

STATE OF CONNECTICUT

Connecticut Department of Public Health Dr. Katherine A. Kelley State Public Health Laboratory 395 West Street, Rocky Hill, CT 06067

10/05/2017

Dear Healthcare Provider,

Due to recent changes in the national Zika virus testing guidelines (available at http://www.ct.gov/dph/lib/dph/infectious_diseases/zikavirus/dph_zika_virus_testing_protocol_fnl.pdf), the Connecticut Department of Public Health (DPH) has made modifications to the protocol for how specimens should be submitted to the State Public Health Laboratory (SPHL) for Zika virus testing. Specimens will no longer require approval from the Epidemiology and Emerging Infections Program before being accepted for testing. However, testing of clinical specimens for viral RNA by RT-PCR and IgM antibodies by ELISA will continue to be offered at the SPHL <u>only</u> for patients who meet specific clinical and exposure criteria as defined in the link provided above. Therefore, in order to determine acceptability for testing, the SPHL will require additional information that is not currently provided on the Clinical Test Requisition, Form OL-9B. Please use the attached Zika Clinical Test Requisition submission form just for Zika virus testing to provide the required information.

- **Pregnant women:** collect serum (\geq 3 ml) and urine (\geq 1 ml) within 12 weeks of potential exposure
- Infants: collect serum (≥ 1 ml) and urine (≥ 1 ml) within 2 days of birth
- Urine should be collected in a sterile container with a tight fitting screw cap (preferably film secured)
- Specimens can be stored refrigerated (2-8°C) and shipped to the SPHL with adequate ice packs if they are received at the SPHL within 3 days of collection.
- If transit time is expected to exceed 3 days, freeze specimens (-70°C) and ship on dry ice to ensure they remain frozen while in transit to the SPHL.
- When submitting **placental tissue** include sections of the placental disk (3 full thickness pieces from the middle third and one from the margin), fetal membranes 5x12 cm strip, and pathologic lesions when possible. Fix specimens in formalin with a volume 10x the mass of tissue. Specimens can be shipped at room temperature.

Specimens should be packaged and shipped as *Category B Biological Substances* in accordance with the Department of Transportation Hazardous Materials Regulations (49 CFR Part 171-180).

For questions regarding **Zika virus testing criteria** please call the DPH Epidemiology and Emerging Infections Program at **860-509-7994**. For questions regarding **specimen handling** please call the SPHL at **860-920-6662 or 920-6506**.

Name and Address of Authorized Submitter

ZIKA VIRUS CLINICAL TEST REQUISITION

STATE OF CONNECTICUT Dr. Katherine A. Kelley State Public Health Laboratory 395 West Street, Rocky Hill, CT 06067 CLIA ID 07D0644555 / CT License CL-0197 Phone 860-920-6500 CLIENT SERVICES 860-920-6635

♦ DENOTES <u>REQUIRED</u> INFORMATION

ACCESSION LABEL

FOR CTDPH LABORATORY USE ONLY

LAB PROFILE Number:

Section 1: Patient Information (Please Print Clearly)							
♦ Name (Last, First, M.I.) or Identifier:							
♦ Street Address:				♦ City, State, Zip:			
♦ Date of Birth:	Gender:		ale				
Section 2: Specimen Informatio	n						
Submitter Sample ID:		Date Collected:			Time Collected:		
♦ Specimen Source/Type:							
Blood Serum	Urine			Amniotic Fluid			
□ Fetal/Placental Tissue; specify ty	pe						
• Specimen Storage (Prior to Delivery):		□ Refrigerated (2-8° C)		□ Frozen (<-20° C)) Ambient Temperature		
Specimen Transport/Delivery:		□ Cold (Ice pack)		□ Frozen (Dry Ice) □ Ambient Temperature		perature	
For questions regarding specimen handling please call the Virology laboratory of the CT SPHL at 860-920-6662							
♦ <u>REQUIRED INFORMATION FOR ZIKA VIRUS TESTING</u> Testing of clinical specimens for viral RNA by RT-PCR and IgM antibodies by ELISA is offered at the CT SPHL ONLY							
for patients who meet clinical at Pregnant Yes No Estima International travel during pregnar Unprotected sex during pregnancy wit Zika Virus associated Symptoms: / Date of Symptom Onset Fetus or infant with Congenital Zik *Reference: CDC Defines Cong FOR INFANTS: Mother's Zika viru Ordering Healthcare Provider:	ted due ncy? th someo Arthralgia ca Virus enital Z	date: Yes □ No Locat one who has travelled a □ Conjunctivitis (i Syndrome* □ Yes ïka Syndrome – Me	d withir non-pu s □ N edscaj	n the previous 2 weel rulent)			
Section 3: ZIKA VIRUS TESTING ONLY							
Zika Virus Testing (IgM MAC-ELISA and Flavivirus Trioplex rRT-PCR) For pregnant women, current Zika virus testing guidelines* recommend PCR on serum and urine and IgM on serum, all done concurrently on specimens collected within 12 weeks of potential exposure. For infants, guidelines recommend PCR testing of urine and serum collected within 2 days of birth. * http://www.ct.gov/dph/lib/dph/infectious diseases/zikavirus/dph zika virus testing protocol fnl.pdf For Laboratory Use Only Comments							