Arbovirus Panel (Encephalitis Viruses)

Test	Detection of virus specific IgM and neutralizing antibodies to arbovirus
Description	infection. The arbovirus(es) suspected must be specified.
Test use	To aid in the diagnosis of current or past infection with arboviruses
Test	Virology
Department	Phone: (860) 920-6662, FAX: (860) 920-6661
Methodology	IgM capture ELISA, IgG ELISA, additional procedures as required
Availability	Specimen is referred to the Centers for Disease Control and Prevention in Fort
	Collins, CO for testing.
	0.5 mL serum and/or 1 mL cerebral spinal fluid
	NOTE: If specimen collection occurs within 8 days after symptom onset, a
Specimen	convalescent serum specimen will be requested. Acute and convalescent
Requirements	serum specimens, if available, should be sent together. Acute specimens
	should be collected 3-10 days after symptom onset. Collect convalescent
Callestian	specimen 2-3 weeks after acute sample.
Collection	To obtain collection kit, refer to Collection Kit Ordering Information
kit/Container	
Collection Instructions	Standard venipuncture technique
	Aseptically obtained spinal fluid
Specimen	Keep serum cold or frozen Transport with an iso pack coolant
Handling & Transport	Transport with an ice pack coolant
Unacceptable	Unlabeled specimen
Conditions	Specimens that have leaked or containers that have broken in transit
conditions	Whole blood
Requisition	Clinical Test Requisition (select Arbovirus Panel (Encephalitis Viruses)). The
Form	arbovirus(es) suspected must be specified.
	Name and address of submitter (and/or Horizon profile #)
Required	Patient name or identifier, town of residence (city, state, zip), date of birth
Information	Specimen source/type, date collected, test requested
	Date of onset of patient symptoms
	Pertinent travel history (3 months prior to symptom onset)
	Please ensure patient name on the requisition matches that on the specimen.
	 Testing requires approval of Epidemiology and Emerging Infections,
	(860) 509-7994.
Limitations	 If initial serological testing is positive, further confirmatory tests are
	done which may delay reporting of the final results.
Additional	Testing is limited to the second interaction with this second states and / second side is the second
	Testing is limited to those patients exhibiting symptoms and/or travel history

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