

TEST DOWN NOTIFICATION

NOTIFICATION DATE: August 6, 2009

EFFECTIVE DATE: Immediately

**Cysticercosis, IgG Antibody, Western Blot Assay, Serum
#81354**

EXPLANATION: Due to reagent issues, this test will be down until further notice. Specimens received for Unit Code #81354 will be sent to ARUP Laboratories for #91947 Cysticercosis Antibody, IgG by Western Blot.

REFERRAL ALTERNATIVE TEST: #91947, Cysticercosis Antibody, IgG by Western Blot, to ARUP Laboratories.

METHODOLOGY: Western Blot (WB)

REFERENCE VALUES: Negative - No significant level of detectable *T. solium* IgG antibody.

Positive - IgG antibody to *T. solium* detected, which may suggest current or past infection.

SPECIMEN REQUIREMENTS: Draw blood in a serum gel tube(s) and separate serum from cells as soon as possible. Transfer 1 mL of serum to a plastic vial and ship refrigerate.

LIST FEE: \$215.85

CPT CODE: 86682

ANALYTIC TIME: 1-8 days

DAY(S) SET-UP: Thursdays

QUESTIONS: Contact your Mayo Medical Laboratories' Regional Manager
Kim J. Baker, Mayo Medical Laboratories' Technologist Support
Telephone: 800-533-1710

1-800-533-1710

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| PATIENT NAME TESTING, 91947 | | PATIENT NUMBER | | AGE 64 | SEX F | ACCESSION # W2193564 |
| ORDERING PHYSICIAN | | CLIENT ORDER # | | | | ACCOUNT # LIAISONS |
| COLLECTION 08/05/09 03:44 P | RECEIVED 08/05/09 03:44 P | REPORT PRINTED 08/06/09 02:49 P | | SPECIMEN INFORMATION DATE OF BIRTH: 6/20/1945 | | |
| DATE TIME | DATE TIME | DATE TIME | | | | |
| Test Client Attn: Mayo Liaisons 200 First Street SW Rochester, MN 55905 507-284-8202 | | | | | | |

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| HI | LO | REF RANGE | PERFORM SITE * |
| TEST REQUESTED | | | |

Cysticercosis Ab, IgG by WB

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| Cysticercosis Ab, IgG by WB | Negative | REF |
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Interpretation: Cysticercosis Ab, IgG by WB

Negative No significant level of detectable T. solium IgG antibody.

Positive IgG antibody to T. solium detected, which may suggest current or past infection.

This test uses a kit designated by the manufacturer as "for research use, not for clinical use." The performance characteristics of this test were validated by ARUP Laboratories, Inc. The U.S. Food and Drug Administration (FDA) has not approved this test. The results are not intended to be used as the sole means for clinical diagnosis or patient management decisions. ARUP is authorized under Clinical Laboratory Improvement Amendments (CLIA) and by all states to perform high-complexity testing.

Test Performed by: ARUP Laboratories, Inc.
 500 Chipeta Way
 Salt Lake City, UT 84108

* PERFORMING SITE

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| PATIENT NAME TESTING, 91947 | ORDER STATUS Final | COLLECTION DATE AND TIME 08/05/09 03:44 P |
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