

Since January 2020 Elsevier has created a COVID-19 resource centre with free information in English and Mandarin on the novel coronavirus COVID-19. The COVID-19 resource centre is hosted on Elsevier Connect, the company's public news and information website.

Elsevier hereby grants permission to make all its COVID-19-related research that is available on the COVID-19 resource centre - including this research content - immediately available in PubMed Central and other publicly funded repositories, such as the WHO COVID database with rights for unrestricted research re-use and analyses in any form or by any means with acknowledgement of the original source. These permissions are granted for free by Elsevier for as long as the COVID-19 resource centre remains active. James F Donohue

Division of Pulmonary & Critical Care Medicine, University of North Carolina School of Medicine, Chapel Hill, NC 27599, USA

jdonohue@med.unc.edu

I declare that I have no conflict of interest.

- 1 Hogg JC. Pathophysiology of airflow limitation in chronic obstructive pulmonary disease. *Lancet* 2004; **364**: 709–21.
- 2 Rennard SI. Treatment of stable chronic obstructive pulmonary disease. Lancet 2004; **364:** 791–802.
- 3 Bowler RP, Barnes PJ, Crapo JD. The role of oxidative stress in chronic obstructive pulmonary disease. J COPD 2004; 1: 255–77.
- 4 Boman G, Backer U, Larsson S. Oral acetylcysteine reduces exacerbation rate in chronic bronchitis. *Eur J Respir Dis* 1983; **64:** 405–15.
- 5 Sinn DD, McAlister FA, Man SF, et al. Contemporary management of chronic obstructive pulmonary disease. JAMA 2003; 290: 2301–16.
- 6 Anthonisen NR, Connett JE, Murray RP. Smoking and lung function of lung health study participants after 11 years. Am J Respir Crit Care Med 2002; 166: 675–79.
- 7 Dirksen A, Diskman JH, Medsen F, et al. A randomized clinical trial of α_1 antitrypsin augmentation therapy. Am J Respir Crit Care Med 1999; **160**: 1468–72.
- 8 The Alpha-1 Antitrypsin Deficiency Registry Study Group. Survival and

 FEV_1 decline in individuals with severe deficiency of α_1 -antitrypsin. Am J Respir Crit Care Med 1998; **158**: 49–59.

- 9 Highland KB, Strange C, Heffner JE. Long-term effects of inhaled corticosteroids on FEV, in patients with chronic obstructive pulmonary disease: a meta-analysis. Ann Intern Med 2003; **138**: 969–73.
- 10 Anthonisen NR, Connett JE, Kiley JP, et al. Effects of smoking intervention and the use of an inhaled anticholinergic bronchodilator on the rate of decline of FEV₁. JAMA 1994; 272: 1497–505.
- 11 The TORCH Study Group. The TORCH (Towards a Revolution in COPD Health) survival study protocol. *Eur Respir J* 2004; **24**: 206–10.
- 12 Decramer M, Bartolome C, Tashkin D, et al. Clinical trial considerations in assessing long-term functional impacts of tiotropium in COPD: the Uplift Trial. J COPD 2004; 1: 303–12.
- 13 Celli BR, Cote CG, Marin JM, et al. The body-mass index, airflow obstruction, dyspnea, and exercise capacity index in chronic obstructive pulmonary disease. N Engl J Med 2004; **350**: 1005–12.
- 14 Zamel N, McClean P, Zhu J, et al. Effect of cilomilast (airflo) on trapped gas volume and indices of hyperinflation in patients with chronic obstructive pulmonary disease. Am J Respir Crit Care Med 2002: 165: A226 (abstr).
- 15 Grassi C, Morandini GC. A controlled trial of intermittent oral acetylcysteine in the long-term treatment of chronic bronchitis. *Eur J Clin Pharmacol* 1976; **5–6:** 393–96.
- 16 Black PN, Morgan-Day A, McMillan TE, Poole PJ, Young RP. Randomised, controlled trial of N-acetylcysteine for the treatment of acute exacerbations of chronic obstructive pulmonary disease [ISRCTN21676344]. BMC Pulm Med 2004; 4: 13.

Extending the benefits of deworming for development

See Articles page 1561

In today's *Lancet*, Charles King and colleagues, in a metaanalysis of functional disability due to schistosomiasis, show that schistosome infection is associated with significant anaemia, chronic pain, diarrhoea, reduced exercise tolerance, and malnutrition.

Previous assessments of the public-health relevance of schistosomiasis have focused mainly on symptomatic morbidity and late-stage disease.¹ Van der Werf et al² estimated clinical morbidity associated with schistosome infection in sub-Saharan Africa—eg, 70 million



Schoolchildren in Laos being treated for worms at school deworming day organised by teachers

Photo is by Carlo Urbani, who first described severe acute respiratory syndrome in Vietnam and died of the disease on March 29, 2003. He was the WHO focal point for parasitic diseases in the Western Pacific. cases of haematuria, 18 million cases of major bladderwall abnormalities, and 10 million cases of major hydronephrosis associated with *Schistosoma haematobium*.² To this burden of disease, King and colleagues convincingly add much subtle morbidity.

Disability-adjusted life years (DALYs) are increasingly used as a non-monetary measure of the impact of mortality and morbidity caused by a disease. Subtle functional disability is very relevant in soil-transmitted helminthiasis, another group of highly prevalent helminths.³ Regular treatment is clearly linked with physical and cognitive development, educational outcome, and economic development.⁴⁵ Consequently, the estimated DALYs lost due to these infections have been rated higher than those lost to schistosomiasis.

In 2001, a WHO Expert Committee concluded that the current figure for DALYs lost to schistosomiasis was considerably underestimated, and recommended that the figure should be revised to take into account the subtle morbidity induced by this disease.⁶ King and colleagues provide this missing information. As a result, we can readjust the disability weight currently assigned to schistosomiasis—and the resulting DALYs lost—to a much higher level. King's results should trigger a better quantification of the development impact of schistosomiasis. Beyond this, their analysis should encourage a

comprehensive re-evaluation of the burden on human and economic development of a group of highly prevalent but still concealed communicable diseases of the poor, including soil-transmitted helminthiasis, lymphatic filariasis, onchocerciasis, cysticercosis, echinococcosis, foodborne trematode infections, and trachoma.

The past 20 years of schistosomiasis control have been characterised by two major advances. The first is the acknowledgment that even in areas where reinfection is intense, regular chemotherapy can effectively control morbidity. The second is the endorsement in 2001 by the World Health Assembly of a novel public-health strategy for the integrated control of soil-transmitted helminthiasis and schistosomiasis.⁷ The aim of this strategy, tailored specifically for areas with high transmission, is to remove the disease burden by regular treatment of high-risk groups within a broader context of preventive measures such as improvement of living conditions and hygienic behaviour.⁸⁻¹⁰

When possible, regular treatment should be delivered through existing channels for the sake of sustainability. School health-programmes, also targeted at non-enrolled school-age children, are an excellent vehicle for the delivery of integrated interventions to a fundamentally high-risk group. Recently, we have also seen a multiplication of country experiences for the delivery of deworming to preschool children, packaged with vaccinations and/or vitamin A distribution.¹¹ The communitydirected treatment approach used in the onchocerciasis control-programme might be an option for delivery of combined treatment packages to remote communities.

We need to strengthen the links between deworming programmes and other chemotherapy-based programmes against endemic diseases affecting poor people. The delivery channels we mention above provide realistic opportunities for the health system to extend its capacity for the packaging and delivery of a series of simple health interventions to those most in need.¹² The combined delivery of antiparasitic treatment is likely to be highly cost effective because most drugs are today cheap or donated.

However, in the current context, two concerns need to be raised. The first is the need to ensure sustainability of delivery, because regular treatment will have to be delivered for a long time before improvement of living conditions will eventually provide a permanent solution. The second concern is the potential limitation of a chemotherapy-based strategy should drug resistance arise. We therefore believe that appropriate tools need to be developed and mechanisms put in place to enable monitoring of any reduction in drug efficacy so that strategic changes can be made in a timely manner.¹³ We also believe that research for new drugs and new control tools, such as the possible development of a hookworm vaccine, should be pursued.¹⁴

King and colleagues have added a further dimension to the effect that chemotherapy against schistosomiasis may have on disability. We believe that this novel information adds strength to the process of development of a comprehensive public-health strategy to control the burden of chronic endemic diseases in the developing world. We also hope that such a strategy will yield a high return on investment in terms of contribution towards reaching the Millennium Development Goals.¹⁵

*Lorenzo Savioli, Dirk Engels, Hiroyoshi Endo Parasitic Diseases and Vector Control (LS, DE), and Communicable Diseases Control, Prevention and Eradication (HE), WHO, 1211 Geneva 27, Switzerland saviolil@who.int

We thank David Brandling-Bennett, David W T Crompton, Alan Fenwick, Peter Hotez, David Molyneux, and Martin G Taylor for their advice and suggestions. We declare that we have no conflict of interest.

- Gryseels B. The relevance of schistosomiasis for public health. Trop Med Parasitol 1989; 40: 134–42.
- 2 Van der Werf MJ, de Vlas SJ, Brooker S, et al. Quantification of clinical morbidity associated with schistosome infection in sub-Saharan Africa. Acta Trop 2003; 86: 125–39.
- 3 Chan MS. The global burden of intestinal nematode infections—fifty years on. Parasitol Today 1997; 13: 438–43.
- 4 Bundy DAP, Barcelona D, Beegle K, et al. School health and nutrition programs. In: Jamison DT, Alleyne G, Breman J, et al, eds. Disease control priorities in developing countries, 2nd edn. New York: World Bank and Oxford University Press (in press).
- 5 Bleakley H. Disease and development: evidence from the American South. J Eur Econ Assoc 2003; 1: 376–86.
- 6 WHO. Prevention and control of schistosomiasis and soil-transmitted helminthiasis: report of a WHO Expert Committee. Geneva: WHO, 2002.
- 7 WHO. World Health Assembly Resolution 54.19. Schistosomiasis and soil-transmitted helminth Infections. May 21, 2001: http://www.who.int/ wormcontrol/about_us/en/ea54r19.pdf (accessed Jan 31, 2005).
- 8 Savioli L, Stansfield S, Bundy DAP, et al. Schistosomiasis and soil-transmitted helminth infections. Trans R Soc Trop Med Hyg 2002; 96: 577–79.
- 9 Savioli L, Engels D, Roungou JB, Fenwick A, Endo H. Schistosomiasis control. Lancet 2004; 363: 658.
- 10 Engels D, Savioli L. Public health strategies for schistosomiasis control. In: Secor WE, Colley DG, eds. Schistosomiasis: world class parasites, vol 10. New York: Springer, 2005.
- 11 WHO/UNICEF. How to add deworming to vitamin A distribution. Document WHO/CDS/CPE/PVC/2004.11. Geneva: WHO, 2004.
- 12 Molyneux DH. "Neglected" diseases but unrecognised successes. Lancet 2004; **364**: 380–83.
- 13 Albonico M, Engels D, Savioli L. Monitoring drug efficacy and early detection of drug resistance in human soil-transmitted nematodes. Int J Parasitol 2004; 34: 1205–10.
- 14 Hotez PJ, Brooker S, Bethony JM, Bottazzi ME, Loukas A, Xiao S. Hookworm. N Engl J Med 2004; 351: 799–807.
- 15 Editorial. Thinking beyond deworming. Lancet 2004; 364: 1993-94.