

CONFIDENTIAL WHEN COMPLETED VCH: Fax the completed report form to BCCDC (604) 707-2516 c/o CDPACS-Zika All other Health Authorities: Enter information into Panorama Case definitions are in Section L below					
PERSON REPORTING					
Health Authority: ☐ FHA ☐ FNHA	□ IHA	□ NHA	□ VCH	□ VIHA	
Name: Last First	Phone Nu	mber: ()	-	ext.	
Email:	Fax Numb	oer ()	-	ext.	
	Date case	report form com	pleted:	YYYY/MM/DD	
Date report received by health authority:					Record in:
· · · · · · · · · · · · · · · · · · ·	YYYY/MM	/DD	_		>Investigation >Investigation Details >>>Reporting Notifications as Report Date (Received)
A. PERSONAL INFORMATION					
Name:	First		Middle		Record or review and update in >Subject
Date of Birth: Ger	nder: Male	☐ Female ☐	Undifferentiated	☐ Unknown	>>Client Details >>>Personal
Health Card Number:		Alternate Name	e(s):		Information Select this address
Phone Number (home/work/mobile): ()		-	ext.		as "Client Home Address at Time of
Address: Unit # Street #	Str	reet Name		City	Initial Investigation" in >Investigation
Postal Code: Province:		Country of Resid	dence (if not Canada):		>>Investigation Details >>>Investigation Information
B. TRAVEL DURING EXPOSURE PERI	IOD				ea.e
Has the case returned from a country with ongo					
widespread Zika transmission in the previous 2		∕es □ No	☐ Unknown		
Origin (city, prov/state, country):			Departure Date:		5
Destination (city, prov/state, country):			Return Date:	YYYY/MM/DD	Record in >Investigation >>Investigation
Did the case notice any insect bites during t	travel?	∕es □ No	 □ Unknown	YYYY/MM/DD	Details >>>Links &
Did the case seek travel medical services p departure?	orior to	∕es □ No	☐ Unknown		Attachments >>>Zika virus case investigation form
List all cities/countries visited:					
Travel details (e.g. activities, mode of travel):					

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C. TRANSMIS	SSION RISK FACTORS						
		Yes	No	Asked but Unknown	Declined to Answer	Not Assessed	
Female cases							Record in >Investigation
	egnant? risk factor and client warning for post-natal fo IC Women's hospital	llow					>>Investigation >>Investigation Details >>>Links & Attachments >>>Zika virus case
	Expected Due Date:						investigation form NOTE: Expected Due
Male cases	Male cases		MM/DD				Date related to female cases or male cases
who is pregna		Ш					with a pregnant sexual partner can be captured
risk factor and clie	contact record for the pregnant woman, add a nt warning for post-natal follow up; refer case ner to BC Women's hospital						in Pregnancy Details If the case or contact is pregnant, see Section
	Expected Due Da	ate:					M.1 for data standards to create a risk factor in
		YYYY/N	YYYY/MM/DD				
	Pregnant contact informati	ion: Name:					>>Risk Factors and a client warning in >Subject >>Client Warnings
				.ast	First		
		Date of birth:		PHN:			If the contact is pregnant, record in
			YYYY/MM/D	D			>Investigation >>Exposure Summary
	of sexual transmission with cas guidance, see Section M.2	e **			as a Transmission Event (Classification = Contact – person under investigation)		
		Yes	No	Asked but Unknown	Declined to Answer	Not Assessed	
All cases							If case has donated
Has the case donated blood, tissue or organ(s) in Canada in the previous 3 weeks?		in \square					blood, tissue or organs(s), record in >Investigation
Date(s)	of donation:						>>Exposure Summary as a Transmission
		YYYY/N	/IM/DD				Event (see Section M.3)
Facility	where donation occurred:						(555 555)
Other risk factor, specify:							
D. OTHER EX	POSURES						
	th a male partner who has returne pread Zika transmission in the pre		h □ Yes	□ No	□ Unknown		Record in >Investigation >Investigation Details >>Links & Attachments
If yes, Contact name Date		Date of last known co	ontact	Other details			>>>Zika virus case investigation form
							NOTE: Contact tracing beyond pregnant contacts is not required for surveillance; if a contact is identified as a case, link the records through a
							TE/AE OR update the classification
							of contacts with an existing investigation >Investigation >sinvestigation Details >>Disease Summary
	eived blood, tissue or organ(s) in 0 ing Zika transmission in the previous		☐ Yes	□ No	☐ Unknown		If case has received blood, tissue or
Date(s) of rec						organs(s), record in >Investigation >>Exposure Summary	
					as an Acquisition Event (see Section M.3)		



E. LABORATO	ORY INFORMA	TION					
Specimen Collected	Collection Date (YYYY/MM/DD)	Test Performed			Result		
		□ PCR	☐ Positive	☐ Negative	☐ Indeterminate	☐ Pending	
		Lab Report Date:	Y/MM/DD	Reporting Lab:			
		☐ Serology IgM	☐ Positive	☐ Negative	☐ Indeterminate	☐ Pending	
□ Blood		IgG	☐ Positive	☐ Negative	☐ Indeterminate	☐ Pending	
		Lab Report Date:	Y/MM/DD	Reporting Lab:			
		☐ Neutralizing assay	☐ Positive	☐ Negative	☐ Indeterminate	☐ Pending	Receive through Electronic-Lab inbox, or
		Lab Report Date:	Y/MM/DD	Reporting Lab:			manually record in >Investigation >>Lab
		□ PCR	☐ Positive	☐ Negative	☐ Indeterminate	☐ Pending	>>>Lab Quick Entry NOTE: In Result Name
☐ Urine		Lab Report Date:	Y/MM/DD	Reporting Lab:			type 'Zlka' to get a list of Zlka results
		□ PCR	☐ Positive	☐ Negative	☐ Indeterminate	☐ Pending	Record Causative Agent in >Investigation
Nasopharyngeal swab		Lab Report Date:	Y/MM/DD	Reporting Lab:			>>Investigation Details >>>Disease Summary
		□ PCR		☐ Negative	☐ Indeterminate	☐ Pending	
		Lab Report Date:	Y/MM/DD	Reporting Lab:			
		☐ Serology IgM	☐ Positive	☐ Negative	☐ Indeterminate	☐ Pending	
☐ Other		IgG	☐ Positive	☐ Negative	☐ Indeterminate	☐ Pending	
Specify:		Lab Report Date:	Y/MM/DD	Reporting Lab:			
		☐ Neutralizing assay	☐ Positive	☐ Negative	☐ Indeterminate	☐ Pending	
		Lab Report Date:	Y/MM/DD	Reporting Lab:			
F. PHYSICIAN	I						
Physician Name:		Last		First			Record in >Investigation >>Investigation Details
Physician Phone:	()	-	1 1131	ext.		>>>External Sources If ordering physician is
(For pregnant cas	es or cases with p	pregnant contacts)					NOT providing perinatal care, record contact information for the
Physician Name:		l and		Final			professional providing perinatal care (e.g. physician, midwife) in
Physician Phone:	(Last	-	First	ext		>Subject >Client Details >>Health Services

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G. CLINICAL PRESENTATION AND COMPLIC	CATIONS					
Onset date if the case has (or had) symptoms compatible from last possible exposure (e.g. travel, sexual):	le with Zika virus	infection 2 v	veeks			
YYYY/MM/DD						
The case is asymptomatic at time of reporting and has not experienced symptoms compatible with Zika virus infection in the 2 weeks from last possible exposure (e.g. travel, sexual)						
Sign / Symptom	Yes	No	Asked but Unknown	Declined to Answer	Not Assessed	
Arthralgia (painful joints)						
Conjunctivitis						Record in
Diarrhea						>Investigation >>Signs and
Fever						Symptoms
Paralysis, flaccid						Select "Set as Onset" and record onset date of earliest symptom
Guillain-Barré Syndrome (GBS)*						For asymptomatic
Headache						cases, set 'Present' field to 'No'
Malaise						
Myalgia (muscle pain)						
Rash						
Other, specify:						
*Check only if formally assessed and diagnosed by a medical professiona	l					
H. RELEVANT IMMUNIZATIONS						
						>Investigation >>Investigation Details
Has the case received a Yellow Fever vaccine in the pa	ıst? □ Yes	□ No	☐ Unknown			>>>Links & Attachments >>>>Zika virus case investigation form
						AND Record details of
If y	ves, Date(s) of i	mmunizatio	า:			relevant immunizations (e.g. Yellow Fever) in
			YYYY/I	MM / DD		Immunization module (see Section M.4)
I. OUTCOME AT TIME OF REPORTING						
☐ Fully Recovered ☐ Not yet recovered/reco	vering \Box Fa	ital <i>If died</i>	date of death:			
			- 1- 11'6	YYYY/MI	M/DD	Record in >Investigation
☐ Other, specify below ☐ Unknown	⊔ P€	ermanent dis	ability, specify b	eiow		>> Outcome (see Section M.5)
Specify other outcome / permanent disability:						
J. CLASSIFICATION						
☐ Confirmed ☐ Person u	nder investigation	(to be used for	case management pu	ırposes in Panorama	if desired)	Record/Update in >Investigation
See Section L for case definitions.						>>Disease Summary
K. GENERAL COMMENTS						
						>Investigation >>Investigation Details
						>>>Links & Attachments
						>>>>Zika virus case investigation form

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L. CASE DEFINITIONS		
Zika Virus Disease		Reportable?
Confirmed case	Laboratory confirmation of infection:	Yes
	Detection of Zika virus-specific nucleic acid by reverse-transcriptase PCR from an appropriate clinical specimen (e.g. blood, urine)	
	OR	
	Demonstration of specific IgM antibodies in an appropriate clinical specimen (e.g. blood) by enzyme-immuno assay (e.g. ELISA) AND confirmation through identification of Zika virus-specific neutralizing antibodies (e.g. using PRNT)	
Person under investigation	A person with two or more symptoms compatible with clinical illness with onset during or within 2 weeks of travel to a country with ongoing or widespread transmission1	No
	OR	
	A person who is epidemiologically-linked to a confirmed case or a person under investigation	
	OR	
	A female who was pregnant during or within two months of returning from a country with ongoing or widespread Zika virus transmission	
	OR	
	A male returning from a country with ongoing or widespread Zika virus transmission AND has a female partner who is pregnant, becomes pregnant within 2 months of his return, or intends to become pregnant in the following 2 months	
	OR	
	A person with specific IgM antibodies from an appropriate clinical specimen with pending or inconclusive confirmatory testing (e.g. PRNT)	

Notes:

¹An updated list of affected countries can be found here:

http://www.paho.org/hq/index.php?option=com_content&view=article&id=11603&Itemid=41696

M. PANORAMA DATA ENTRY DETAILS AND REFERENCES

M.1 Pregnancy warning and data entry

For records of cases or contacts who are pregnant:

- Add the client warning "Communicable Disease Alert Potential Vertical Transmission see notes" and create a Note Note Subject: CD Alert- Post Natal Follow Up Note Content: Initiate post-natal CD follow up for baby
- 2) Add the risk factor "Special Population Pregnancy Relevant to Disease Investigation"

If ordering physician is NOT the professional providing perinatal care, record contact information for the professional providing perinatal care (e.g. physician, midwife) under >Subject >>Client Details >>>Health Services

Please notify the Reproductive Infectious Diseases Clinic at BC Women's Hospital of this case (Tel: 604-875-2424 ext. 5212, Fax 604-875-2871).

M.2 Risks associated with sexual transmission

Potential risks of sexual transmission of Zika virus, contraceptive measures, and safer sexual practices should be discussed with cases (where appropriate) Refer to latest BC Zika case management and testing recommendations found at: http://www.bccdc.ca/health-info/diseases-conditions/zika-virus/information-for-health-professionals



M.3 Donation/receipt of blood, organs, or tissue

To report a *transfusion transmissible infection* for a case who has <u>received</u> blood, tissue or organ(s), create an Acquisition Event on the Exposure Summary screen (under Investigation on the left hand navigation) using the Acquisition Event Details screen.

For blood:

Exposure Name: XXX-TTI-Disease Name where XXX is the Health Authority recording/creating exposure (FNHA, IHA, VIHA, FHA, or NHA)

Potential Mode of Acquisition: Transfusion transmitted Nature of Exposure: Received other blood/blood products

Exposure Start: Date of transfusion

note: when exact date is unknown, enter estimate based on available information and select the "Estimated" flag

Exposure Location Name: same as Exposure Name Exposure Setting Type: Facility – non-recreational

Exposure Setting: Hospital

Address: Details for facility where transfusion occurred

For tissue or organs:

Exposure Name: XXX-TTI-Disease Name where XXX is the Health Authority recording/creating exposure (FNHA, IHA, VIHA, FHA, or NHA)

Source description: Tissues/Organs Potential Mode of Acquisition: Other Nature of Exposure: *leave blank* Exposure Start: Date of operation

note: when exact date is unknown, enter estimate based on available information and select the "Estimated" flag

Exposure Location Name: same as Exposure Name Exposure Setting Type: Facility – non-recreational Exposure Setting: Hospital

Address: Details for facility where operation occurred

To report a *transfusion transmissible infection* for a case who has <u>donated</u> blood, tissue or organ(s), create a Transmission Event on the Exposure Summary screen (under Investigation on the left hand navigation) using the Transmission Event Details screen.

For blood

Exposure Name: XXX-TTI-Disease Name where XXX is the Health Authority recording/creating exposure (FNHA, IHA, VIHA, FHA, or NHA)

Mode of Transmission: Transfusion transmitted Nature of Transmission: Donated blood/blood products

Exposure Start: Date donated blood

note: When exact date is unknown, enter estimate based on available information and select the "Estimated" flag

Exposure Location Name: same as Exposure Name Exposure Setting Type: Facility – non-recreational Exposure Setting: Canadian Blood Services Address: Details for facility where blood was donated

For tissue or organs:

Exposure Name: XXX-TTI-Disease Name where XXX is the Health Authority recording/creating exposure (FNHA, IHA, VIHA, FHA, or NHA)

Source description: Tissues/Organs Mode of Transmission: Other Nature of Transmission: *leave blank*

Transmission Event Date/Time > Exposure Start: Date of operation

note: When exact date is unknown, enter estimate based on available information and select the "Estimated" flag

Exposure Location Name: same as Exposure Name Exposure Setting Type: Facility – non-recreational Exposure Setting: Hospital

Address: Details for facility where operation occurred

Training Materials (https://panoramacst.gov.bc.ca): Exposures-Reference Guide-Investigations

System Guidelines (https://panoramacst.gov.bc.ca): Transfusion Transmissible Infections-Data Capture Guideline – Investigations, Exposures-Data Capture Guideline – Investigations, Transfusions, Transfusions, Exposures-Data Capture Guideline – Investigations, Exposures-Data Capture Guideline – Investigations, Transfusions,

Investigations

M.4 Vaccination entry into Immunization module

If the case has been previously vaccinated and the vaccination has been recorded in Panorama, the record can be viewed in the Immunization module

If the case has been previously vaccinated and the vaccination was not recorded, enter the data in the Immunization module as per Panorama Training Materials

Note: When exact date is unknown, enter estimate based on available information and select the "Estimated" flag (in 'Add historical details' screen)

M.5 Outcome at time of reporting

Asymptomatic cases can be classified as Other with 'Asymptomatic' in text box