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 IgG- class antibodies to <i>Strongyloides</i> 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 | | |
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| 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 <i>Entamoeba histolytica, Ascaris, Taenia solium, Fasciola spp, Echinococcus spp, Shistosoma spp</i>, and <i>Toxocara</i> (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 <i>Strongyloides stercoralis</i> 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. | | |