Soil-transmitted helminth infections: ascariasis, trichuriasis, and hookworm

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The three main soil-transmitted helminth infections, ascariasis, trichuriasis, and hookworm, are common clinical disorders in man. The gastrointestinal tract of a child living in poverty in a less developed country is likely to be parasitised with at least one species of soil-transmitted helminth, with resultant impairments in physical, intellectual, and cognitive development. The benzimidazole anthelmintics, mebendazole and albendazole, are commonly used to remove these infections. The use of these drugs is not limited to treatment of symptomatic soil-transmitted helminth infections, but also for large-scale prevention of morbidity in children living in endemic areas. As a result of data showing improvements in child health and education after deworming, and the burden of disease attributed to soil-transmitted helminths, the worldwide community is awakening to the importance of these infections. Concerns about the sustainability of periodic deworming with benzimidazole anthelmintics and the emergence of resistance have prompted efforts to develop and test new control tools.

Introduction

The soil-transmitted helminths are a group of parasitic nematode worms causing human infection through contact with parasite eggs or larvae that thrive in the warm and moist soil of the world’s tropical and subtropical countries. As adult worms, the soil-transmitted helminths live for years in the human gastrointestinal tract. More than a billion people are infected with at least one species (table 1).1 Of particular worldwide importance are the roundworms (Ascaris lumbricoides), whipworms (Trichuris trichiura), and hookworms (Necator americanus or Ancylostoma duodenale). They are considered together because it is common for a single individual, especially a child living in a less developed country, to be chronically infected with all three worms. Such children have malnutrition, growth stunting, intellectual retardation, and cognitive and educational deficits.1

The soil-transmitted helminths are one of the world’s most important causes of physical and intellectual growth retardation. Yet, despite their educational, economic, and public-health importance (panel), they remain largely neglected by the medical and international community. This neglect stems from three features: first, the people most affected are the world’s most impoverished, particularly those who live on less than US$2 per day; second, the infections cause chronic ill health and have insidious clinical presentation; and third, quantification of the effect of soil-transmitted helminth infections on economic development and education is difficult. Over the past 5 years, however, the worldwide community has begun to recognise the importance of these infections after revised estimates showed that their combined disease burden might be as great as those of malaria or tuberculosis.2 Studies have also highlighted the profound effect of soil-transmitted helminth infection on school performance and attendance and future economic productivity.14 Such infections might also increase host susceptibility to other important illnesses such as malaria, tuberculosis, and HIV infection.16 In 2001, the World Health Assembly passed a resolution urging member states to control the morbidity of soil-transmitted helminth infections through large-scale use of anthelmintic drugs for school-aged children in less developed countries. A response to this resolution could establish one of the

<table>
<thead>
<tr>
<th>Disease</th>
<th>Estimated population infected (millions)</th>
<th>Geographic region</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ascaris lumbricoides</td>
<td>Common roundworm infection</td>
<td>807–1221</td>
</tr>
<tr>
<td>Trichuris trichiura</td>
<td>Whipworm infection</td>
<td>604–795</td>
</tr>
<tr>
<td>Necator americanus and Ancylostoma duodenale</td>
<td>Hookworm infection</td>
<td>576–740</td>
</tr>
<tr>
<td>Strongyloides stercoralis</td>
<td>Threadworm infection</td>
<td>30–100</td>
</tr>
<tr>
<td>Enterobius vermicularis</td>
<td>Pinworm infection</td>
<td>4–28% of children</td>
</tr>
<tr>
<td>Toxocara canis and Toxocara cati</td>
<td>Visceral and ocular larva migrans</td>
<td>2–80% of children</td>
</tr>
</tbody>
</table>

Pathogens of minor or local importance

<table>
<thead>
<tr>
<th>Disease</th>
<th>Geographic region</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ancylostoma braziliense</td>
<td>Cutaneous larva migrans</td>
</tr>
<tr>
<td>Uncinaria stenocephala</td>
<td>Cutaneous larva migrans</td>
</tr>
<tr>
<td>Ancylostoma caninum</td>
<td>Eosinophilic enteritis</td>
</tr>
<tr>
<td>Ancylostoma ceylanicum</td>
<td>Hookworm infection</td>
</tr>
<tr>
<td>Baylisascaris procyonis</td>
<td>Eosinophilic meningitis</td>
</tr>
<tr>
<td>Oesophagostomum bifurcum</td>
<td>Nodular worm infection</td>
</tr>
<tr>
<td>Strongyloides fuelleborni</td>
<td>Swollen belly syndrome</td>
</tr>
<tr>
<td>Ternidens diminutus</td>
<td>False hookworm infection</td>
</tr>
</tbody>
</table>

Table 1: Soil-transmitted helminth infections of human beings
largest worldwide health initiatives ever undertaken. However, such widespread and frequent use of anthelmintics could lead to drug resistance or at least a decline in effectiveness of these front-line drugs in the long-term battle with soil-transmitted helminths.

The parasites
Adult hookworms of the genera *Necator* and *Ancylostoma* parasitise the upper part of the human small intestine, whereas ascaris roundworms parasitise the entire small intestine and adult trichuris whipworms live in the large intestine, especially the caecum (table 2). The parasites can live for several years in the human gastrointestinal tract. Human beings are regarded as the only major definitive host for these parasites, although in some cases ascaris infections can also be acquired from pigs. The soil-transmitted helminths vary greatly in size, and female worms are larger than males (figure 1). After mating, each adult female produces thousands of eggs per day (figure 2), which leave the body in the faeces. People become infected with *Trichuris trichiura* and *Ascaris lumbricoides* by ingesting the fully developed eggs. After ingestion of trichuris eggs, the released larvae moult and travel to the colon where they burrow into the epithelia and develop into adult whipworms within about 12 weeks. Ascaris larvae penetrate the intestinal mucosa and after an obligatory extraintestinal migration, they enter the liver then the lungs, before passing over the epiglottis to re-enter the gastrointestinal tract and develop into egg-laying adult worms about 9–11 weeks after egg ingestion.

*N americanus* and *A duodenale* hookworm eggs hatch in soil. The larvae moult twice to become infective third-stage larvae, which are non-feeding but motile organisms that seek out higher ground to improve the chance of contact with human skin. After skin penetration, they enter subcutaneous venules and lymphatic vessels to access the host’s afferent circulation. Ultimately, the larvae become trapped in pulmonary capillaries, enter the lungs, pass over the epiglottis, and migrate into the gastrointestinal tract. About 5–9 weeks are needed from skin penetration until development of egg-laying adults. *A duodenale* larvae are also orally infective, and lactogenic transmission during breastfeeding has been postulated. Soil-transmitted helminths do not reproduce within the host. This feature is crucial for understanding of the epidemiology and clinical features of soil-transmitted helminth infections, as well as the approaches to their control.

**Epidemiology and burden of disease**
Soil-transmitted helminth infections are widely distributed throughout the tropics and subtropics (table 3). Climate is an important determinant of transmission of these infections, with adequate moisture and warm temperature required for egg development and larval development and migration.

<table>
<thead>
<tr>
<th>Species</th>
<th>Length (mm)</th>
<th>Daily egg output per female worm</th>
<th>Location in host</th>
<th>Lifespan (years)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Large common roundworm</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ascaris lumbricoides</td>
<td>150–400</td>
<td>200 000</td>
<td>Small intestine</td>
<td>1</td>
</tr>
<tr>
<td>Whipworm</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Trichuris trichiura</td>
<td>30–50</td>
<td>3000–5000</td>
<td>Caecum and colon</td>
<td>1·5–2·0</td>
</tr>
<tr>
<td>Hookworm</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Necator americanus</td>
<td>7–13</td>
<td>9000–10 000</td>
<td>Upper small intestine</td>
<td>5–7</td>
</tr>
<tr>
<td>Ancylostoma duodenale</td>
<td>8–13</td>
<td>25 000–30 000</td>
<td>Upper small intestine</td>
<td>5–7</td>
</tr>
</tbody>
</table>

Table 2: Characteristics of the soil-transmitted helminths: adult worms of greatest public-health significance
essential for larval development in the soil. Equally important determinants are poverty and inadequate water supplies and sanitation. In such conditions, soil-transmitted helminth species are commonly coendemic. There is evidence that individuals with many helminth infections have even heavier infections with soil-transmitted helminths. Because morbidity from these infections and the rate of transmission are directly related to the number of worms harboured in the host, intensity of infection is the main epidemiological index used to describe soil-transmitted helminth infection. Intensity of infection is measured by the number of eggs per gram of faeces, generally by the Kato-Katz faecal thick-smear technique. For *A lumbricoides* and *T trichiura*, the most intense infections are in children aged 5–15 years, with a decline in intensity and frequency in adulthood. Whether such age dependency indicates changes in exposure, acquired immunity, or a combination of both remains controversial. Although heavy hookworm infections also occur in childhood, frequency and intensity commonly remain high in adulthood, even in elderly people. Soil-transmitted helminth infections are often referred to as being “overdispersed” in endemic communities, such that most worms are harboured by a few individuals in an endemic area. There is also evidence of familial and household aggregation of infection, with the relative contribution of genetics and common household environment debated.

Estimates of annual deaths from soil-transmitted helminth infection vary widely, from 12 000 to as many as 135 000. Because these infections cause more disability than death, the worldwide burden, as for many neglected tropical diseases, is typically assessed by disability-adjusted life years (DALY). Since the first DALY estimates were provided, there has been much variability in quoted estimates (table 4), partly because of different emphases on the cognitive and health effects. The lower estimates assume that most hookworm cases do not result in severe anaemia or pronounced protein loss by the host, whereas the higher estimates show the long-term results of infection such as malnutrition and delayed cognitive development, especially in children. For these reasons, school-aged children have been the major targets for anthelmintic treatment, and the scale of disease in this age group was pivotal in leveraging support for school-based control.

<table>
<thead>
<tr>
<th>Infection</th>
<th>1990</th>
<th>1990</th>
<th>2001</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ascariasis</td>
<td>1.8</td>
<td>10.5</td>
<td>1.2</td>
</tr>
<tr>
<td>Trichuriasis</td>
<td>1.8</td>
<td>6.4</td>
<td>1.6</td>
</tr>
<tr>
<td>Hookworm</td>
<td>1.5</td>
<td>22.1</td>
<td>1.8</td>
</tr>
<tr>
<td>Total</td>
<td>5.0</td>
<td>39.0</td>
<td>4.7</td>
</tr>
</tbody>
</table>

Table 4: Estimates of DALY lost to soil-transmitted helminth infections.


Table 3: Worldwide estimates of number of soil-transmitted helminth infections by region (millions of cases).

LAC=Latin America and Caribbean; SSA=sub-Saharan Africa; MENA=middle-east and north Africa; SAS=south Asia; EAP=East Asia and the Pacific Islands.
There is evidence to support the high disease-burden estimates from soil-transmitted helminth infections, and highlight the importance of hookworm as a threat to maternal and child health. For example, cross-sectional evidence from Africa and Asia shows that 30–54% of moderate to severe anaemia in pregnant women is attributable to hookworm.\textsuperscript{11,13} and intervention studies suggest that antenatal anthelmintics substantially increase maternal haemoglobin concentrations as well as birthweight and infant survival.\textsuperscript{14} In childhood, hookworm contributes to moderate and severe anaemia in school-aged children,\textsuperscript{19} and there is increasing recognition of a similar contribution in preschool children.\textsuperscript{20,21} These features of hookworm disease need to be better incorporated into DALY estimates. Because hookworms are the most widespread species of soil-transmitted helminth in sub-Saharan Africa,\textsuperscript{6} where iron stores are low, this consequence of infection could substantially alter the perception of the public-health importance of hookworm. In light of their nutritional and educational effects, soil-transmitted helminth infections clearly need to be reassessed, as has lately been done for schistosomiasis.\textsuperscript{38}

**Host-parasite interactions**

Despite their large size and ability to elicit potent immune responses, soil-transmitted helminths are refractory to host immunity, establishing chronic infections during the host’s life, and, in the case of hookworm, intensity of infection actually rises with the age of the host.\textsuperscript{22} These organisms have complex life cycles within the human host, undergoing a succession of developmental stages, which can carry stage-specific antigens, and pass through a range of host tissues (skin, lungs, and gut).\textsuperscript{36} Soil-transmitted helminths are thought to survive within the host not just by warding off immune attack, but instead by aggressively subverting the host immune response to create niches that optimise successful residence, feeding, and reproduction.\textsuperscript{40} Soil-transmitted helminths induce production of cytokines (interleukin-4, interleukin-5, interleukin-10, and interleukin-13), parasite-specific immunoglobulin, and non-specific immunoglobulin E, and expansion and mobilisation of mast cells, eosinophils, and basophils.\textsuperscript{40} This constellation of responses is known as the T-helper-2 (Th2) immune response. It is important in allergy and clinical immunology in general.\textsuperscript{40} Whether the Th2 response brings about the elimination or the maintenance of the parasite is debated. The functional effector mechanisms driven by the Th2 response to infection with soil-transmitted helminths include eosinophil-mediated larval killing, production of specific and polyclonal immunoglobulin E, mast-cell degranulation, goblet-cell hyperplasia, and increased mucus secretion.\textsuperscript{40} Different subsets of effector cells might operate against different nematode species;\textsuperscript{41} for example, mast cells seem to be central to protective responses against hookworm and ascars but not in the expulsion of trichuris.\textsuperscript{42} Although immunity to hookworm at the population level is not apparent, a negative association between concentrations of interleukin-5 and the likelihood of being reinfected with \textit{N americanus} after anthelmintic treatment has been found, suggesting that the effect of interleukin-5 (probably mediated by eosinophils) is directed against incoming larvae.\textsuperscript{43,44} Similarly, inverse associations between secretion of interleukin-5 and interleukin-13 and susceptibility to reinfection were noted in patients infected with \textit{A lumbricoides} or \textit{T trichiura} infections.\textsuperscript{45,46}

The survival of soil-transmitted helminths suggests that they succeed by achieving some form of balanced parasitism, in which transmission is maintained and acute morbidity avoided. This ideal homeostatic state almost certainly needs an environment rich in regulatory mechanisms. Interleukin-10 is the most abundantly produced regulatory cytokine in soil-transmitted helminth infection. However, its role in maintaining the chronicity of soil-transmitted helminth infection is unclear.\textsuperscript{47} Geiger and colleagues,\textsuperscript{47} reported that interleukin-10 responses to crude ascars antigen were high in individuals infected with ascars or trichuris. Whereas Turner and co-workers\textsuperscript{48} reported that interleukin-10 concentrations declined with intensity of ascars infection in older individuals, Jackson and colleagues\textsuperscript{49} showed that higher interleukin-10 concentrations correlated with heavier ascars infection in older people. These downregulatory immune mechanisms might also benefit the host by blocking progression to atopic reactions.\textsuperscript{50} The immune response to these infections has long been known to share key features with the allergic response, especially the enhanced Th2 response. In view of these immunological features and the complementary geographic distribution of soil-transmitted helminth infection and allergic disease, many studies have investigated whether the Th2 response to soil-transmitted helminths protects or pre-empts the host from developing allergic manifestations linked to Th2, a theory known as the hygiene hypothesis.\textsuperscript{51}

Much of the survival success of soil-transmitted helminths can be attributed to their secretomes, which interact with host tissues and maintain the parasitic existence (table 5). Of particular importance are the secretions that modulate the host’s immune response. As helminth proteomes are being matched to increasing gene sequence datasets,\textsuperscript{74,75} a molecular snapshot of the mixture released into host tissues by these parasites is being gradually revealed. One constituent, natural-killer-cell-binding protein, is secreted by adult \textit{N americanus} and binds specifically to natural-killer cells and induces them to secrete interferon-y.\textsuperscript{52} This finding is the first evidence of a pathogen-derived protein that binds selectively to natural-killer cells and the first report of a nematode-derived product that induces abundant secretion of cytokines from natural-killer cells. The researchers suggested that interferon-y production in the gut would counteract the development of a potentially host-protective Th2 response that might eliminate the parasite.\textsuperscript{52} Other secreted proteins from adult hookworms
modulate immune responses. The dog hookworm, *A caninum*, secretes neutrophil inhibitory factor, which binds to the integrins CD11b/CD18 and blocks adhesion of activated human neutrophils to vascular endothelial cells as well as the release of hydrogen peroxide from activated neutrophils.\(^6\) This protein is in the pathogenesis-related protein superfamily, cysteine-rich secreted proteins that are abundantly expressed by all parasitic nematodes investigated so far. They seem to have diverse roles in nematode parasitism by binding to host cells. Other hookworm pathogenesis-related proteins combat haemostasis by binding to platelets and inhibiting their activation.\(^6\) The observations that these proteins are released by third-stage larvae after stimulation with human serum suggests their importance in the mechanisms with potential as anti-helminth vaccines or therapies for other disorders (experimentally proven or suggested by the cited authors).

<table>
<thead>
<tr>
<th>Species</th>
<th>Molecule</th>
<th>Known or putative function</th>
<th>Therapeutic potential</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hookworms</td>
<td>ASP2</td>
<td>Pathogenesis-related protein of unknown function but secreted on host entry by third-stage larvae</td>
<td>Hookworm vaccine antigen</td>
<td>52-54</td>
</tr>
<tr>
<td></td>
<td>NIF</td>
<td>Binds CD11b/CD18 and blocks neutrophil migration</td>
<td>Treatment for cerebral ischemia</td>
<td>55-56</td>
</tr>
<tr>
<td></td>
<td>NKBP</td>
<td>Binds natural killer cells and induces interferon-γ production</td>
<td>Potential adjuvant</td>
<td>57</td>
</tr>
<tr>
<td></td>
<td>Haemoglobinases</td>
<td>Cascade of mechanically distinct proteases that digest haemoglobin in the worm’s gut</td>
<td>Hookworm vaccine antigens</td>
<td>58-60</td>
</tr>
<tr>
<td></td>
<td>CPI</td>
<td>Pathogenesis-related protein that inhibits platelet activation and adhesion by blocking function of gpIIb/IIIa and gpIa/IIa</td>
<td>Potential hookworm vaccine candidate</td>
<td>61,62</td>
</tr>
<tr>
<td></td>
<td>AcAPs</td>
<td>Novel and potent anticoagulant that inhibit factor Xa, factor VIIa, and tissue factor VIIa/TF</td>
<td>Thrombosis and disseminated intravascular coagulation</td>
<td>63-65</td>
</tr>
<tr>
<td></td>
<td>Eotaxin-cleaving protease</td>
<td>Secreted metalloprotease that digests eotaxin and prevents eosinophil recruitment</td>
<td></td>
<td>66</td>
</tr>
<tr>
<td></td>
<td>Haemolysin</td>
<td>Haemolytic protein that forms pores in erythrocyte membranes allowing hemoglobin to be released</td>
<td>Potential hookworm vaccine candidate</td>
<td>67</td>
</tr>
<tr>
<td>Ascaris</td>
<td>PI 3</td>
<td>Pepsin inhibitor that protects worms from digestion</td>
<td></td>
<td>68</td>
</tr>
<tr>
<td></td>
<td>PC</td>
<td>Phosphorylcholine linked to secreted glycoconjugates suppress lymphocyte proliferation</td>
<td></td>
<td>69</td>
</tr>
<tr>
<td>Trichuris</td>
<td>TT47</td>
<td>Forms pores in caecal epithelial cells, allowing parasite to keep anterior end in syncytilal environment</td>
<td></td>
<td>70</td>
</tr>
<tr>
<td></td>
<td>ES products</td>
<td>Promote Th2/Treg response that dampens intestinal inflammation</td>
<td>Therapy for Crohn’s disease and ulcerative colitis</td>
<td>71,72</td>
</tr>
<tr>
<td></td>
<td>TsMIF</td>
<td>Inhibits migration of PBMCs by competing with host macrophage inhibitory factor</td>
<td></td>
<td>73</td>
</tr>
</tbody>
</table>

Table 5: Selected molecules secreted by soil-transmitted helminths, their known or putative functions and their potentials as anti-helminth vaccines or therapies for other disorders (experimentally proven or suggested by the cited authors)
the suppression of lymphocyte responses in ascarisis and filarial nematode infections. Moreover, secreted ascaris glycosphingolipids inhibit lipopolysaccharide-induced production of Th1 cytokines such as interferon-γ in a phosphorylcholine-dependent manner, further highlighting the diverse molecular interactions of the immunmodulatory secretory products of the soil-transmitted helminths.

T trichiura secretes large amounts of a protein called TT47 that forms ion-conducting pores in lipid bilayers, allowing the parasite to invade the host gut and maintain its anterior end in a syncytial environment in the caecal epithelium. Unlike T trichiura, the swine whipworm T suis does not develop to maturity in people, although the larvae can briefly colonise individuals without causing disease. The secreted products of trichuris are potent inducers of anti-inflammatory cytokines. This attribute has led to the use of T suis to treat proinflammatory autoimmune disorders such as Crohn’s disease, in which helminth larvae are thought to create an anti-inflammatory environment in the gut that combats the proinflammatory (Th1-biased) immune response associated with this disease. The specific secreted molecules in T suis that induce the anti-inflammatory response are unknown, although potential candidates include one that mimics the effects of the human chemokine, macrophage migration inhibitory factor.

Clinical features
The clinical features of soil-transmitted helminth infections can be classified into the acute manifestations associated with larval migration through the skin and viscerae, and the acute and chronic manifestations resulting from parasitism of the gastrointestinal tract by adult worms (table 6).

Early larval migration
Migrating soil-transmitted helminth larvae provoke reactions in many of the tissues through which they pass. For example, ascaris larvae that die during migration through the liver can induce eosinophilic granulomas. In the lungs, ascaris larval antigens cause an intense inflammatory response consisting of eosinophilic infiltrates that can be seen on chest radiographs. The resulting verminous pneumonia is commonly accompanied by wheezing, dyspnoea, a non-productive cough, and fever, with blood-tinged sputum produced during heavy infections. Children are more susceptible to pneumonitis, and the disease is more severe on reinfection. In some regions—such as Saudi Arabia—verminous pneumonia is seasonal and occurs after spring rains. Small numbers of affected children develop status asthmaticus, leading to the idea that A lumbricoides and its zoonotic counterpart, Toxocara canis, are occult environmental causes of asthma.

Several cutaneous syndromes result from skin-penetrating larvae. Repeated exposure to N americanus and A duodenale hookworm third-stage larvae results in ground itch, a local erythematous and papular rash accompanied by pruritus on the hands and feet. By contrast, when zoonotic hookworm third-stage larvae—typically A braziliense—enter the skin, they produce cutaneous larva migrans, which is characterised by the appearance of serpiginous tracks on the feet, buttocks, and abdomen. After skin invasion, hookworm third-stage larvae travel through the vasculature and enter the lungs, although the resulting pneumonitis is not as great as in ascaris infection. Oral ingestion of A duodenale larvae can result in Wakana syndrome, which is characterised by nausea, vomiting, pharyngeal irritation, cough, dyspnoea, and hoarseness.

Intestinal parasitism
Generally only soil-transmitted helminth infections of moderate and high intensity in the gastrointestinal tract produce clinical manifestations, with the highest-intensity infections most common in children. The numerical threshold at which worms cause disease in children has not been established, because it depends on the underlying nutritional status of the host. Each of the major soil-transmitted helminths produces characteristic disease syndromes.

Ascarasis
The presence of large numbers of adult ascaris worms in the small intestine can cause abdominal distension and pain (figure 3). They can also cause lactose intolerance and malabsorption of vitamin A and possibly other nutrients, which might partly cause the nutritional and growth failure. In young children, adult worms can aggregate in the ileum and cause partial obstruction because the lumen is small. Various grave consequences can ensue, including intussusception, volvulus, and complete obstruction, leading to bowel infarction and intestinal perforation. The resulting peritonitis can be fatal, although if the child survives, the wandering adult worms can die and cause a chronic
granulomatous peritonitis. Typically, a child with obstruction because of ascaris has a toxic appearance with signs and symptoms of peritonitis. In some cases, a mass can be felt in the right lower quadrant. Adult worms can enter the lumen of the appendix, leading to acute appendicular colic and gangrene of the appendix tip, resulting in a clinical picture indistinguishable from appendicitis. Adult ascaris worms also tend to move in children with high fever, resulting in the emergence of worms from the nasopharynx or anus. Hepatobiliary and pancreatic ascariasis results when adult worms in the duodenum enter and block the ampullary orifice of the common bile duct, leading to biliary colic, choledysitis, cholangitis, pancreatitis, and hepatic abscess. By contrast with intestinal obstruction, hepatobiliary and pancreatic ascariasis occurs more commonly in adults—especially women—than in children, presumably because the adult biliary tree is large enough to accommodate an adult worm.

**Trichuriasis**

Adult whipworms live preferentially in the caecum, although in heavy infections, whipworms can be seen throughout the colon and rectum. The adult parasite leads both an intracellular and an extracellular existence, with the anterior end embedded in epithelial tunnels within the intestinal mucosa and the posterior end located in the lumen. Inflammation at the site of attachment from large numbers of whipworms results in colitis. Longstanding colitis produces a clinical disorder that resembles inflammatory bowel disease, including chronic abdominal pain and diarrhoea, as well as the sequelae of impaired growth, anaemia of chronic disease, and finger clubbing. Trichuris dysentery syndrome is an even more serious manifestation of heavy whipworm infection, resulting in chronic dysentery and rectal prolapse. Whipworm infection can also exacerbate colitis caused by infection with *Campylobacter jejuni*.

**Hookworm infection**

In hookworm infection, the appearance of eosinophilia coincides with the development of adult hookworms in the intestine. The major pathology of hookworm infection, however, results from intestinal blood loss as a result of adult parasite invasion and attachment to the mucosa and submucosa of the small intestine. Hookworm disease occurs when the blood loss exceeds the nutritional reserves of the host, thus resulting in iron-deficiency anaemia. The presence of more than 40 adult worms in the small intestine is estimated to be sufficient to reduce host haemoglobin concentrations below 11 g/dL, although the exact number depends on several factors including the species of hookworm—*A duodenale* causes more blood loss than *N americanus*—and the host iron reserves. The clinical manifestations of hookworm disease resemble those of iron-deficiency anaemia from other causes. The chronic protein loss from heavy hookworm infection can result in hypoproteinaemia and oedema. Because children and women of reproductive age have reduced iron reserves, they are at particular risk of hookworm disease. The severe iron-deficiency anaemia that can arise from hookworm disease during pregnancy can have adverse results for the mother, the fetus, and the neonate.

**Diagnosis and treatment**

In their definitive host, each adult female whipworm or hookworm produces thousands of eggs per day, and each female ascaris worm produces upwards of 200 000 eggs daily (table 2). Because many soil-transmitted helminth infections present without specific signs and symptoms, the clinician typically needs some index of suspicion, such as local epidemiology or country of origin, to request a faecal examination. In some cases, especially of hookworm infection, persistent eosinophilia is a common presenting finding. Several egg concentration techniques—eg, formalin-ethyl acetate sedimentation—can detect even light infections. The Kato-Katz faecal-thick smear and the McMaster method are used to measure the intensity of infection by estimating the number of egg counts per gram of faeces. Ultrasonography and endoscopy are useful for diagnostic imaging of the complications of ascariasis, including intestinal obstruction and hepatobiliary and pancreatic involvement.

The treatment goal for soil-transmitted helminth infections is to remove adult worms from the gastrointestinal tract (table 7). The drugs most commonly used for the removal of soil-transmitted helminth infections are mebendazole and albendazole. These benzimidazole drugs bind to nematode β-tubulin and inhibit parasite microtubule polymerisation, which causes death of adult worms through a process that can take several days. Although both albendazole and mebendazole are deemed broad-spectrum anthelmintic agents, important therapeutic differences affect their use in clinical practice. Both agents are effective against ascaris in a single dose. However, in hookworm, a single
Table 7: Treatment of soil-transmitted helminth infections

<table>
<thead>
<tr>
<th>Infection</th>
<th>Drug</th>
<th>Dose</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ascarisis</td>
<td>Albendazole†</td>
<td>400 mg once</td>
</tr>
<tr>
<td></td>
<td></td>
<td>400 mg once</td>
</tr>
<tr>
<td></td>
<td>Mebendazole</td>
<td>100 mg twice a day for 3 days</td>
</tr>
<tr>
<td></td>
<td></td>
<td>100 mg twice a day for 3 days</td>
</tr>
<tr>
<td></td>
<td>Pyrantel pamoate</td>
<td>11 mg/kg (maximum dose 1 g) for 3 days</td>
</tr>
<tr>
<td></td>
<td></td>
<td>11 mg/kg (maximum dose 1 g) for 3 days</td>
</tr>
<tr>
<td>Hookworm</td>
<td>Albendazole⁷</td>
<td>400 mg once</td>
</tr>
<tr>
<td></td>
<td></td>
<td>400 mg once</td>
</tr>
<tr>
<td></td>
<td>Mebendazole</td>
<td>100 mg twice a day for 3 days</td>
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</tr>
<tr>
<td></td>
<td></td>
<td>11 mg/kg (maximum dose 1 g) for 3 days</td>
</tr>
<tr>
<td>Trichuriasis</td>
<td>Mebendazole</td>
<td>100 mg twice a day for 3 days</td>
</tr>
<tr>
<td></td>
<td></td>
<td>100 mg twice a day for 3 days</td>
</tr>
<tr>
<td></td>
<td>Levamisole</td>
<td>2.5 mg/kg once</td>
</tr>
<tr>
<td></td>
<td></td>
<td>2.5 mg/kg once</td>
</tr>
<tr>
<td></td>
<td>Pyrantel pamoate</td>
<td>11 mg/kg (maximum dose 1 g) for 3 days</td>
</tr>
<tr>
<td></td>
<td></td>
<td>11 mg/kg (maximum dose 1 g) for 3 days</td>
</tr>
<tr>
<td></td>
<td>Levamisole</td>
<td>2.5 mg/kg; repeat after 7 days in heavy infection</td>
</tr>
<tr>
<td></td>
<td></td>
<td>2.5 mg/kg; repeat after 7 days in heavy infection</td>
</tr>
<tr>
<td></td>
<td>Albendazole⁷</td>
<td>400 mg once</td>
</tr>
<tr>
<td></td>
<td></td>
<td>400 mg once</td>
</tr>
<tr>
<td></td>
<td>Mebendazole</td>
<td>100 mg once</td>
</tr>
<tr>
<td></td>
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<td></td>
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<td>2.5 mg/kg once</td>
</tr>
<tr>
<td></td>
<td>Albendazole⁷</td>
<td>400 mg for 3 days</td>
</tr>
<tr>
<td></td>
<td></td>
<td>400 mg for 3 days</td>
</tr>
</tbody>
</table>

“Modified from the Medical Letter on Drugs and Therapeutics, Drugs for Parasitic Infections.” For children of 1–2 years the dose of albendazole is 200 mg instead of 400 mg, based on a recommendation in the Report of the WHO informal consultation on the use of praziquantel during pregnancy and lactation and albendazole/mebendazole in children under 24 months.

A dose of mebendazole has a low cure rate and albendazole is more effective. Conversely, a single dose of albendazole is not effective in many cases of trichuriasis. For both trichuriasis and hookworm infection, several doses of benzimidazole anthelmintic drugs are commonly needed. Another important difference between the two drugs is that mebendazole is poorly absorbed from the gastrointestinal tract so its therapeutic activity is largely confined to adult worms. Albendazole is better absorbed, especially when ingested with fatty meals, and the drug is metabolised in the liver to a sulphoxide derivative, which has a high volume of distribution in the tissues. For this reason, albendazole is used for the treatment of disorders caused by tissue-migrating larvae such as visceral larva migrans caused by Toxocara canis. Systemic toxic effects, such as those on the liver and bone marrow, are rare for the benzimidazole anthelmintic drugs in the doses used to treat soil-transmitted helminth infections. However, transient abdominal pain, diarrhoea, nausea, dizziness, and headache commonly occur.

Because the benzimidazole anthelmintic drugs are embryotoxic and teratogenic in pregnant rats, there are concerns about their use in children younger than 12 months and during pregnancy. Overall, the experience with these drugs in children younger than 6 years is scarce, although evidence suggests they are probably safe. A review of the use of the benzimidazole anthelmintic drugs in children aged 12–24 months concluded that they can be used “if local circumstances show that relief from ascariasis and trichuriasis is justified.” Both pyrantel pamoate and levamisole are regarded as alternative drugs for the treatment of hookworm and ascaris infections, although the former is not effective for the treatment of trichuriasis and they are administered by bodyweight.

Morbidity control through deworming

The use of anthelmintic drugs nowadays is not restricted to the treatment of symptomatic soil-transmitted helminth infections; the drugs are now used also for large-scale morbidity reduction in endemic communities. Increasing evidence suggests that chronic infection with soil-transmitted helminths results in impaired childhood growth and poor physical fitness and nutritional status. The causal link between chronic infection and impaired childhood development is extrapolated from the recorded improvement in these features after deworming. The mechanisms underlying these associations are thought to involve impairment of nutrition, although there is little specific evidence to support this assumption.

Regular treatment with benzimidazole anthelmintic drugs in school-age children reduces and maintains the worm burden below the threshold associated with disease. The benefits of regular deworming in this age group include improvements in iron stores, growth and physical fitness, cognitive performance, and school attendance. In younger children, studies have shown improved nutritional indicators such as reduced wasting, malnutrition, and stunting, and improved appetite. Although some investigators still find this relation controversial. Relevant to these findings, administration of anthelmintic drugs to children infected with soil-transmitted helminths from 1 year of age is now deemed appropriate. The patents on anthelmintic drugs recommended by WHO have expired, and the drugs can be produced at low cost by generic manufacturers. The cost of drug delivery is also low because after simple training, teachers could be involved in deworming. If children in endemic areas are treated once or twice during pregnancy, there are substantial improvements in maternal anaemia and birthweight and infant mortality at 6 months. In areas where hookworm infections are endemic, anthelmintic treatment is recommended during pregnancy except in the first trimester.

An important factor in treatment is reinfection. After community-wide treatment, rates of hookworm infection reach 80% of pretreatment rates within 30–36 months. A L. loa infection reached 55% of pretreatment rates within 11 months and T. trichiura infection reached 44% of pretreatment rates within 17 months. Despite reinfection, however, regular treatment to reduce the worm burden consistently could prevent some of the sequelae associated with chronic infection.

Drug resistance against the front-line anthelmintics is widespread in nematodes of livestock as a result of frequent treatment of animals kept in close proximity and with little gene flow. If such conditions were replicated in human nematodes, drug resistance would
soon arise.124 Human nematodes have longer reproducing times, are subjected to less frequent treatment (the treatment interval is longer than the parasites’ generation time), and the treatment is targeted at certain populations, thereby sparing a circulating pool of sensitive alleles, which should reduce selection pressure.125 Nevertheless, the effectiveness of drugs must be closely monitored, especially in areas where drug pressure is high, such as regions where mass anthelmintic chemotherapy is also administered for the elimination of lymphatic filariasis. Development of sensitive methods for the early detection of anthelmintic resistance are part of the research agenda, with special attention being given to in-vitro tests and molecular biology techniques that could be adaptable to field conditions.9 Because no new anthelmintic drugs are in late-stage development at present, the effectiveness of available products needs to be preserved.

New control methods
Concerns about the sustainability of periodic deworming with benzimidazole anthelmintic drugs and the emergence of resistance with widespread use have prompted efforts to develop and test new control tools. Nitazoxanide, a nitroimidazole compound that is increasingly used in children with giardiasis and cryptosporidiosis, is also being explored as a broad-spectrum antiparasitic agent with anthelmintic properties.126 Tribendimidine has low toxicity, yet broad-spectrum activity against many soil-transmitted helminths.127 In randomised studies in China, tribendimidine was equivalent to mebendazole and albendazole for the treatment of A lumbricoides, T trichiura, and hookworm infections, and better than these drugs for N americanus infection.128 A study comparing tribendimidine with albendazole for the treatment of hookworm is under way in Africa. Combination therapy with drugs with differing modes of action is an alternative strategy to improve efficacy and lower the risk of resistance.9 For example, combinations of levamisole with mebendazole and of pyrantel with oxantel are more effective than any single drug.129

Vaccination remains the method of choice to control soil-transmitted helminth infection, because it offers the possibility of a simple, single step for the interruption of infection, disease, and transmission. Several substantial obstacles impede vaccine development against soil-transmitted helminths,19 including the lack of good animal models and a poor understanding of the events that permit soil-transmitted helminths to endure for years in their human host in the face of a potent immune response. Nevertheless, a hookworm vaccine consisting of the recombinant larval antigen ASP2 is effective in animal models (dogs and hamsters) and has shown a protective association in immunoepidemiology studies in two continents.30,32,33,130 The Na ASP-2 hookworm vaccine is now undergoing clinical development in human beings.131

Conclusions
Soil-transmitted helminth infections in people will remain a worldwide public-health threat for as long as poverty persists in the developing world. The UN agencies have appropriately recognised the health and educational effect of these infections in children, and have taken steps to distribute anthelmintic drugs in schools and to undertake chemotherapy programmes on an unprecedented scale. Large-scale deworming is necessary to reduce the worldwide morbidity of these infections, but without improved water supplies and sanitation this approach cannot be relied on for sustainable reductions in parasite frequency or intensity of infection. The infrastructure that has been established for deworming of children in schools is expected, however, to facilitate introduction of new anthelmintic vaccines and other control tools,131 and some of the proposed interventions for the integrated control of endemic neglected tropical coinfections such as lymphatic filariasis, onchocerciasis, schistosomiasis, and trachoma.132 Such strategies could result in substantial reductions in the worldwide disease burden in the years to come.

Conflict of interest statement
P. Hotez is an inventor on an international patent application: PCT/US02/33106 (filed Nov 11, 2002) “Hookworm vaccine”. The patent was filed in the USA, Brazil, India, China, and Mexico. If awarded, the patent would belong to the George Washington University with an exclusive licence to the Human Hookworm Vaccine Initiative of the Albert B Sabin Vaccine Institute, a non-profit (501c3) organisation devoted to increasing the use of vaccines worldwide. The Human Hookworm Vaccine Initiative is funded mainly by the Bill and Melinda Gates Foundation. Because hookworm is a neglected disease affecting the poorest people in less developed countries, a hookworm vaccine is not expected to have commercial value or income-generating potential. The rationale for filing a patent is to ensure that the vaccine is developed for those who need it and to encourage vaccine manufacturers in less developed countries to work with the Sabin Vaccine Institute for manufacture of the hookworm vaccine. The first-generation hookworm vaccine, the Na-ASP-2 vaccine was developed entirely in the non-profit sector through the Human Hookworm Vaccine Initiative of the Albert B Sabin Vaccine Institute. P Hotez is a co-chair of the Scientific Advisory Council of the Albert B Sabin Vaccine Institute, but he receives no compensation for this activity. He is also a member of the academic advisory board for the Pfizer Postdoctoral Fellowship in Infectious Diseases. This provides a postdoctoral fellowship to a highly-qualified infectious diseases specialist conducting basic and translational research in infectious diseases at an academic medical centre. This activity is unrelated to anything discussed in this Seminar. J Bethony is the recipient of an International Research Scientist Award (RO1) from Fogarty International Center of the National Institutes of Health. S Brooker is the recipient of a Wellcome Trust Advanced Research Fellowship (073656). A. Loukas is the recipient of an RD Wright Career Development Award from the National Health and Medical Research Council of Australia. M Albonico is supported by the Fondazione Ivo de Carneri. All the authors receive funding from the Bill and Melinda Gates Foundation through the Human Hookworm Vaccine Initiative of the Albert B Sabin Vaccine Institute. No funding source had any role in the writing of this Seminar.

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immunosuppressed patients and very young children.

On the basis of our findings, M A Phadke and N A Kshirsagar consider the tuberculin skin test, although less specific, more sensitive and suitable for resource-limited settings than the new interferon-gamma-based assays. Both types of test identify individuals with latent tuberculosis infection; however, only a few people latently infected also have (or will have in the future) active disease. Thus figure 1 of our paper does not show that the blood tests are less sensitive, but indicates that the skin test is less specific for diagnosing latent tuberculosis.

Furthermore, table 2 suggests, albeit in few cases, that the new blood tests might in fact be more sensitive than the skin test in patients with active disease. Poor sensitivity of the skin test in immunosuppressed patients is a well-known limitation of this century-old diagnostic tool; however, an ELISPOT-based blood assay was more sensitive than the skin test for diagnosing tuberculosis in malnourished, HIV-positive children in a high-prevalence setting. Although these new diagnostics hold great promise, further studies are clearly needed, in particular in high-risk groups.

Beate Kampmann and co-workers’ data confirm those reported by Connell and co-workers, who showed that up to 70% of children with a diagnosis of latent tuberculosis infection based on the results of the tuberculin skin test would have been misdiagnosed had they been tested with a whole-blood interferon gamma assay. Nonetheless, an ELISPOT-based assay provided valid results in all neonates tested in a contact-tracing setting, and a new in-tube format of the whole-blood interferon gamma assay gave determinate results in all children admitted to hospital in rural India.

The need for more accurate diagnostic tools in children, in particular for those living in endemic areas, is crucial for the global control of tuberculosis. For this reason, we should not miss the unique opportunity provided by the new interferon-gamma-based blood tests. However, we agree that validating studies in highly vulnerable groups, such as young children, are urgently needed before advocating the universal use of one particular blood test.

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Urban and colleagues’ excellent Seminar on ground helminths (May 6, p 1521), I feel several points should be addressed.

Although it is true that the greatest burden of these pathogens is felt in developing countries, the average practitioner in developed countries is most likely to diagnose ground helminth infections in a specific population: returning travellers. Here, the acute manifestations resulting from helminthic migration discussed briefly by Bethony and colleagues are the main concern. Fever, cough, myalgia, and fatigue are common, with eosinophilia being the sole finding suggestive of helminthic infection. Ground itch—the typical rash of human hookworm infection—is often intensely pruritic, and can continue for weeks.

Since many other disorders more familiar to Western practitioners (eg, atopy and lymphatic and solid cancers) can be associated with similar symptoms and eosinophilia, many patients undergo the added morbidity of multiple radiological and invasive diagnostic procedures.

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### Soil-transmitted helminth infections

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### Table: Geohelminthic infections causing eosinophilia in travellers

<table>
<thead>
<tr>
<th>Mode of transmission</th>
<th>Diagnosis</th>
<th>Mode of transmission</th>
<th>Diagnosis</th>
<th>Treatment</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ascaris lumbricoides</td>
<td>Soil 2 months</td>
<td>Nematodirus stercoralis</td>
<td>Soil 3 months</td>
<td>Albendazole, mebendazole, ivermectin</td>
</tr>
<tr>
<td>Human hookworm spp</td>
<td>Soil 6–8 weeks</td>
<td>Soil 4 weeks</td>
<td>Soil 3 months</td>
<td>Albendazole, mebendazole, ivermectin</td>
</tr>
<tr>
<td>Strongyloides stercoralis</td>
<td>Soil 6–8 weeks</td>
<td>Soil 4 weeks</td>
<td>Soil 3 months</td>
<td>Albendazole, mebendazole, ivermectin</td>
</tr>
<tr>
<td>Trichuris trichura</td>
<td>Soil 2 months</td>
<td>Soil 4 weeks</td>
<td>Soil 3 months</td>
<td>Albendazole, mebendazole, ivermectin</td>
</tr>
<tr>
<td>Zoonotic hookworms, ascarids (cutaneous, visceral larva migrans)</td>
<td>Soil 6–8 weeks</td>
<td>Soil 4 weeks</td>
<td>Soil 3 months</td>
<td>Albendazole, mebendazole, ivermectin</td>
</tr>
<tr>
<td>Angiostrongyloides cantonensis (eosinophilic gastroenteritis)</td>
<td>Soil 2 months</td>
<td>Soil 4 weeks</td>
<td>Soil 3 months</td>
<td>Albendazole, mebendazole, ivermectin</td>
</tr>
</tbody>
</table>

*Table: Geohelminthic infections causing eosinophilia in travellers*
The thought-provoking article by Desmond Sheridan (May 20, p 1698) raises some interesting questions. Academic medicine does not only require good scientists and researchers, but good clinicians, who contribute immensely to clinical medicine and the translation of laboratory research into clinical practice. In a culture of “publish or perish”, those who want to do both clinical and academic medicine find it difficult to find a balance between the two, mainly because of lack of time and resources. At least in the USA, people who work in academic institutions have to compete with doctors in the private sector for revenue generation. Thus academic medicine is going further and further away from the bedside and into the conference room. Those who are exceptional clinicians should be recognised as such and should have a role in academic medicine. We should involve clinicians under the umbrella of academics, for a large part of academic medicine is teaching at the bedside and training medical students, which can only come with clinical skills and spending time with patients.

Academic medicine should provide two systems: one for researchers and one for pure clinicians, who should support each other in academic endeavours. Unless we recognise those who spend most of their time looking after patients, we will lose them to the private sector and end up with a glut of researchers and no one to teach clinical skills and bedside medicine to the coming generation.

I declare that I have no conflict of interest.

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1 Sheridan DJ. Reversing the decline of academic medicine in Europe. Lancet 2006; 367: 1698–701.

Desmond Sheridan1 mentions several signs and solutions associated with academic medicine in Europe. I would like to call attention to the role of general practice in academic medicine and in medical research. During the past decade, changes in medical school curricula that put greater emphasis on early exposure to patients, clerkships with community-based clinicians,7 and longitudinal clinical experiences1 were seen as ways to stimulate a renewed interest in primary-care practice. But is this sufficient for general practice to be an attractive academic discipline?

There is growing evidence that absence of research in primary care could lead to overinvestigation of patients, inappropriate treatment, and diagnostic delay through wrong-track referral.4

Four “evidence gaps” in primary care involve the effectiveness of interventions delivered mainly in primary care, the applicability of hospital-based research to primary care, the implementation of best evidence in primary-care practice, and the basic science of illness and its care in the community.4

Despite changes in the healthcare system and in education, students and residents encounter