11 Rieder HL, Watson JM, Raviglione MC, Forssbohm M, Migliori GB, Schwoebel V, Leitch AG, Zellweger J-P. Surveillance of tuberculosis in Europe. Eur Respir J 1996;9(5):1097-104.

12 Chaulk CP, Moore-Rice K, Rizzo R, Chaisson RE. Eleven years of community-based directly observed therapy for tuberculosis. JAMA 1995;274(12):945-51.

13 Weis SE, Slocum PC, Blais FX, King B, Nunn M, Matney GB, Gomez E, Foresman BH. The effect of directly observed therapy on rates of drug resistance and relapse in tuberculosis. N Engl J Med 1994;330(17):1179-84.

14 Cohn DL, Catlin BJ, Peterson KL, Judson FN, Sbarbaro JA. A 62-dose, 6-month therapy for pulmonary and extrapulmonary tuberculosis: a twice-weekly, directly observed, and cost-effective regimen. Ann Int Med 1990;112:407-15.

15 Snider DE, Graczyk J, Bek E, Rogowski J. Supervised six-months treatment of newly diagnosed pulmonary tuberculosis using isoniazid, rifampin, and pyrazinamide with and without streptomycin. Am Rev Resp Dis 1984;130:1091-4.

16 Farmer PE, Kim JY. Community-based approaches to the control of multidrug-resistant tuberculosis: introducing 'DOTS-plus'. BMJ 1998,317:671-4.

17 Farmer PE. Managerial successes, clinical failures. Int J Tuberc Lung Dis 1999;3(5):365-7.

The global impact of drug-resistant tuberculosis. Boston,
 MA: Harvard Medical School and the Open Society Institute, 1999.
 Farmer PE, Shin SS, Bayona J, Kim JY, Furin JJ, Brenner JG.

Making DOTS-Plus Work, Chapter 19. In: Bastian I, Portaels F,

editors. Multidrug-resistant tuberculosis. Kluwer Academic Publications, The Netherlands, 2000.

20 World Health Organization. Coordination of DOTS-Plus pilot projects for the management of MDR-TB. Geneva: World Health Organization, 1999a. WHO/CDS/CBC/TB/99.262.

21 Kim JY, Bayona J, Furin JJ, Shin SS, Farmer PE. Making DOTS-Plus work: laboratories, drug procurement, planning, and evaluation. In: The Global Impact of Drug-Resistant Tuberculosis. Boston, MA: Harvard Medical School and the Open Society Institute. 1999.

22 Banatvala N, Matic S, Kimerling M, Farmer PE, Goldfarb A. Tuberculosis in Russia. Lancet 1999;354:1036.

23 Farmer PE, Robin S, Ramilus SL, Kim JY. Tuberculosis, poverty, and 'compliance': lessons from rural Haiti. Semin Respir Infect 1991;6(4):373-9.

24 Farmer PE. Social scientists and the new tuberculosis. Soc Sci Med 1997;44(3):347-58.

25 Farmer PE. Managerial successes, clinical failures. Int J Tuberc Lung Dis 1999;3(5):365-7.

26 Garrett L. The Coming Plague. New York: Farrar, Straus and Giroux, 1994.

27 Bifani PJ, Plikaytis BB, Kapur V, Stockbauer K, Pan X, Lutfey ML, Mogzaheh SL, Eisner W, Daniel TM, Kaplan MH, Crawford JT, Musser JM, Kreiswirth BN. Origin and interstate spread of a New York City multidrug-resistant Mycobacterium tuberculosis clone family. JAMA 1996;275(6):452-7.

The effect of school screening on surgery for adolescent idiopathic scoliosis

Reanalysis is needed

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e read the paper of Wiegersma et al.¹ regarding the effect of school screening on surgery for adolescent idiopathic scoliosis with much interest. It is based on a comparison of the number of cases of surgery for idiopathic scoliosis in Dutch regions in which the youth health care (YHC) department screens and does not screen for this disorder. The topic is important as scoliosis is disabling and may require extensive surgery. An intervention which effectively prevents extensive surgery at reasonable costs should thus be encouraged. Regarding school screening for idiopathic scoliosis (and subsequent conservative treatment) Wiegersma et al.¹ concluded that this does not reduce population rates for scoliosis surgery and should thus be reconsidered. Unfortunately, this conclusion seems to be based on a flawed interpretation by the authors of their own data.

Central in the study of Wiegersma et al.¹ are the data as presented in *table 1* of their paper, which is reproduced

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epidemiologist, Netherlands Organisation of Applied Scientific Research (TNO), Institute of Prevention and Health, P.O. Box 2215, 2301 CE Leiden, The Netherlands, tel +31 71 5181770, fax +31 71 5181903, e-mail. SA.Reijneveld@pg.tno.nl here. In their calculations they first assumed that screening for scoliosis of children aged 12 or 13 years will prevent surgery for scoliosis in the age group 12–19 years. On the basis of this, they concluded that screening for scoliosis

 Table 1 Distribution of cases and referents across YHC

 departments which screen (screening) and do not screen

 (non-screening) for idiopathic scoliosis

| Cases by age | | Non- | | |
|-----------------|-----------|-----------|------|-----------|
| (years) | Screening | screening | ORª | 95% CI |
| 12 | 6 | 7 | 1.00 | 0.74-1.35 |
| 13 | 15 | 15 | 0.97 | 0.71-1.33 |
| 14 | 14 | 27 | 0.86 | 0.61-1.23 |
| 15 | 17 | 25 | 0.87 | 0.57-1.32 |
| 16 | 8 | 19 | 0.71 | 0.40-1.26 |
| 17 | 7 | 9 | 0.74 | 0.34-1.62 |
| 18 | 2 | 10 | 0.30 | 0.07-1.34 |
| 19 | 0 | 1 | - | |
| Total cases | 69 | 113 | | |
| Referents total | 413 152 | 676.840 | | |

a: Odds ratio (OR) regarding the number of surgery cases among children of this age and older (but <20 years); the odds ratios and 95% confidence intervals (CI) for each age group refer to the number of cases and referents of that age and older has no preventive effect regarding surgery for scoliosis: the odds ratio (OR) comparing regions in which screening is performed with regions in which this is not done is 1.00 (whereas only values between 0 and 1 indicate a preventive effect). However, their assumption in constructing this table is that screening of children aged 12–13 years should have an immediate effect for children aged 12 or 13 years as well. This is very unlikely: it means that after detection of early scoliosis, a conservative treatment should be started immediately and should have immediate effects too. In some instances, the screening should even have an effect before the child could actually have been screened.

Wiegersma et al.¹ recognised that this line of thinking is logically impossible and, therefore, performed a second calculation in which they excluded children with surgery sooner than 1 year after possible screening. However, this calculation seems to be seriously flawed. It is based on 58 cases with surgery in the screening regions and 92 cases in the non-screening regions. We cannot derive these numbers of cases in any way from their tables: to be unbiased, regarding cases and controls from similar age groups should be excluded. We do not know the exact age at which children in the non-screening regions would have been screened if there had been screening in these regions. However, assuming like Wiegersma et al.¹ do that screening occurs in the age group 12-13 years and that the effects of a conservative treatment can only be expected after at least 1 year, children aged 12-14 years should be excluded from the analysis. In this case the resulting OR (95% confidence interval) is 0.87 (0.57-1.32).^{2,3} This implies a somewhat preventive effect, though clearly without statistical significance due to the small number of cases involved. If the cut-off is set at a higher age, this preventive effect increases though it remains without statistical significance (table 1, last columns).

On the basis of this reanalysis we conclude that the data of Wiegersma et al.¹ indicate that screening for idiopathic scoliosis may have some preventive effect. However, their study is too small to yield an accurate estimate of this effect. Furthermore, the design of their study will lead to an underestimation of this preventive effect. Firstly, children may move from a screening to a non-screening region (and vice versa) in the period during which the effect of the screening on surgery rates would occur. This will always lead to an underestimation of the actual effects of this screening. Secondly, in the study period almost all Dutch children aged 12, 13 or 14 years received either a screening for scoliosis or a preventive examination by a YHC.⁴ Examination of the trunk is usually also part of the latter too. This implies that they compared the effect of specific screening for scoliosis with the effect of a combined examination, again leading to an underestimation of the net effect of such a screening. Thirdly, even in regions labelled as screening, some children will not be invited or not participate, again giving an underestimation. We invite Wiegersma et al.¹ to reformulate their conclusion in the sense that their study gives some, though inconclusive, evidence for effectiveness of screening on idiopathic scoliosis.

REFERENCES

1 Wiegersma PA, Hofman A, Zielhuis GA. The effect of school screening on surgery for adolescent idiopathic scoliosis. Eur J Public Hlth 1998;8:237-41.

2 Woolf B. On estimating the relation between blood group and disease. Ann Human Genet 1955;19:251-3.

3 Rothman KJ. Modern epidemiology. Boston: Little Brown, 1986.

4 Burgmeijer RJF, Van Geenhuizen YM, Filedt Kok-Weimar T, De Jager AM. Growing adult: evaluation of child health care 1996 (in Dutch). Leiden/Maarssen: TNO Prevention and Health/KPMG, 1997.

The effect of school screening on surgery for adolescent idiopathic scoliosis

Response to readers' comments

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Sir,

We were pleasantly surprised to learn that more than 1 year after its publication our article still generates enough interest to give rise to letters to the editor. Furthermore, we are grateful to the authors of the letter for bringing forward the interesting point concerning the age differ-

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ence at surgery and its possible explanation. It provides new insights into the reasons behind the inadvertently adverse effects of youth health care activities discussed in other publications,^{1–3} although we doubt this was the authors' intention.

Before explaining this in more detail, we will first address the comments of the authors. In their letter Reijneveld and Hirasing state that we should have excluded the 12–14 year olds *in toto*, because the effect of screening of scoliosis on surgery could only be expected 1 year after such a screening. This, of course, is a rather curious line



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