



Pregnancy and Zika Virus Disease Surveillance Form

These data are considered confidential and will be stored in a secure database at the Centers for Disease Control and Prevention

Please return completed form by sending an encrypted email to ZIKApregnancy@cdc.gov or by fax to the secure number: 404-718-2200. Pregnancy & Birth Defects Task Force phone number: 770-488-7100

Mother's Zika virus infection (ADB follow-up)		
Mother's name: Last First MI		Maiden name (if applicable)
State/Territory ID:	DOB: ____/____/____	State/Territory of residence:
County of residence:	Ethnicity: ☐ Hispanic or Latino ☐ Not Hispanic or Latino	
Race (check all that apply): ☐ American Indian or Alaska Native ☐ Asian ☐ Black or African-American ☐ Native Hawaiian or other Pacific Islander ☐ White		
Indication for maternal Zika virus testing: ☐ Exposure history, no known fetal concerns ☐ Exposure history and fetal concerns		
Date of Zika virus symptom onset: ____/____/____ OR- ☐ Asymptomatic <i>If date not known, trimester of symptom onset</i> _____		
Hospitalized for Zika virus disease ☐ No ☐ Yes Maternal Death ☐ No ☐ Yes		
Symptoms of mother's Zika virus disease: (check all that apply) ☐ Fever ____°F (if measured) ☐ Rash ☐ Arthralgia ☐ Conjunctivitis ☐ Other Clinical Presentation _____		
If symptomatic, gestational age at onset: _____ weeks <i>If gestational age not known, trimester of symptom onset</i> _____		Travel history: ☐ No ☐ Yes
Was Zika virus infection acquired in place of residence ☐ No ☐ Yes, if yes, skip to the section on Mother's pregnancy		
If TRAVEL DURING PREGNANCY , answer questions below. If not, skip to non-traveling woman		
Country(s) of exposure (1) _____	Travel start ____/____/____	Travel end ____/____/____
Mother's sexual partner(s)? please check all that apply ☐ Male ☐ Female		
Did any male sexual partner(s) travel on this trip? ☐ No ☐ Yes ☐ Unknown		
If yes, did any male partner(s) have an illness that included fever, rash, arthralgia, or conjunctivitis during or within 2 weeks of travel? ☐ No ☐ Yes ☐ Unknown		
If yes, was there unprotected sexual contact while male partner(s) had illness? ☐ No ☐ Yes ☐ Unknown		
If male partner(s) traveled, did he have a test that showed lab evidence of Zika? ☐ No ☐ Yes ☐ Unknown		
Country(s) of exposure (2) _____	Travel start ____/____/____	Travel end ____/____/____
Mother's sexual partner(s)? please check all that apply ☐ Male ☐ Female		
Did any male sexual partner(s) travel on this trip? ☐ No ☐ Yes ☐ Unknown		
If yes, did any male partner(s) have an illness that included fever, rash, joint pain, or pink eye during or within 2 weeks of travel? ☐ No ☐ Yes ☐ Unknown		
If yes, was there unprotected sexual contact while male partner(s) had illness? ☐ No ☐ Yes ☐ Unknown		
If male partner(s) traveled, did he have a test that showed lab evidence of Zika? ☐ No ☐ Yes ☐ Unknown		
Country(s) of exposure (3) _____	Travel start ____/____/____	Travel end ____/____/____
Mother's sexual partner(s)? please check all that apply ☐ Male ☐ Female		



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Did any male sexual partner(s) travel on this trip? ☐ No ☐ Yes ☐ Unknown	
If yes, did any male partner(s) have an illness that included fever, rash, joint pain, or pink eye during or within 2 weeks of travel? ☐ No ☐ Yes ☐ Unknown	
If yes, was there unprotected sexual contact while male partner(s) had illness? ☐ No ☐ Yes ☐ Unknown	
If male partner(s) traveled, did he have a test that showed lab evidence of Zika? ☐ No ☐ Yes ☐ Unknown	
NON-TRAVELLING WOMAN: other possible exposures?	
☐ Sexual partner w/travel history, symptomatic, lab evidence of Zika	
☐ Sexual partner w/travel history, symptomatic, no test results	
☐ Sexual partner w/travel history, <u>asymptomatic</u> , lab evidence Zika	
☐ Other, please describe _____	
☐ Unknown exposure history	
Mother's pregnancy (DRH/DBDDD follow-up)	
Last menstrual period (LMP): ____/____/____	Estimated delivery date: ____/____/____
Estimated delivery date based on (check all that apply): ☐ LMP ____/____/____	☐ U/S (1 st trimester)
☐ U/S (2 nd trimester)	☐ U/S (3 rd trimester)
History: # pregnancies ____ # living children ____ # miscarriages ____ # elective terminations ____	
Prior fetus/infant with microcephaly: ☐ No ☐ Yes If yes, genetic cause: ☐ No ☐ Yes	
Gestation: ☐ Single ☐ Twins ☐ Triplets+	
Underlying maternal illness:	
Diabetes ☐ No ☐ Yes Maternal PKU ☐ No ☐ Yes Hypothyroidism ☐ No ☐ Yes Hypertension ☐ No ☐ Yes	
Substance use during this pregnancy: Alcohol use ☐ No ☐ Yes Cocaine use ☐ No ☐ Yes Smoking ☐ No ☐ Yes	
Other underlying illness: _____	
Complications of pregnancy:	
Toxoplasmosis ☐ Negative ☐ Positive ☐ Unknown	Cytomegalovirus ☐ Negative ☐ Positive ☐ Unknown
Herpes Simplex ☐ Negative ☐ Positive ☐ Unknown	Rubella ☐ Negative ☐ Positive ☐ Unknown
Syphilis ☐ Negative ☐ Positive ☐ Unknown	
Fetal genetic abnormality ☐ No ☐ Yes, <i>diagnosis</i> _____ ☐ Unknown	
Gestational diabetes ☐ No ☐ Yes Pregnancy-related HTN ☐ No ☐ Yes Intrauterine death of a twin ☐ No ☐ Yes	
Other _____	
Medications during pregnancy: ☐ No ☐ Yes (please list type and see guide for further instructions)	
Did this pregnancy end in miscarriage or intrauterine fetal demise (IUFD)? ☐ No ☐ Yes Date: ____/____/____ Gestational age ____ weeks	Was this pregnancy terminated? ☐ No ☐ Yes Date: ____/____/____ Gestational age ____ weeks
Maternal Prenatal Imaging and Diagnostics	
Date(s) of Ultrasound(s):	
Overall Fetal Ultrasound Results: ☐ Normal ☐ Abnormal	



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____/____/____ ☐ check if date approximated if date not known, gestational age _____ weeks	☐ reported by patient/healthcare provider ☐ ultrasound report
	Head Circumference _____cm ☐ Normal ☐ Abnormal (by physician report)
	Biparietal diameter _____cm Femur Length _____cm Abdominal circumference _____cm
	☐ Symmetrical intrauterine growth restriction (IUGR) (<5% EFW)
	☐ Asymmetrical IUGR (HC<FL or HC <AC)
	Intracranial calcifications ☐ No ☐ Yes Ventriculomegaly ☐ No ☐ Yes
	Cerebral atrophy ☐ No ☐ Yes Ocular anomalies ☐ No ☐ Yes
	Cerebellar abnormalities ☐ No ☐ Yes Arthrogryposis ☐ No ☐ Yes
Lissencephaly ☐ No ☐ Yes Pachygyria ☐ No ☐ Yes	
Hydranencephaly ☐ No ☐ Yes Porencephaly ☐ No ☐ Yes	
Corpus callosum abnormalities ☐ No ☐ Yes Hydrops ☐ No ☐ Yes	
Ascites ☐ No ☐ Yes Other ☐ No ☐ Yes, describe	
Description of abnormal ultrasound findings:	
____/____/____ ☐ check if date is approximated if date not known, gestational age _____ weeks	Overall Fetal Ultrasound Results: ☐ Normal ☐ Abnormal
	☐ reported by patient/healthcare provider ☐ ultrasound report
	Head Circumference _____cm ☐ Normal ☐ Abnormal (by physician report)
	Biparietal diameter _____cm Femur Length _____cm Abdominal circumference _____cm
	☐ Symmetrical IUGR (<5% EFW) ☐ Asymmetrical IUGR (HC<FL or HC <AC)
	Intracranial calcifications ☐ No ☐ Yes Ventriculomegaly ☐ No ☐ Yes
	Cerebral atrophy ☐ No ☐ Yes Ocular anomalies ☐ No ☐ Yes
	Cerebellar abnormalities ☐ No ☐ Yes Arthrogryposis ☐ No ☐ Yes
Lissencephaly ☐ No ☐ Yes Pachygyria ☐ No ☐ Yes	
Hydranencephaly ☐ No ☐ Yes Porencephaly ☐ No ☐ Yes	
Corpus callosum abnormalities ☐ No ☐ Yes Hydrops ☐ No ☐ Yes	
Ascites ☐ No ☐ Yes Other ☐ No ☐ Yes, describe	
Description of abnormal ultrasound findings:	
____/____/____ ☐ check if date is approximated if date not known, gestational age _____ weeks	Overall Fetal Ultrasound Results: ☐ Normal ☐ Abnormal
	☐ reported by patient/healthcare provider ☐ ultrasound report
	Head Circumference _____cm ☐ Normal ☐ Abnormal (by physician report)
	Biparietal diameter _____cm Femur Length _____cm Abdominal circumference _____cm
☐ Symmetrical IUGR (<5% EFW) ☐ Asymmetrical IUGR (HC<FL or HC <AC)	
Intracranial calcifications ☐ No ☐ Yes Ventriculomegaly ☐ No ☐ Yes	
Cerebral atrophy ☐ No ☐ Yes Ocular anomalies ☐ No ☐ Yes	
Cerebellar abnormalities ☐ No ☐ Yes Arthrogryposis ☐ No ☐ Yes	



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_____ weeks	Lissencephaly	☐ No ☐ Yes	Pachygyria	☐ No ☐ Yes
	Hydranencephaly	☐ No ☐ Yes	Porencephaly	☐ No ☐ Yes
	Corpus callosum abnormalities	☐ No ☐ Yes	Hydrops	☐ No ☐ Yes
	Ascites	☐ No ☐ Yes	Other	☐ No ☐ Yes, describe

Description of abnormal ultrasound findings:**For additional ultrasounds, please request a supplementary ultrasound form****Fetal MRI performed:** ☐ No ☐ Yes (please answer questions below)

____/____/____ ☐ check if date is approximated if date not known, gestational age _____ weeks	Overall Fetal MRI Results: ☐ Normal ☐ Abnormal			
	☐ reported by patient/healthcare provider ☐ ultrasound report			
	Head Circumference ____cm ☐ Normal ☐ Abnormal (by physician report)			
	Biparietal diameter ____cm Femur Length ____cm Abdominal circumference ____cm			
	☐ Symmetrical IUGR (<5% EFW) ☐ Asymmetrical IUGR (HC<FL or HC <AC)			
	Intracranial calcifications ☐ No ☐ Yes		Ventriculomegaly ☐ No ☐ Yes	
	Cerebral atrophy ☐ No ☐ Yes		Ocular anomalies ☐ No ☐ Yes	
	Cerebellar abnormalities ☐ No ☐ Yes		Arthrogryposis ☐ No ☐ Yes	
Lissencephaly ☐ No ☐ Yes		Pachygyria ☐ No ☐ Yes		
Hydranencephaly ☐ No ☐ Yes		Porencephaly ☐ No ☐ Yes		
Corpus callosum abnormalities ☐ No ☐ Yes		Hydrops ☐ No ☐ Yes		
Ascites ☐ No ☐ Yes		Other ☐ No ☐ Yes, describe		

Description of abnormal MRI findings:**Amniocentesis performed:** ☐ No ☐ Yes (date: ____/____/____)Zika virus testing: ☐ Not performed ☐ Yes, if yes test results: ☐ negative for Zika ☐ lab evidence of ZikaNon-Zika infection detected ☐ No ☐ Yes if yes, what infection(s) detected _____Genetic abnormality detected ☐ No ☐ Yes Please Describe:**Provider Information****Provider name:** ☐ Dr. ☐ PA ☐ RN ☐ Mr. ☐ Ms. _____
Last First MI**Phone:** _____ **Email:** _____ **Date of form completion** ____/____/____**Name of person completing form: (if different from provider)** _____
Last First MI**Hospital/facility:** _____**Phone:** _____ **Email:** _____ **Date of form completion** ____/____/____**Health Department Information**



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Name of person completing form: _____

Phone: _____ Email: _____ Date of form completion ____/____/____

FOR INTERNAL CDC USE ONLY

Mother ID: _____

State/Territory ID: _____

Zika T ID: _____

R number: _____ Mother infection type: ☐ Confirmed ☐ Probable ☐ Possible

Public reporting burden of this collection of information is estimated to average 15 minutes per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. An agency may not conduct or sponsor, and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to CDC/ATSDR Reports Clearance Officer; 1600 Clifton Road NE, MS E-11, Atlanta, Georgia 30333; ATTN: PRA (0920-1101).