Congenital Zika syndrome (CZS) in infants in Canada

CANADIAN PAEDIATRIC SURVEILLANCE PROGRAM

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REPORTING INFORMATION (To be completed by CPSP staff) Report number: Month of reporting: Province: Today's date:

Please complete the following sections for the case identified above.

Confidentiality of information will be assured.

CASE DEFINITION FOR CONGENITAL ZIKA SYNDROME

Report any infant less than 12 months of age who presents with:

 Microcephaly defined as head circumference less than 2 standard deviations for gestational age and sex according to the standardized reference percentile*

OR

Other congenital anomalies and malformations consistent with congenital Zika syndrome including malformations of the
central nervous system, such as intracranial calcifications, structural brain or eye abnormalities, or other congenital central
nervous system-related abnormalities (not explained by another etiology[†])

AND

 A maternal history that includes an epidemiologic linkage[‡] to Zika virus OR a positive or inconclusive Zika virus laboratory test

OR

- An infant with a positive or inconclusive Zika virus laboratory test
- If there is a case of severe microcephaly suspected to be associated with Zika virus then a questionnaire for the severe microcephaly study <u>and</u> the congenital Zika syndrome study should be completed (i.e., if the case meets both case definitions).
- † Other etiologies that should be considered include other congenital infections such as syphilis, toxoplasmosis, rubella, cytomegalovirus, varicella zoster, parvovirus B19, and herpes simplex virus. An assessment of potential genetic and other teratogenic causes of the congenital anomalies should also be considered.
- ‡ Epidemiological linkage means: travelled to, or resided in, an area with active Zika virus transmission during her pregnancy; OR had unprotected sex during pregnancy with a partner who resided in, or traveled to, an area with active Zika virus transmission.

CASE INCLUSION CRITERIA

There must be at least one check in Column 1 AND one check in Column 2 to meet the case definition.

Zika virus criteria	Column 1 Yes	Clinical criteria (infant up to 12 months of age)	Column 2 Yes
Born to mother with positive Zika virus test		Has microcephaly (according to definition above)	
Born to mother with inconclusive Zika virus test (ZIKA plaque reduction neutralization test with inconclusive result)		(according to definition above)	
Born to mother with <u>epidemiological link</u> with <u>no</u> <u>Zika virus test or unknown test result</u>		Has any congenital anomaly or malformation of concern	
Infant has positive Zika virus test			

SECTION 1 - DEMOGRAPHIC AND BACKGROUND CLINICAL INFORMATION

1.1	Date of first vis	:/	/	/	
			DD	MM	YYYY
1.2	Date of birth:	//	_		
		DD MM YYYY			

.5 E			juous geni	talia					
	Province/territory of residen	nce:							
	thnicity (check all that app	oly): First N	lations	Inuit	Métis	White	South Asi	an (e.g., East	Indian
P	Pakistani, Sri Lankan, etc.)	• •						. •	
	Southeast Asian (e.g., Vietr				•				tc.)
	Korean Japanese							_	,
	Node of delivery:			_					
	Spontaneous vaginal delive	erv	Assisted v	aginal deliver	v A	Assisted breech	/breech ext	raction	
	Caesarian section	•							
	APGAR scores: 1 minute			•					
	Gestational age at delivery:			·	o minato.				
	Birth type: Singleton			rder multinle	Or	der of hirth	Unknov	vn	
			r lighter of g):			rence (cm):	_	VII	
	• , ,			At birt		` '			
_									
А	t visit:	At visit:		At visi	t:				
ECTIO:	N 2 – PREGNANCY AND	FYPOSIII	RES INFO	RMATION					
			(LO IIII O			_	_		
1 M	laternal disorder/disease		Present		Ag		of diagnosis DD/YYYY)		
ļ	Hypertension		Yes	NoUnk	nown	Age:		///_	
	Gestational diabetes melli	tus	Yes	No Unk		_			
	Pre-gestational diabetes		Yes	No Unk		•		//_	
	Sickle cell disease		Yes	No Unk				//_	
	Elevated maternal PKU		Yes	NoUnk				//_	
	Severe malnutrition		Yes	No Unk		·		//_	
ſ	Placental insufficiency		Yes	No Unk	known	Age:	or _	//_	
ſ	Rhesus disease		Yes	No Unk	known	Age:	or _	//_	
i	Abnormal ultrasound findi	ngs	Yes	No Unl	known	Age:	or _	//_	
(Other pregnancy complica	ations	Yes	No Unk	known	Age:	or _	//_	
(Chronic conditions of the r	mother:	Yes	No Unk	known	Age:	or _	//	
I	If yes, please specify cond	dition(s): _							
_									
	Exposure history during		•			_			
	Smoking			Unknowr		f yes, consump	•	•	
	Alcohol use			Unknow		f yes, consump		•	
	Illicit drug use	Yes	No_	Unknowr	า [Drug:	Consur	notion:	
	Known teratogen exposure			Unknowr		Specify:			

	during or immediate					
	-		-	-		
Symptoms: Fev	er Rash	_ Arthralgia/arth	ritis Conju	nctivitis	Myalgia	_ Headache
•		•	-			
	have a sexual part		to other countrie	es during preg	nancy or thr	ee months prior to
•	es No Un	known				
If yes, please s	specify:					
Country			proximate date a	arrived Ap	proximate of DD/MM/YY	date departed
			_ //		/ /_	
					//_	
Sexual partner	illnoss prosont duri	ng or immodiatel	_ / /			 lo Unknown
•	•	•		•		OIRHOWH
	=		_			Headache
		_				rieauacrie
rtotro orbitar pa						
Maternal pre-n Tests	atal tests (If no te	sts done, check ormed	NO as appropr Date of t	-	Res	ult
16212	Perio	ormea	(DD/MM/YY		Kes	uit
RT-PCR Zika:	V N-	I la la succe	, ,			
BloodUrine	Yes No Yes No		///			egative Unknown
Placenta	Yes No			——— Ро ——— Ро		egative Unknown egative Unknown
Zika serology	Yes No	Unknown	////////	lgl	M: Positive	_ Negative Unknowr
		Unknown	///			_ Negative Unknown
Zika PRNT	Yes No	Unknown	///	Po	sitive N	egative Unknown
(plaque					conclusive*_	-
reduction neutralization						rentiate between Zika,
test)					J	er flaviviruses
Dengue	Yes No	Unknown	//	lgl	M: Positive	_ Negative Unknowr
serology	Yes No	Unknown	////////	lg(G: Positive	_ Negative Unknown
Congenital	Yes No	Unknown	/ /	Po	sitive N	egative Unknown
infection screen					oositive, spec	
						bella CMV
						V Syphilis pecify:
Matamalaat						
Maternal post-	natai tests (if tes	is not done, che	eck nere ai	na go to 4.3 (Shiid Zika te	ests section below.)
•			after delivery? V	es No		
•	above tests done	and/or repeated	and delivery: 1		_	
•		and/or repeated	Date of test		 esult	
Were any of the		and/or repeated	•	Re	esult	Unknown
Were any of the		and/or repeated	Date of test	Re	esult Negative	

4.3 Child Zika virus tests

	Tests Performed		ed	Date of test (DD/MM/YYYY)			Result						
	RT-PCR ZIKA: • Blood • Urine • CSF Zika serology Zika PRNT	Yes N Yes N Yes N Yes N	No U No U No U No U	nknown_ nknown_ nknown_ nknown_ nknown_ nknown_		/ //	/ / /	Positive I Positive I Positive_ I IgM: Positive_ IgG: Positive_ Positive_ I Inconclusive*_ * unable to diff dengue	Negative Negative Negativ Negativ	_ Unkno _ Unkno /e Ur /e Un _ Unkno	own nknown own		
4.4	Child congenita												
			•			-	he following:	nilis VZV	⊔ 1\7				
	•	ease specify					• •	IIII5 VZV	IIIV_				
	• •							ital infection(s) If the ch	ild tests ı	nositive for		
								refer to whic			00011110 101		
				Perfo	ormed		Normal	Abnorr	nal	Da	ate of test:		
								(give re	(give result)		(DD/MM/YYYY)		
	PCR		Yes	_ No	Unknov	wn				/_	/		
	IgM		Yes	_ No	Unknov	wn				/_	/_		
	IgG		Yes	_ No	Unknov	wn				/_	/		
	Other:		Yes	_ No	Unknov	wn				/_	/		
	Other bas	sis for											
SECT 5.1	diagnosis ION 5 – CLINICAI Cranial morpho	L INFORMA	TION			Presen	t	At k	oirth <i>or</i> da				
	Microcephaly (>	2 SD and < 3	SD for gas	tational				Δt hirth	n or	(DD/MM/Y	,		
	age and sex accord percentile)				Yes	. No	_ Unknown		i <i>0i</i>	/	/		
	Severe microcep and sex according percentile)				Yes	No	Unknown_	At birth	n or	/	/		
	Overlapping crai	nial sutures			Yes	No	Unknown_	At birth	n or _	/	/		
	Overlapping crain Prominent occip				Yes Yes			 ^			/ /		
	0	ital bone					Unknown_	 ^	n or		/ / /		
	Prominent occip Excess nuchal s	ital bone kin			Yes	No	Unknown_ Unknown_	At birth	n or n or	/	/		
	Prominent occip	ital bone kin proportion			Yes	No	Unknown_ Unknown_ Unknown_	At birth At birth At birth	n or n or	/_ /	//////		
5.2	Prominent occip Excess nuchal s Craniofacial disp	ital bone kin proportion ssion			Yes Yes	No No No	Unknown_ Unknown_ Unknown_ Unknown_	At birth At birth At birth At birth	or or n or n or n or	/////	///////		
5.2	Prominent occip Excess nuchal s Craniofacial disp Biparietal depres	ital bone kin proportion ssion			Yes Yes	No No No Present	Unknown_ Unknown_ Unknown_ Unknown_	At birth At birth At birth At birth At birth	or or n or n or n or	/	///////		
5.2	Prominent occip Excess nuchal s Craniofacial disp Biparietal depres Ocular anomalia	ital bone kin proportion ssion			Yes Yes Yes	No No No Present	Unknown_ Unknown_ Unknown_ Unknown_	At birth At birth At birth At birth At birth At birth	or or or or or or oirth or da	////	///////		
5.2	Prominent occip Excess nuchal s Craniofacial disp Biparietal depres Ocular anomalia Microphthalmia	ital bone kin proportion ssion			Yes Yes Yes Yes	No No No Present	Unknown_ Unknown_ Unknown_ Unknown_ Unknown_ Unknown_	At birth	or	/// te first ic (DD/MM/Y	///////		
5.2	Prominent occip Excess nuchal s Craniofacial disp Biparietal depres Ocular anomalia Microphthalmia Coloboma	ital bone kin proportion ssion es			Yes Yes Yes Yes Yes	No No No Present No No No	Unknown_ Unknown_ Unknown_ Unknown_ Unknown_ Unknown_ Unknown_ Unknown_	At birth	or or or or or or or da or or or or	te first ic (DD/MM/Y	///////		

	Focal pigmentary mottling of the retina	Yes	No	Unknown	At birth	_ or	/	/
	Optic nerve atrophy/anomalies	Yes	No	Unknown	At birth	_ or	/	/
	Retinal lesions, including well-defined chorioretinal atrophy and gross pigmentation	Yes	No	Unknown	At birth	_ or	/	/
5.3	Congenital contractures		Present		At bir	th <i>or</i> da		identified I/YYYY)
	Arthrogryposis	Yes	No	Unknown	At birth	_ or	•	/
	Clubfootunilateralbilateral	Yes	No	Unknown	At birth	_ or	/_	/
	Bilateral congenital hip dislocation	Yes	No	Unknown	At birth	_ or	/	/
	Dislocation or partial dislocation of one or both knees	Yes	No	Unknown	At birth	_ or	/	/
5.4	Neurological sequelae		Present		At bir	th <i>or</i> da		identified I/YYYY)
	Motor disabilitiesfinegross	Yes	No	Unknown	At birth	_ or	•	/
	Hypertonia and spasticity	Yes	No	Unknown	At birth	_ or	/_	/
	Hypotonia	Yes	No	Unknown	At birth	_ or	/	/
	Tremors	Yes	No	Unknown	At birth	_ or	/	/
	Abnormal posturing (e.g., opisthotonus)	Yes	No	Unknown	At birth	_ or	/_	/
	Cognitive disabilities, specify:	Yes	. No	Unknown	At birth	_ or	/_	/
	Irritability/excessive crying	Yes	No	Unknown	At birth	_ or	/	/
	Seizuresgeneralfocal	Yes	No	Unknown	At birth	_ or	/	/
	Swallowing difficulties	Yes	No	Unknown	At birth	_ or	/_	/
	Failure to thrive	Yes	No	Unknown	At birth	_ or	/	/
	Vision impairment, specify:	Yes	No	Unknown	At birth	_ or	/_	/
	Hearing impairment/loss, specify:	Yes	No	Unknown	At birth	_ or	/	/
5.5	Brain imaging/neurophysiology		Present		At bir	th <i>or</i> da		identified I/YYYY)
	Intracranial calcifications	Yes	No	Unknown	At birth	_ or	/	•
	Neural tube defect	Yes	No	Unknown	At birth	_ or	/_	/
	Congenital hydrocephalus	Yes	No	Unknown	At birth	_ or	/	/
	Increased fluid spaces (ventricular and extraaxial)	Yes	. No	Unknown	At birth	_ or	/	/
	Cortical thinning with abnormal gyral patterns (most consistent with polymicrogyria)	Yes	No	Unknown	At birth	_ or	/	/
	Hypoplasia or absence of the corpus callosum	Yes	No	Unknown	At birth	_ or	/	/
	Decreased myelination	Yes	No	Unknown	At birth	_ or	/	/
	Cerebellar or cerebella vermis hypoplasia	Yes	No	Unknown	At birth	_ or	/_	/
	Thinning of the cord and reduction in the ventral roots (via MRI)	Yes	No	Unknown	At birth	_ or	/	/
	Wallerian degeneration of the long descending tracts	Yes	No	Unknown	At birth	_ or	/	/
	Abnormal EEG	Yes	No	Unknown	At birth	_ or	/	/

5.6	Has the patient achieved age? (Please complete				wing dor	mains, approp	oriate for age/o	corrected gestational	
	Gross motor	Yes	No Unkn	own	N/A. sc	pecify			
	Fine motor		No Unkn						
	Social		No Unkn						
	Language	Yes	No Unkn	own					
5.7	Other health systems affe	ected:							
	Cardiovascular system		_ Abnormal	Unknow	n	If abnormal,	specify:		
	Respiratory system								
	Gastrointestinal system								
		Abdominal Splenome	tenderness galy Omph	_ Jaundic	e He Gast	epatomegaly_ troschesis	Hernia Other, spec	sify:	_
5.8	Other abnormalities: Ye								_
SECT	TION 6 – FAMILY HISTORY	Y							
6.1	Consanguinity of parents	: Degree of	relationship: No	one 1 ^s	st 2 nd	d 3 rd	>3 rd		
6.2	Family history (up to 2 nd c	degree relativ	ves, e.g., grand	parent, sib	ling, etc.):			
	Microcephaly	-	Yes No_	Unkno	wn	If yes, speci	ify relative:		
	Other congenital anomali	ies,	Yes No	Unkno	wn	If yes, speci	ify relative:		
	specify					, , - ,			
	History of developmental specify	delays,	Yes No_	Unkno	own	If yes, speci	ify relative:		
	Previous pregnancy with anomaly(ies)		Yes No_	Unkno	own	If yes, speci	ify relative:		
	History of miscarriage		Yes No_	Unkno	wn	If yes, spec	ify relative:		
	History of stillbirth		Yes No_	Unkno	wn	If yes, speci	ify relative:		
			d by the CPSP ntacted by the				•		
POSSI	BLE COHORT STUDY OP	PORTUNITY	′						
consen	rate study may be conducted to f the family. This process unity? By checking "yes" yous) to the research team res	s is separate u are giving	from the CPSF permission for t	P. Are you the CPSP	intereste	d in obtaining	further inform	nation on this	

Yes___ No___

SECTION 7 – REPORTING PHYSICIAN

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

Thank you for completing this form.

(CZS 2017/03)