Lab Dept: Serology

Test Name: STRONGYLOIDES ANTIBODY, IGG

General Information

Lab Order Codes: STRNG

Synonyms: N/A

CPT Codes: 86682 – Antibody, helminth, not elsewhere specified

Test Includes: Strongyloides IgG antibody reported as Negative, Equivocal or Positive.

Logistics

Test Indications: Screening for the presence of IgG-class antibodies to *Strongyloides*

This test is not useful for monitoring patient response to therapy as IgGclass antibodies to *Strongyloides* may remain detectable following

resolution of infection.

Lab Testing Sections: Serology - Sendouts

Referred to: Mayo Clinic Laboratories (MML Test: STRNG)

Phone Numbers: MIN Lab: 612-813-6280

STP Lab: 651-220-6550

Test Availability: Daily, 24 hours

Turnaround Time: 1 - 4 days

Special Instructions: N/A

Specimen

Specimen Type: Blood

Container: SST (Gold, marble or red) tube

Draw Volume: 1.5 mL (Minimum: 1.2 mL) blood

Processed Volume: 0.5 mL (Minimum: 0.4 mL) serum

Collection: Routine venipuncture

Special Processing: Lab Staff: Centrifuge specimen. Remove serum and aliquot into a plastic

screw-capped tube. Store and ship at refrigerated temperatures. Forward

promptly.

Specimen stable refrigerated (preferred) or frozen for 30 days.

Patient Preparation: None

Sample Rejection: Warm specimens; gross lipemia, hemolysis or icterus; bacterially

contaminated; heat-inactivated; mislabeled or unlabeled specimen

Interpretive

Reference Range: All ages: Negative

Critical Values: N/A

Limitations:

• False positive results may occur with other heminth infections, including prior exposure to Entamoeba histolytica, Ascaris, Taenia solium, Fasciola

spp, Echinococcus spp, Shistosoma spp, and Toxocara (per assay

manufacturer).

• This assay should not be used alone to establish a diagnosis of strongyloidiasis. Results should be correlated with other laboratory findings

and through clinical evaluation.

• False negative results may occur during acute or localized infection. A single negative result should not be used to rule out infection.

This assay should not be used to monitor response to therapy.

• The seroprevalence of IgG-class antibodies to Strongyloides stercoralis

ranges from 0 to 6.1% in the United States.

Methodology: Enzyme-Linked Immunosorbent Assay (ELISA)

References: Mayo Clinic Laboratories December 2024

Updates: 10/28/2015: Testing no longer forwarded to ARUP. Testing now performed

onsite at MML. No longer reported with IV units, specimen requirements

updated.

12/22/2017: Collection container update.

12/27/2024: Updated indications and sample rejection. Added specimen

stability.