



### Effective as of May 1, 2023

## Additional ordering and billing information

Information when ordering laboratory tests that are billed to Medicare/Medicaid

Information regarding Current Procedural Terminology (CPT)

Test Number	Mnemonic	Test Name	New Test	Test Name Change	Specimen Requirements	Methodology	Performed/Reported	Note	Interpretive Data	Reference Interval	Component Charting Name	Component Change	Reflex Pattern	Result Type	Ask at Order Prompt	Numeric Map	Unit of Measure	CPT Code	Pricing Change	Inactivation w/ Replacement	Inactivation w/o Replacement
0099408	AL U	Aluminum, Urine (Inactive as of 05/01/23)																			x
2002483	HCV AB	Hepatitis C Virus Antibody by CIA (Change effective as of 05/01/23: Refer to 2010784)																		x	
2007220	ECHINO IGG	Echinococcus Antibody, IgG			x	x			х	х						х	x				
3006254	JCV AB	JC Virus Antibody by ELISA, Serum with Reflex to Inhibition Assay	x																		
3016444	PHOSPHO T	Phospho-Tau/Total- Tau/A Beta42, CSF	x																		



### **TEST CHANGE**

Echinococcus Antibody, IgG

2007220, ECHINO IGG

Specimen Requirements:								
Patient Preparation:								
Collect: Serum separator tube (SST <u>) or plain red.</u> }-								
Specimen Preparation:	Separate serum from cells ASAP or within 2 hours of collection. Transfer 1 mL serum to an ARUP standard transport tubeStandard Transport Tube. (Min: 0.15 mL) Parallel testing is preferred and convalescent specimens must be received within 30 days from receipt of acute specimens. Mark specimens plainly as acute or convalescent.							
Transport Temperature:	<u>Preferred transport temp:</u> Refrigerated. <u>Also acceptable: Frozen</u>							
Unacceptable Conditions: Contaminated, <u>heat-inactivated</u> , <u>grossly hemolyzed</u> , or severel lipemic specimens.								
Remarks:								
Stability: After separation from cells: Ambient: 48 hours; Refrigerated: 2 weeks; Frozen: 1 monthyear (avoid repeated freeze/thaw cycles)								
Methodology:	Semi-Quantitative Enzyme-Linked Immunosorbent Assay (FLISA)							
Performed: Mon, Thu								
Reported: 1-5 days								
Note:								
CPT Codes: 86682								
New York DOH Approval Status: This test is New York DOH approved.								
Interpretive Data:								
Patients with collagen vascular diseases, hepatic cirrhosis, schistosomiasis, and other parasitic infections can produce false-positive results. There is a strong cross-reaction between echinococcosis- and cysticercosis-positive sera.								
Seroconversion between acute and convalescent sera is considered strong evidence of recent								

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specimens where both tests are done in the same laboratory at the same time.

infection. The best evidence for infection is a significant change on two appropriately timed  ${\bf r}$ 



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Component	Interpretation
Echinococcus	0-8
Antibody IgG	UNegative:
	No significant level of
	Echinococcus IgG
	antibodies detected. 9-11
	UEquivocal:
	Recommend repeat
	testing in 2-4 weeks with
	fresh sample. 12 U or
	greaterPositive: IgG
	antibodies to
	Echinococcus detected,
	indicating current or past
	infection.

#### Reference Interval:

Test Number  Number  Number  Number  Number  Components Reference  Interval					4	A			
level of Echinococcus lgG-antibody detected.  0.9-1.1 IV Questionable presence of Echinococcus lgG-antibody detected. Repeat testing in 10- 14-days may be helpful.  1.2-IV or Gresence of greater greater greater greater or greater current or past	Test	0.0-0.8	Negative - No	Components	Reference	<u>e</u>			
level of Echinococcus lgG-antibody detected.  0.9-1.1 IV Questionable presence of Echinococcus lgG-antibody detected. Repeat testing in 10- 14-days may be helpful.  1.2-IV or Gresence of greater greater greater greater or greater current or past	Number	· <del>IV</del>	significant		Interval				
lgG antibody detected.  0.9-1.1 IV Equivocal-Questionable presence of Echinococcus lgG antibody detected. Repeat testing in 10-14 days may be helpful.  1.2 IV. or Presence of greater lgG antibody to Echinococcus detected, suggestive of current or past			level of						
detected.  0.9-1.1 IV Questionable presence of Echinococcus IgG antibody detected. Repeat testing in 10-14 days may be helpful.  1.2.IV Positive-Presence of greater IgG antibody to Echinococcus detected, suggestive of current or past			Echinococcus						
0.9-1.1  IV  Questionable presence of Echinococcus IgG antibody detected. Repeat testing in 10-14 days may be helpful.  1.2-IV  or  greater gr			IgG antibody						
IV Questionable presence of Echinococcus IgG antibody detected. Repeat testing in 10-14 days may be helpful.  1.2.IV Positive-Presence of greater IgG antibody to Echinococcus detected, suggestive of current or past			detected.						
presence of Echinococcus IgG antibody detected. Repeat testing in 10- 14 days may be helpful.  1.2.IV or greater greater teg antibody to Echinococcus detected, suggestive of current or past		0.9-1.1	Equivocal -						
Echinococcus IgG antibody detected. Repeat testing in 10- 14 days may be helpful.  1.2 IV or Greater Greater  Echinococcus detected, suggestive of current or past		₩	Questionable						
Echinococcus IgG antibody detected. Repeat testing in 10- 14 days may be helpful.  1.2 IV or Greater Greater  Echinococcus detected, suggestive of current or past			presence of						
detected. Repeat testing in 10- 14 days may be helpful.  1.2.1V. Positive- Presence of greater gG antibody to Echinococcus detected, suggestive of current or past									
detected. Repeat testing in 10- 14 days may be helpful.  1.2.1V. Positive- Presence of greater gG antibody to Echinococcus detected, suggestive of current or past			IgG antibody						
testing in 10- 14 days may be helpful.  1.2 IV or resence of greater lgG antibody to Echinococcus detected, suggestive of current or past									
1.2 IV. or Positive Presence of greater to  Echinococcus detected, suggestive of current or past			Repeat						
be helpful.  1.2.IV. Positive- or Presence of greater greater telegicantibody			testing in 10-						
1.2 IV or greater greater  Echinococcus detected, suggestive of current or past			14 days may						
Presence of IgG antibody to Echinococcus detected, suggestive of current or past			<del>be helpful.</del>						
greater to Echinococcus detected, suggestive of current or past		1.2 IV	Positive -						
to Echinococcus detected, suggestive of current or past		or	Presence of						
to Echinococcus detected, suggestive of current or past		greater	IgG antibody						
detected, suggestive of current or past									
suggestive of current or past			Echinococcus						
current or past			detected,						
past			suggestive of						
			current or						
infection			past						
in court			infection.						
Echinococcus 8 U or				Echinococcus	8 U or				
Antibody IgG less									

HOTLINE NOTE: There is a numeric map change associated with this test. Refer to the Hotline Test Mix for interface build information.

HOTLINE NOTE: There is a unit of measure change associated with this test. Refer to the Hotline Test Mix for interface build information.

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**NEW TEST** 

**Click for Pricing** 

JC Virus Antibody by ELISA, Serum with Reflex to Inhibition Assay

3006254, JCV AB

Specimen Requirements: **Patient Preparation:** Collect: Plain red or serum separator tube (SST). Also acceptable: Lavender (K2EDTA) Specimen Preparation: Transfer 1 mL serum or plasma to an ARUP standard transport tube. (Min: 0.5 mL) Test is not performed at ARUP; separate specimens must be submitted when multiple tests are ordered. **Transport Temperature:** Refrigerated. Also acceptable: Room temperature or frozen. **Unacceptable Conditions:** Remarks: Stability: Ambient: 1 week; Refrigerated: 2 weeks; Frozen: 3 months Methodology: Enzyme-Linked Immunosorbent Assay (ELISA) Performed: Varies Reported: 3-7 days Note: If antibody result is indeterminate, then a confirmation (inhibition) assay will be added. **CPT Codes:** 86711; if reflexed, add 86711 New York DOH Approval Status: This test is New York DOH approved. Interpretive Data: Reference Interval: By report

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HOTLINE NOTE: Refer to the Hotline Test Mix for interface build information.



**NEW TEST** 

**Click for Pricing** 

Phospho-Tau/Total-Tau/A Beta42, CSF

3016444, PHOSPHO T

Specimen Requirements:

**Patient Preparation:** 

Collect: CSF

Specimen Preparation: Transfer 2 mL CSF to an ARUP standard transport tube. (Min:

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0.5 mL).

Transport Temperature: Frozen

Unacceptable Conditions: Specimens with cell count greater than 500 erythrocytes/mm3.

Remarks:

Stability: Ambient: 72 hours, Refrigerated: 21 days, Frozen: 4 months

Methodology: Enzyme-Linked Immunosorbent Assay (ELISA)

Performed: Varies

Reported: 7-19 days

Note:

CPT Codes: 83520 x3

New York DOH Approval Status: This test is New York DOH approved.

Interpretive Data:

Reference Interval:

Test Components Reference Interval

HOTLINE NOTE: Refer to the Hotline Test Mix for interface build information.



# **Inactivations**

The following will be discontinued from ARUP's test menu on May 1, 2023 Replacement test options are indicated when applicable.

Test Number	Test Name	Refer to Replacement Test
0099408	Aluminum, Urine (Inactive as of 05/01/23)	
2002483	Hepatitis C Virus Antibody by CIA (Change effective as of 05/01/23: Refer to 2010784)	Hepatitis C Virus Antibody by CIA with Reflex to HCV by Quantitative NAAT (2010784)