Although there is limited evidence associating low levels of prenatal alcohol exposure with risks to human fetal development, (24) the Australian Guide to Reduce Health Risks for Drinking Alcohol(10) states that maternal alcohol consumption can harm the developing fetus and recommends that for women who are pregnant or planning a pregnancy, not drinking is the safest option(10).

The level of risk to the fetus from prenatal alcohol exposure is highest when there is high, frequent maternal alcohol intake. The level of risk for the fetus is likely to be low if a woman has consumed only small amounts of alcohol (such as one or two drinks per week) before she knew she was pregnant or during pregnancy.(10)

A diagnosis of FASD is not appropriate where there is *confirmed absence* of prenatal alcohol exposure, but a diagnosis of FASD with three sentinel facial features can be made when prenatal alcohol exposure is unknown (see Table 1). (3)

Assessment of prenatal alcohol exposure requires clinical judgement and careful evaluation of a range of information that may provide confirmation of maternal alcohol use and allow quantification of intake.

Evidence of confirmed prenatal alcohol exposure may include:

- Information reported by the birth mother about her alcohol consumption during the index pregnancy, ideally using a validated tool;
- Reports by others, including a relative, partner, household or community member who
 had direct observation of drinking during the index pregnancy; or
- Documentation in child protection, medical, legal or other records of maternal alcohol consumption, alcohol-related disorders, and problems directly related to drinking during the index pregnancy, including alcohol-related injury and intoxication.

Assessing the reliability of evidence:

 If recalled information from different informants is in direct conflict (confirmed absence and confirmed presence) and reliable information on exposure is not available, alcohol exposure should be recorded as unknown. (4)

- The reliability of information on prenatal alcohol exposure may reflect the timing of pregnancy awareness.
- A history of alcohol dependence without evidence of consumption during the index pregnancy is not sufficient to indicate confirmed exposure but should raise suspicion of risk.(3, 4)

Alcohol Use Disorders Identification Test - Consumption (AUDIT-C)

When detailed information on maternal alcohol use is available, consumption during pregnancy should be assessed using the AUDIT-C questions(25) as included on the *Australian FASD Diagnostic Assessment Form* (Appendix A1) and reproduced in Table 2.

The AUDIT-C questions provide a standardised method for the assessment of maternal alcohol use and are based on a validated sex-specific version of the instrument.(26, 27) The use of a sex-specific threshold of 5 or more drinks on one occasion for question 3 of the AUDIT-C reflects known levels of maternal alcohol consumption associated with increased risk of FASD and other harms.(10, 28, 29) Five or more drinks on an occasion (consumption of 50+ g of alcohol) is sometimes referred to as a binge.(29)

Derivation of the AUDIT-C score, although not essential for diagnosis, allows the clinician to categorise the **level of fetal risk associated with maternal drinking**.

Information on the definition of a standard drink for different types of alcoholic drinks should be provided prior to using the AUDIT-C. Appendix B shows standard drink sizes for commonly consumed drinks. A complete guide is available at: http://www.alcohol.gov.au/internet/alcohol/publishing.nsf/Content/drinksguide-cnt)

Some guiding principles for taking an alcohol history in pregnancy:

A non-judgemental approach is important when taking a history of alcohol consumption in pregnancy.

Some factors to consider:

- A pregnancy may be unplanned and not confirmed for some time, during which time alcohol may have been consumed;
- A woman may have made lifestyle changes once the pregnancy was confirmed, including reducing or stopping alcohol consumption;
- A woman may be unaware that not drinking during pregnancy is the 'safest' option and may have been given incorrect advice by other health professionals;
- Women may be more likely to drink if their partner and household members also drink and this may be explored.

Some questions to begin history taking:

- Was the pregnancy planned or unplanned?
- When did the birth mother realise that she was pregnant?
- Did the birth mother modify her drinking behaviour on confirmation of pregnancy?
- Were there any special occasions (e.g. a wedding) during pregnancy when alcohol was consumed at a high level?

Evidence of maternal alcohol use in the three months prior to and during pregnancy should be assessed, including any special occasions when a large amount of alcohol may have been consumed. The definition of a standard drink should be explained prior to administering the AUDIT-C (Q1-3), using the Standard Drinks Guide (Appendix B).

Table 2 Reported alcohol use, including AUDIT-C Questions

Alcohol usc	in early pregnance	cy (if available)						
Was the pregnancy planned or unplanned? □ Planned □ Unplanned □ Unknown								
When did the	e birth mother realis	se that she was pr	egnant?((weeks)	☐ Unkno	own		
Did the birth	mother drink alcoh	ol before the preg	nancy was confirme	ed?	☐ Yes	□ No	☐ Ur	nknown
	mother modify her ase specify:	drinking behaviou	ır on confirmation c	of pregnancy?	□ Yes	□No	□Un	known
During which	n trimesters was alco	ohol consumed? (t	cick one or more)	□ None	□ 1 st	☐ 2 nd	\square 3 rd	□ Unknown
AUDIT-C qu	estions							
Source of rep	oorted information (on alcohol use:	☐ Birth mother	☐ Other (plea	se specify)			
1. How often	did the birth mothe	er have a drink cor	ntaining alcohol dur	ing this pregnar	ncy?			
Unknown	Never							
		Monthly	2-4 times	2-3 times	4 or r	more times		
	[skip Q2+Q3]	or less	2-4 times a month	2-3 times a week		more times week		
		,						
	[skip Q2+Q3]	or less \square_1	a month \square_2	a week □3	ā	a week □ ₄	pregnand	cy?
	[skip Q2+Q3] □ ₀	or less \square_1	a month \square_2	a week □3	as drinking (a week □ ₄	pregnand	cy?
2. How many	[skip Q2+Q3] □ ₀ v standard drinks dic	or less \Box_1 d the birth mother	a month \Box_{2} have on a typical d	a week □₃ ay when she wa	as drinking (week \Box_4 during this	pregnand	cy?
2. How many Unknown	[skip Q2+Q3] \Box_0 standard drinks did 1 or 2	or less \Box_1 d the birth mother \Box_1 \Box_1	a month \Box_2 have on a typical d \Box_2	a week \square_3 ay when she wa 7 to 9 \square_3	as drinking (10	a week \Box_4 during this or more \Box_4	pregnand	cy?
2. How many Unknown	[skip Q2+Q3] \Box_0 standard drinks did 1 or 2 \Box_0	or less \Box_1 d the birth mother \Box_1 \Box_1	a month \Box_2 have on a typical d \Box_2	a week \square_3 ay when she wa 7 to 9 \square_3	as drinking o 10 uring this p	a week \Box_4 during this or more \Box_4	pregnand	cy?
2. How many Unknown 3. How often	[skip Q2+Q3] \Box_0 y standard drinks did $1 \text{ or } 2$ \Box_0 a did the birth mother	or less \Box_1 d the birth mother \Box_1 \Box_1 er have 5 or more	a month \Box_2 have on a typical d \Box_2 standard drinks on	a week □3 ay when she wa 7 to 9 □3 one occasion du	as drinking o 10 uring this p	week \Box_4 during this or more \Box_4 regnancy?	pregnand	cy?
2. How many Unknown 3. How often	[skip Q2+Q3] \Box_0 y standard drinks did $1 \text{ or } 2$ \Box_0 a did the birth mother	or less \Box_1 d the birth mother \Box_1 \Box_1 er have 5 or more Less than	a month \Box_2 have on a typical d \Box_2 standard drinks on	a week □3 ay when she wa 7 to 9 □3 one occasion du	as drinking o 10 uring this p	a week during this or more 4 regnancy?	pregnand	cy?

Assessing prenatal alcohol exposure: Summary

Assessment of prenatal alcohol exposure requires clinical judgement and careful evaluation of a range of information that may provide confirmation of maternal alcohol use and quantification of intake.

Evidence of exposure can be evaluated to estimate the overall level of risk using the following broad risk categories:

- i. **No exposure** (confirmed absence), no risk of FASD;
- ii. Unknown exposure (alcohol use is unknown);
- **iii. Confirmed exposure** (AUDIT-C score =1-4; or confirmed use, but exposure less than high risk level for FASD; or confirmed use, but not known if exposed at a high risk level for FASD); and
- iv. **Confirmed-high risk exposure** (AUDIT-C score = 5+; confirmed use, exposure at high risk level for FASD).

Confirmed high risk exposures for FASD can be considered to include, at any time during pregnancy:

- i. An AUDIT-C score of 5 or more
- ii. Reported consumption of **5 or more standard drinks on one occasion** (e.g. AUDIT-C question 3)
- iii. Other reliable evidence of high consumption

Other prenatal and post-natal exposures

Neurodevelopment impairment observed among individuals being assessed for FASD may be associated with exposures other than alcohol. It is important to determine whether any observed impairments can be explained by other causes or events (e.g. prenatal complications, genetic factors including chromosomal abnormalities, head injuries, early life trauma (including social and emotional abuse), problems with vision or hearing, or substance abuse by the patient).

All relevant prenatal and postnatal exposures or events, including prenatal exposure to prescription and non-prescription drugs, should be documented during the diagnostic assessment, and evaluated based on their likely influence. Other exposures should be considered when determining the appropriate diagnosis and management plan.

There may not be a single explanation for the observed neurodevelopmental impairment, and it is important that the diagnostic assessment process considers the effects of other adverse prenatal and postnatal exposures. (3)

In addition to vision and hearing testing, other clinically indicated investigations may include chromosome microarray analysis and Fragile X testing, and other tests such as full blood count, ferritin, vitamin B_{12} , metabolic screen, creatinine kinase, lead, and thyroid function.

Table 1 Diagnostic criteria and categories for Fetal Alcohol Spectrum Disorder (FASD)

FETAL ALCOHOL SPECTRUM DISORDER								
	Diagnostic categories							
Diagnostic criteria	FASD with 3 Sentinel Facial Features	FASD with < 3 Sentinel Facial Features						
Prenatal alcohol exposure	Confirmed or unknown	Confirmed						
Neurodevelopmental domains - Brain structure/Neurology - Motor skills - Cognition - Language - Academic Achievement - Memory - Attention - Executive Function, including impulse control and hyperactivity - Affect Regulation - Adaptive Behaviour, Social Skills or Social Communication	Severe impairment in at least 3 neurodevelopmental domains	Severe impairment in at least 3 neurodevelopmental domains						
Sentinel facial features - Short palpebral fissure - Smooth philtrum - Thin upper lip	Presence of 3 sentinel facial features	Presence of 0, 1 or 2 sentinel facial features						

Key components of the FASD diagnostic assessment include documentation of:

- History presenting concerns, obstetric, developmental, medical, mental health, behavioural, social;
- Birth defects dysmorphic facial features, other major and minor birth defects;
- Adverse prenatal and postnatal exposures, including alcohol;
- Known medical conditions including genetic syndromes and other disorders;
- Growth

Infants and young children under 6 years of age and older adolescents and adults warrant special consideration during the FASD diagnostic assessment process. (16) There are also circumstances where an individual may be considered to be 'at risk' of FASD. These special clinical considerations are discussed in detail in Section B: Neurodevelopmental Impairment.

☐ Female	☐ Male	□ Oth	er
/ /	Age at ass	essment:	
☐ Birth mother ☐ Foster carer ☐ Other			
☐ Department of C	hild Protection	☐ Juvenile justice	☐ Not applicable
□ No	☐ Yes		
/ /			
regiver, teacher; strengths	and needs; age-app	oropriate abilities e.g. beh	avioural regulation,
	/ / Birth mother Foster carer Other Department of C	/ / Age at ass	/ / Age at assessment: Birth mother

Obstetric history:
Developmental history:
Mental health and other behavioural problems:
Patient's medical history:
Social history: e.g. foster care, living arrangements.

MATERNAL ALCOHOL USE

Evidence of maternal alcohol use in the three months prior to and during pregnancy should be assessed, including any special occasions when a large amount of alcohol may have been consumed. The definition of a standard drink should be explained prior to administering the AUDIT-C (Q1-3). A Standard Drinks Guide can be downloaded.

http://www.health.gov.au/internet/alcohol/publishing.nsf/Content/drinksguide-cnt

	Alcohol use in early pregnancy (if available)									
a. Was the pregnancy planned or unplanned? □ Planned □ Unplanned □ Unknown										
b. At what gestation did the birth mother realise that she was pregnant?(weeks) Unknown										
c. Did the birth mother drink alcohol before the pregnancy was confirmed?										
d. Did the birth mother modify her drinking behaviour on confirmation of pregnancy? Yes Unknown										
	If Yes please specify: e. During which trimesters was alcohol consumed? (tick one or more) □ None □ 1st □ 2nd □ 3rd □ Unknown									
AUDIT-C Reported alcohol use (if available)										
1. How often d	id the birth mother ha	ive a drink containi	ng alcohol during this p	regnancy?						
Unknown	Never	Monthly	2-4 times	2-3 times	4 or more times					
	[skip Q2+Q3]	or less	a month	a week	a week					
	\square_0	\square_1	\square_2	\square_3	\square_4					
2. How many st	tandard drinks did the	birth mother have	on a typical day when s	she was drinking d	luring this pregnancy?					
Unknown	1 or 2	3 or 4	5 or 6	7 to 9	10 or more					
	\square_0	\square_1	\square_2	\square_3	\square_4					
					_					
			ard drinks on one occas		· .					
Unknown	Never	Less than	Monthly	Weekly	Daily or					
		monthly			almost daily					
	\square_0	\square_1	\square_2	\square_3	□4					
AUDIT-C score	during this pregnance	y: (Q1+Q2+Q3)=								
		Scores: 0=No ex	posure 1-4= Confirme	d exposure 5+= C	Confirmed high-risk exposure					
Other evidence	e of exposure									
Is there eviden	ce that the birth moth	er has ever had a p	roblem associated with	alcohol misuse o	r dependency?					
□ No □ Yes	(identify below, including	source of information)								
☐ Alcohol depe	endency (specify)									
☐ Alcohol-rela	ted illness or hospitali	sation (specify)								
☐ Alcohol-rela	ted injury (specify)									
☐ Alcohol-rela	ted offence (specify)									
☐ Other (specify	')									
Information from records: e.g. medical records, court reports, child protection records.										
l <u> </u>	Is there evidence that the birth mother's partner has ever had a problem associated with alcohol misuse or dependency? No Yes (identify below, including source of information)									
Alcohol exposu	ire summary									
Source of repor	rted information on a	cohol use:	rth mother	r (specify)						
In your judgem	ent what is the reliab	lity of the informat	ion on alcohol exposure	e: 🗆 Unknown	☐ Low ☐ High					
In your judgem	ent was there high-ris	k consumption of a	lcohol during pregnanc	y? 🗆 Unknown	☐ Yes ☐ No					
Prenatal alcoho	ol exposure: 🗆 Unkno	own exposure 🗆 N	o exposure Confirm	ned exposure 🗆 (Confirmed-high risk exposure					

OTHER EXPOSURES

Postnatal risk summary:

 \square No known risk

Assess evidence of adverse prenatal and pos	stnatal exposures and events	that need to be consider	ea.
Prenatal			
Other prenatal exposures identified: (if yes, sp	pecify and indicate source of informa	ation)	
☐ Nicotine (e.g. cigarettes, inhalers, e-cigs and chew	ved tobacco) (specify)		
☐ Marijuana (specify)			
☐ Heroin (specify)			
☐ Cocaine (specify)			
☐ Amphetamines (specify)			
☐ Other non-prescription drugs (specify)			
☐ Anti-convulsants (specify)			
☐ Other prescription drugs (specify)			
☐ Don't know			
□ None			
Specify other prenatal risk factors and asses including ionizing radiation, paternal or maternal intelle			exposure to known teratogens,
Other prenatal risk summary:			
. □ No known risk	☐ Unknown risk	☐ Some risk	☐ High risk
Postnatal			
Specify other physical or medical risk factors or neglect, serious head injury, meningitis or other specify other psychosocial risk factors and a	er medical conditions that lead to	o brain damage, child substa	ance abuse)

 \square Unknown risk

 \square High risk

 \square Some risk

GROWTH

Assess birth parameters and postnatal growth, and determine if any deficit exists that is unexplained by genetic potential, environmental influences (e.g. nutritional deficiency) or other known conditions (e.g. chronic illness).

Birth	Gestational age		Birth length		Birth	n weight
Date	weeks		cm	percentile	grams	percentile
	–		_			1
Growth reference	chart used: CDC		□ WHO	☐ Other (specify)		
Postnatal			He	eight	W	eight /
Г	1			T		
Date	Age		cm	percentile	kg	percentile
		•			•	•
Growth reference	chart used: CDC		□ WHO	☐ Other (specify)		
Parental height (if a	available)					
Mother's height (cm) Father's heigh	t (cm)	Sex-specific	target height (cm)	Sex-specific targe	t height (percentile)
drugs, nicotine)	t may explain growth		eror (e.g. macm		negreet, generie contain	on, prematant, oure
Growth summary						
Was an unexplaine	d deficit in height or v	veight <	3 rd percentile	identified at any time	? □ Yes □ No	
If Yes □ height o	r weight ≤10 th and >3 ^r	^d percent	ile 🗆 height	or weight ≤3 rd percent	ile	

SENTINEL FACIAL FEATURES

Assess for the 3 sentinel facial features of Fetal Alcohol Spectrum Disorder: short palpebral fissure length (2 SD or more below the mean), smooth philtrum (rank 4 or 5 on the Lip-Philtrum guide), and thin upper lip (rank 4 or 5 on the Lip-Philtrum guide).

		I (DEL)		Righ	t PFL	Left	PFL	Ме	an PFL
-	issure Lengt				1		1		1
Date	Age	Assessment i	nethod	mm	Z score (SD)	mm	Z score	mm	Z score*
		☐ direct measu	re \Box photo analy	/sis					
		☐ direct measu	re 🔲 photo analy	/sis					
PFL referen	ce chart use	d: 🗆 Stro	omland \Box C	larren 🗆 (Other				
Philtrum									
Date	Age	Assessment r	nethod			UW Lip-Ph	iltrum Guid	e 5-point r	ank
		☐ direct measu	re	/sis					
		☐ direct measu	re 🔲 photo analy	/sis					
		☐ direct measu	re \Box photo analy	/sis					
Upper lip					-				
Date	Age	Assessment r	nethod			UW Lip-Ph	iltrum Guid	e 5-point r	ank
		☐ direct measu	re 🔲 photo analy	/sis					
		☐ direct measu	re	/sis					
		☐ direct measu	re 🔲 photo analy	/sis					
Lip-Philtrur	n Guide [†] use	ed: 🗆 Guid	e 1. Caucasian	☐ Guide	2. African	American			
Sentinel F	acial Featu	res Summar	y						
			PFL 2 SD or more b	elow the mear	, philtrum	rank 4 or 5	, upper lip r	ank 4 or 5)	:
		□ 0							
OTHER PHY	SICAL FINDI	NGS							
Dysmorphi	c facial featu	ures (please spe	cify)						
Othou binth	defeate w	aiau au mainau	/-l						
Other birth	i defects - m	ajor or minor	(please specify)						
Other med	ical conditio	ns:							
Hearing im	pairment:	□ No	☐ Not tested	☐ Yes (specify)				
Vision impa	irment:	\square No	\square Not tested	☐ Yes (specify)				
Known syn	drome or ge	netic disorder	(please specify):						
Other (plea	se specify):								
Investigation	ons:								
_	al microarray:	. □ No	☐ Result pending	☐ Yes (specify	result)				
Fragile X tes	•		☐ Result pending	☐ Yes (specific					
Other invest	igations as inc	dicated: Full blo	ood count, ferritin, m	etabolic screen,	creatinine k	inase, lead, a	and thyroid f	unction	

^{*}University of Washington Palpebral Fissure Length Z-score calculator: http://depts.washington.edu/fasdpn/htmls/diagnostic-tools.htm#pfl

[†]University of Washington Lip-Philtrum Guides: http://depts.washington.edu/fasdpn/htmls/lip-philtrum-guides.htm

NEURODEVELOPMENTAL DOMAINS

1 BRAIN STRUCTURE / NEUROLOGY DOMAIN

BRAIN STRUCTURE

Occipitofrontal Circumference (OFC)

Date	Age	OFC (cm)	Percentile*	Reference used					
Birth:									
*correct for gestational	age when < 2 years old	1	1						
If OFC < 3 rd percen	tile, is it explained b	by other aetiologies e	e.g. infection, metabolic	or other disease?					
No ☐ Yes (spe									
Imaging									
CNS imaging perfo	rmed: 🗆 No	☐ Yes (specify imag	ge modality and date)						
Specify any structu	ral abnormalities:								
If yes, are they exp	lained by other aeti	iologies e.g. injury, in	fection, or metabolic o	other disease? No Yes (specify)					
NEUROLOGY									
	seizure disorders o	r other abnormal hai	rd neurological signs.						
Seizure disorder	seizure disorders o	. other donormal har	a near ological olgilo.						
Seizure disorder pr	esent: No	☐ Yes (specify)							
If yes, are they exp	lained by other aeti	iologies e.g. injury, in	fection, or metabolic o	other disease? No Yes (specify)					
Other neurologica	l diagnoses e.g. cer	ebral palsy, visual im	pairment, sensorineura	l hearing loss					
Other abnormal ne	eurological diagnose	es present: \square No	☐ Yes (specify)						
If ves. are they exp	lained by other aeti	iologies e.g. iniurv. in	ifection, or metabolic o	r other disease?□ No □ Yes (specify)					
, , , , , , , , , , , , ,	, , , , , , , , , , , , , , , , , , , ,		,	,					
Brain Structure/ N	eurology domain sı	ummary							
Brain Structure/ Neurology domain summary Evidence of brain structure/neurology abnormalities of presumed prenatal origin that are unexplained by other causes?									
Evidence of Brain's	tructure/neurology	abnormalities of pre	esumed prenatal origin	that are unexplained by other causes?					

FUNCTIONAL NEURODEVELOPMENTAL DOMAIN SUMMARIES

Assess evidence of significant CNS dysfunction due to underlying brain damage. Required evidence includes severe neurodevelopmental impairment (2 SD or more below the mean or < the 3rd percentile) in domains of brain function based on standardised psychometric assessment by a qualified professional.

2.	ΝЛ	a	т	റ	D	c	vi		ıc
۷.	IVI	v	ı	u	n	3	N	ы	LJ

2. MOTOR SKILLS					
Test/subtest name		Age/ Date	Score	%ile/SD	Interpretation
Other information:					
Motor Skills impairment:	☐ None	☐ Some		Severe	☐ Not assessed
3. COCNUTION					
3. COGNITION		Ago/ Data	Cooro	%ile/SD	Interpretation
Test/subtest name		Age/ Date	Score	%ile/SD	Interpretation
Other information:					
Other mornation.					
Cognition impairments	☐ None	☐ Some		Severe	☐ Not assessed
Cognition impairment:		Joine		Severe	□ Not assessed
4. LANGUAGE					
(Expressive and Receptive)		1	r	1	T
Test/subtest name		Age/Date	Score	%ile/SD	Interpretation
Other information:					
Language impairment	☐ None	☐ Some		Severe	☐ Not assessed

5. ACADEMIC ACHIEVEMENT

5. ACADEIVIIC ACHIEVEIVIENT					
Test/subtest name		Age/ Date	Score	%ile/SD	Interpretation
Other information:					
Academic achievement impairment	□ None	□ Son		☐ Severe	□ Not accessed
Academic achievement impairment	☐ None	☐ 20II	1e	Severe	□ Not assessed
6. MEMORY					
Test/subtest name		Age /Date	Score	%ile/SD	Interpretation
Other information:					
Memory impairment	☐ None	☐ Som	e	☐ Severe	☐ Not assessed
, .					
7. ATTENTION			ı		
Test/subtest name		Age/ Date	Score	%ile/SD	Interpretation
Other information:					
Attention impairment	□ None	☐ Som	e	☐ Severe	☐ Not assessed

8. EXECUTIVE FUNCTION, INCLUDING IMPULSE CONTROL AND HYPERACTIVITY

Test/subtest name	Age/ Date	Score	%ile/SD	Interpretation	
				<u>`</u>	
Other information:	<u> </u>		<u> </u>		
Executive function, including impulse control and			_	_	
□ None	☐ Some	ġ] Severe	☐ Not assessed	
9. AFFECT REGULATION	<u> </u>	<u> </u>	<u> </u>	1	
Test/subtest name	Age/ Date	Score	%ile/SD	Interpretation	
Other information:					
Affect regulation impairment:	ie 🗆 Som	ie	☐ Severe	☐ Not assessed	
10. ADAPTIVE BEHAVIOUR, SOCIAL SKILLS, OR SO			0/11 /00		
Test/subtest name	Age/ Date	Score	%ile/SD	Interpretation	
Other information					
Other information:					
Adaptive behaviour, social skills, or social commu	nication impairm	ent			
□ None	☐ Some] Severe	☐ Not assessed	
NEURODEVELOPMENTAL DOMAINS SUMMARY					
Number of neurodevelopmental domains with evidence of severe impairment:					
			· (_	

DIAGNOSIS:

For derivation of the Australian FASD diagnostic categories, please refer to the Australian FASD Diagnostic Criteria and FASD Diagnostic Pathway Algorithm below (also see Table 1 and Figure 1 in the Guide). Record the diagnosis below. *Indicate as applicable:*

	FASD with 3 sentinel facial features
	FASD with < 3 sentinel facial features
	At risk of FASD
	Incomplete assessment e.g. further investigation/information needed
	Other diagnoses (with or without FASD)
	Other diagnoses (with or without 1735)
Clin	ical notes:

PATIENT DETAILS

	NAME						
	Sex	☐ Female ☐ Ma		☐ Male	☐ Other		
Date of birth (DD/MN	1/YYYY)	/ / Age at ass		e at assessment:			
Racial/ethnic back	ground						
Hospital number (if ap	plicable)						
ALCOHOL EVROCURE CHIMANARY							
ALCOHOL EXPOSURE SUMMARY Source of reported information on alcohol use: Birth mother Other (specify)							
In your judgement what is the reliability of the information on alcohol exposure: \Box Unknown \Box Low \Box High							
In your judgement was there high-risk	consump	tion of a	alcoho	ol during pregi	nancy? 🗆 Unknow	n □ Yes □ No	
Prenatal alcohol exposure: ☐ Unknown	n exposu	re 🗆 No	о ехро	osure 🗆 Confi	rmed exposure 🗆 Co	onfirmed-high risk exposure	
SENTINEL FACIAL FEATURES SUMMARY							
Number of Sentinel Facial Features (PF	L 2 SD or	more b	elow	the mean, phi	ltrum rank 4 or 5, upp	er lip rank 4 or 5):	
□ 0	□ 1	□ 2		□ 3			
NEURODEVELOPMENTAL DOMAINS SUN	ИMARY						
Neurodevelopmental Domain		Impairment					
1 Brain structure/Neurology	□ No			☐ Yes	☐ Not asse	essed	
2 Motor Skills	□ No	ne		☐ Some	☐ Severe	☐ Not assessed	
3 Cognition	□ No	ne		☐ Some	☐ Severe	☐ Not assessed	
4 Language	□ No	ne		☐ Some	☐ Severe	☐ Not assessed	
5 Academic achievement	□ No	ne		☐ Some	☐ Severe	☐ Not assessed	
6 Memory impairment	□ No	ne		☐ Some	☐ Severe	☐ Not assessed	
7 Attention	□ No	ne		☐ Some	☐ Severe	☐ Not assessed	
8 Executive function, including impulse control and hyperactivity	□ No	ne		☐ Some	☐ Severe	☐ Not assessed	
9 Affect regulation	□ No	ne		☐ Some	☐ Severe	☐ Not assessed	
10 Adaptive behavior, Social Skills, or Social Communication	□ No	ne		□ Some	☐ Severe	☐ Not assessed	
Number of neurodevelopmental doma				evere impairm e (specify)	ent 		
Other Prenatal or Post-natal risk/exposu	ure						
	lo knowr	ı risk	□ u	Inknown risk	☐ Some risk	☐ High risk	
Postnatal risk summary:	No know	n risk		Inknown risk	☐ Some risk	☐ High risk	
Growth summary							
Was an unexplained deficit in height or	weight ·	< 3 rd per	centil	e identified at	any time?	s □ No	

DIAGNOSIS:

For derivation of the Australian FASD diagnostic categories, p	please refer to the Australian FASD Criteria and the FASD Diagnost
Pathway Algorithm below (also see Table 1 and Figure 1 in th	he Guide). Record the diagnosis below.

Indicate as applicable:
 □ FASD with 3 sentinel facial features □ FASD with < 3 sentinel facial features □ At risk of FASD □ Incomplete assessment e.g. further investigation/information needed □ Other diagnoses (with or without FASD)
Clinical notes: