

TABLE 245e-2 ROUNDWORM INFECTIONS

| Parasite | Geographic Distribution | Life-Cycle Hosts | | Diagnosis | | | |
|---|---|---------------------------------|-----------------------------|---------------------|-------------------------------|--------------------------------------|--|
| | | Intermediate (Transmission) | Definitive | Parasite Stage | Body Fluid or Tissue | Serologic Tests | Other |
| Intestinal Roundworms | | | | | | | |
| <i>Enterobius vermicularis</i> (pinworm) | Temperate and tropical zones | Fecal-oral | Humans | Ova | Perianal skin | — | “Cellophane tape” test |
| <i>Trichuris trichiura</i> (whipworm) | Temperate and tropical zones | Soil, fecal-oral | Humans | Ova | Feces | — | Rectal prolapse |
| <i>Ascaris lumbricoides</i> (roundworm of humans) | Temperate and tropical zones | Soil, fecal-oral | Humans | Ova | Feces | — | Sx of pulmonary migration |
| <i>Ancylostoma duodenale</i> (Old World hookworm) | Eurasia, Africa, Pacific | Soil to skin | Humans | Ova/larvae | Feces | — | Sx of pulmonary migration, anemia |
| <i>Necator americanus</i> (New World hookworm) | U.S., Africa, worldwide | Soil to skin | Humans | Ova/larvae | Feces | — | Sx of pulmonary migration, anemia |
| <i>Strongyloides stercoralis</i> (strongyloidiasis) | Moist tropics and subtropics | Soil to skin | Humans | Larvae | Feces, sputum, duodenal fluid | EIA | Dissemination in immunodeficiency |
| <i>Capillaria philippinensis</i> | Southeast Asia, Taiwan, Egypt | Raw fish | Birds | Ova, larvae, adults | Feces | — | Malabsorption/ autoinfection, biopsy |
| Tissue Roundworms | | | | | | | |
| <i>Trichinella spiralis</i> (trichinellosis) | Worldwide | Swine/humans | Swine/humans | Larvae | Muscle | EIA | Muscle biopsy |
| <i>Wuchereria bancrofti</i> (filariasis) | Coastal areas in tropics and subtropics | Mosquitoes | Humans | Microfilariae | Blood, lymph nodes | EIA, RAPID, PCR ^b | Nocturnal periodicity ^a |
| <i>Brugia malayi</i> (filariasis) | Asia, Indian subcontinent | Mosquitoes | Humans | Microfilariae | Blood | EIA, RAPID, PCR ^b | Nocturnal |
| <i>Loa loa</i> (African eye worm) | West and Central Africa | Mango flies (<i>Chrysops</i>) | Humans | Microfilariae | Blood | LIPS ^b , PCR ^b | May be visible in eye, diurnal |
| <i>Onchocerca volvulus</i> (river blindness) | Africa, Mexico, Central and South America | Blackflies | Humans | Adults/larvae | Skin/eye | LIPS ^b , PCR ^b | Examine nodules or skin snips |
| <i>Dracunculus medienensis</i> (guinea worm) | Africa | <i>Cyclops</i> | Humans | Adults/larvae | Skin | — | May be visible in lesion |
| <i>Angiostrongylus cantonensis</i> | Southeast Asia, Pacific, Caribbean | Snails/slugs, shrimp/fish | Rats | Larvae | CSF (rarely found) | — | Eosinophilic meningitis |
| Larva Migrans Syndromes | | | | | | | |
| <i>Ancylostoma braziliense</i> (creeping eruption) | Tropical and temperate zones | Soil to skin | Dogs/cats, humans | Larvae | Skin | — | Dog and cat hookworm |
| <i>Toxocara canis</i> and <i>T. cati</i> (visceral larva migrans), <i>Baylisascaris</i> | Tropical and temperate zones | Soil, fecal-oral | Dogs/cats, raccoons, humans | Larvae | Viscera, CNS, eye | EIA | Also caused by roundworms of other species |

^aExcept for infection acquired in the South Pacific, blood should be drawn at midnight. ^bLIPS (the luciferase immunoprecipitation system) for serology and PCR (polymerase chain reaction) are available from the Laboratory of Parasitic Diseases at the National Institutes of Health: 301-496-5398.

Abbreviations: CNS, central nervous system; CSF, cerebrospinal fluid; EIA, enzyme immunoassay; RAPID, rapid immunographic assay (available internationally); Sx, signs/symptoms.

malaria, rapid detection tests (RDTs) are increasingly being used to fill this gap. These are immunochromatographic capture assays with monoclonal antibodies to species-specific antigens (histidine-rich protein 2 [PfHRP2] or aldolase of *Plasmodium falciparum*) or conserved *Plasmodium* antigens (lactate dehydrogenase). The World Health Organization sponsored a major testing program evaluating the different RDTs. Lower performance rates have been reported in a variety of field sites, especially in areas where deletions of the *pfhrp2* gene have been detected. Subpatent infection and identification of *Plasmodium* species can also be confirmed by polymerase chain reaction (PCR), but for this purpose PCR is primarily a research tool. *P. knowlesi*, a simian parasite, has been identified as the cause of an increasing number of infections in Malaysian Borneo and other areas of Southeast Asia; PCR

or another molecular method is required to differentiate *P. knowlesi* from *P. malariae*.

Although most tissue parasites stain with the traditional hematoxylin and eosin, appropriate special stains should also be applied to surgical biopsy specimens. The surgical pathologist who is accustomed to applying silver stains for *Pneumocystis* to induced sputum and transbronchial biopsies may need to be reminded to examine wet mounts and iron-hematoxylin-stained preparations of pulmonary specimens for helminthic ova and *E. histolytica*. The clinician should also be able to advise the surgeon and pathologist about optimal techniques for the identification of parasites in specimens obtained by certain specialized minor procedures (Table 245e-6). For example, the excision of skin snips for the diagnosis of onchocerciasis, the collection of rectal snips