Severe obesity and global developmental delay in preschool children

CANADIAN PAEDIATRIC SURVEILLANCE PROGRAM

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REPORTING INFORMATION

To be completed by the CPSP)	
Report number:	
Month of reporting:	
Province:	
Foday's date:	

Please complete the following sections for the case identified above. If the information asked below is not readily available, please leave it blank. Strict confidentiality of information will be assured.

	CASE DEFINITION FOR SEVERE OBESITY AND GLOBAL DEVELOPMENTAL DELAY IN PRESCHOOL CHILDREN									
	Report any new case of a child	≤5 years of age with:								
	 Severe obesity (SO), defined as body mass index ≥99.9th percentile according to references developed by the World Health Organization and the Canadian Pediatric Endocrine Group. The absolute cut-offs by age and sex are shown below in Appendix 1. 									
	AND									
	 Global developmental dela Gross motor Fine motor Speech/language Cognitive Social/personal Delay in activities of da 	y (GDD), defined as a significa ily living	nt delay in two or more develo	opmental domains, including:						
			Month first seen (mr	n):						
SEC 1.1 1.3 1.4	TION 1 – DEMOGRAPHIC Date of birth: / / DD / MM Postal code (first 3 digits o Population groups (check	/ YYYY nly):	1.2 Sex: Male Fe	male						
	□ Arab	□ Black	□ Chinese	🗆 Filipino						
	Japanese	Korean	Latin American	□ White						
	□ First Nations	🗆 Inuit	□ Métis	Unknown						
	☐ Southeast Asian (e.g., Vietnamese, Cambodian, Laotian)	 ☐ South Asian (e.g., East Indian, Pakistani, Sri Lankan) 	□ West Asian (e.g., Iranian, Afghan)	☐ Other, specify:						
SEC	TION 2 - PATIENT INFOR	MATION								
2.1 2.2	Please attach de-identified Patient's most recent grov Height: cm OR Length: cm OR	inches	Date performed: DD	_// MM YYYY						

Weight: _____ kg OR _____ lbs

Head circumference: _____ cm OR _____ inches

2.3 Current physical features: Blood pressure: ____/ Waist circumference: _____ cm OR _____ inches

2.4 Global developmental delay is defined as a significant delay in ≥2 developmental domains.

(Check all that apply):	Yes	No	Unknown
Gross motor delay			
Fine motor delay			
Speech/language delay			
Cognitive delay			
Social/personal delay			
Delay with activities of daily living			

SECTION 3 – FAMILY HISTORY

 $\hfill\square$ Not obtained in the course of my care of this case

3.1 Have members of your patient's **biological family** been diagnosed with (check all that apply):

Diagnosed	None	Mother	Father	Sibling	Unknown
ADHD					
Alcohol and/or drug problem					
Anxiety and/or depression					
Asthma					
Autism spectrum disorder					
Diabetes					
Global developmental delay					
Heart disease and/or stroke					
High cholesterol					
Hypertension					
Intellectual and/or learning disability					
Obesity					

SECTION 4 – PREGNANCY/BIRTH INFORMATION Not obtained in the course of my care of this case

4.1	Age of mother at pregnancy: years			
4.2	Weight prior to pregnancy: kg Weight prior to delivery:	kg	Unknown_	
4.3	Prescribed medications in pregnancy: Yes No If known, list:			
4.4	Pregnancy/postpartum history (check all that apply):	Yes	No	Unknown
	Tobacco use			
	Alcohol use			
	Illicit drug use			
	Gestational diabetes			
	Hypertension			
	Postpartum depression			
4.5	Birth/postpartum history:			
	Gestational age: weeks Birth weight: kg			
	(Check all that apply):	Yes	No	Unknown
	Complicated birth (If yes, please circle: breech, failure to progress, PROM))		
	Required neonatal intensive care unit care			
	Exclusively breastfed			
	If yes, length of time: months			
	Total time breast fed (exclusive and non-inclusive): months			
	Formula fed			
	If yes, age of introduction: months			

		Yes	No	Unknown
	Combination breast and formula fed			
	If yes, age of introduction of formula months			
4.6	Solid foods – age of introduction: months			
SECT	TION 5 – CLINICAL PRESENTATION	ourse o	f my ca	ire of this case
5.1	Since what age have you been following this patient?		_ years	months
5.2	At what age was the diagnosis of global developmental delay made?		_ years	months
5.3	At what age did you first have concerns about weight?		_ years	months
5.4	Please describe your role in this patient's care.			
	Primary care paediatrician Consulting paedia	atrician		
	Consulting developmental paediatrician Consulting geneti	cist		
	Consulting endocrinologist Other, specify:			

SECTION 6 - CLINICAL FINDINGS AND INVESTIGATIONS

6.1 Health problems (check all that apply) – current or past:

	Yes	No	Unknown
ADHD			
Asthma or recurrent wheeze			
Blount's disease (or bowing of legs)			
Bullying			
Depression and/or anxiety			
Diabetes			
Diagnosed sleep apnea			
Fatty liver disease			
Fractures			
Gastroesophageal reflux			
Hearing loss			
High blood pressure			
Hyperlipidemia			
Hypothyroidism			
Insulin resistance and/or acanthosis nigricans			
Nutrient deficiencies			<i>If yes</i> , specify:
Recurrent otitis media			
School and/or behavioural problems			
Seizures			
Snoring			
Any other health problems			<i>If yes</i> , specify:

6.2 Current medications and non-prescription products:

	Yes	No	Unknown	If yes, please list: (drug, dose, frequency)
Medications				
Non-prescription medications				
Natural health products				
Vitamin or mineral supplements				
Other				

				SO and	GDD in preschool childre	en questionnaire — Page 4
6.3	Have genetic tests been ordered?	Yes	No	Pending	_ Unknown	
	(See Appendix 2 if you would like	more info	ormation	on how to obtain	microarray testing.)	
	If yes:					
	a) Has a microarray been ordered	l? Yes_	No	Pending	Unknown	
	b) Please record the result of gen		-		•	Unknown
	<i>If abnormal</i> , what was th	e result o	of the sp	ecific genetic test	(see Appendix 3)?	
					(OR attach de	-identified results)
6.4	Has the patient been diagnosed w			ental disorders, ir nknown	cluding:	
	Autism spectrum disorder	163 1	10 0	IIKIIOWII		
	Cerebral palsy					
	Post-encephalitis/meningitis					
	Other (please list):					
6.5	Has central nervous system imagi	ng been d	complete	d? Yes I	No Unknown	
	If yes, provide results (check all the	at apply):				
	CT head Results:					
	MRI brain Results:	·				
	Ultrasound head Results:					
6.6	Is the patient currently enrolled in a	any obesi	ty mana	gement program?	Yes No	_ Unknown
6.7	Which of the following other clinicia	ans/servi			the patient's care?	
	(check all that apply):	Yes	No	Unknown		
	Autism assessment					
	Autism intervention					
	Child development program					
	Clinical geneticist					
	Developmental paediatrician					
	Dietitian					
	Family physician					
	General paediatrician					
	Home visiting program					
	Neurologist					
	Obesity program					
	Occupational therapy					
	Paediatric endocrinologist					
	Physiotherapist					
	Psychology or psychiatry					
	Public health nurse					
	Social worker					
	Speech therapy					
	opecent merapy					

6.8 What has been the biggest challenge in providing care for this patient?

____ I agree to be contacted by the CPSP for further information on this questionnaire.

____ I do not wish to be contacted by the CPSP for further information on this questionnaire.

SECTION 7 – REPORTING PHYSICIAN

First name	Surname	
Address		
City	Province	Postal code
Telephone number	Fax number	
E-mail	Date completed	

Thank you for completing this form.

(SOGDD 2018/02)

AGE (years)	BOYS BMI ≥99.9 th percentile	GIRLS BMI ≥99.9 th percentile
6 months	22.3	22.3
1	21.6	21.6
2	20.6	20.6
3	20.0	20.3
4	19.9	20.6
5	20.3	21.1

Appendix 1: Age- and sex-specific cut-offs for severe obesity using World Health Organization Growth Standards*

*de Onis M, Garza C, Victora CG, Onyango AW, Frongillo EA, Martines J. The WHO Multicentre Growth Reference Study: planning, study design, and methodology. Food and nutrition bulletin. 2004;25(1 Suppl):S15–26.

Appendix 2: Microarray Testing

For more information on how to obtain microarray testing, please contact the Severe obesity and global developmental delay in preschool children study coordinator at 780-394-0066.

Appendix 3: Potential Results of Abnormal Genetic Testing

Melanocortin pathway defects	Oligogenic genomic disorders, including disorders of imprinting	Obesity with retinitis pigmentosa
Leptin deficiency	Prader-Willi syndrome	Bardet-Biedl syndrome
Leptin receptor deficiency	Beckwith-Wiedemann syndrome	Alström syndrome
Pro-opiomelanocortin deficiency	Wilms tumor, aniridia, genitourinary anomalies, mental retardation, and obesity syndrome	Cohen syndrome
Prohormone convertase 1/3 deficiency	Albright hereditary osteodystrophy	Monogenic obesity disorders not otherwise classified
Melanocortin 4 receptor deficiency	Fragile X syndrome with Prader-Willi phenotype	SIM1 deficiency
Brain-derived neurotrophic factor deficiency	Maternal uniparental disomy chromosome 14	Oligogenic disorder, not otherwise classified
Neurotrophic tyrosine kinase, receptor type 2 deficiency	Chromosomal microdeletion 16p11.2	Chromosomal aneuploidies
Other monogenic syndromes of melanocortin pathway, specified	Carpenter syndrome	Other aneuploidy, specified
Other	Other	