

devastating sexually transmitted infection. The phased implementation will allow further refinement of the structure and process of the screening programme over the coming months; however, successful implementation will only be achieved with a sustained commitment to joint working among stakeholders at local and national levels.

ACKNOWLEDGEMENTS

All members of the Chlamydia Advisory Group contributed to the preparation of this editorial.

Sex Transm Infect 2004;**80**:331–333.
doi: 10.1136/sti.2004.009787

Authors' affiliations

K A Fenton, HIV and Sexually Transmitted Infections Department, Health Protection Agency, London, UK
H Ward, Department of Infectious Disease Epidemiology, Imperial College London

Correspondence to: Dr Kevin A Fenton, HIV and Sexually Transmitted Infections Department, Health Protection Agency, Communicable Disease Surveillance Centre, 61 Colindale Avenue, London NW9 5EQ, UK; Kevin.fenton@hpa.org.uk

Syndromic approach

Current status of syndromic management of sexually transmitted infections in developing countries

B Vuylsteke

The syndromic approach has been a major step forward in rationalising and improving management of STI

D iagnosis of a presumed sexually transmitted infection (STI) has traditionally been based on either clinical diagnosis, which is often inaccurate and incomplete, or laboratory diagnosis, which is complex, very expensive, and may delay treatment. As early as the 1970s, public health physicians, particularly those working in Africa, became interested in testing simple clinical tools for controlling and treating STIs.¹ This resulted in the design and promotion of “syndromic management” guidelines for STIs by the World Health Organization in 1991.² The syndromic approach does not require identification of the underlying aetiology. Instead, it is based on the identification of a syndrome—that is, a group

REFERENCES

- 1 **House of Commons**. Select Committee on Health. *Third report on sexual health*. Available at www.parliament.the-stationery-office.co.uk/pa/cm200203/cmselect/cmhealth/69/6902.htm Last accessed on 1 March 2004.
- 2 **LaMontagne DS**, Fenton KA, Randall S, *et al*. Establishing the National Chlamydia Screening Programme in England: results from the first full year of screening. *Sex Transm Infect* 2004;**80**:335–41.
- 3 **Low N**, McCarthy A, Macleod J, *et al*. The chlamydia screening studies: rationale and design. *Sex Transm Infect* 2004;**80**:342–8.
- 4 **Adams EJ**, Charlett A, Edmunds WJ, *et al*. Chlamydia trachomatis in the United Kingdom: a systematic review and analysis of prevalence studies. *Sex Transm Infect* 2004;**80**:354–62.
- 5 **Adams EJ**, LaMontagne DS, Johnston AR, *et al*. Modelling the healthcare costs of an opportunistic Chlamydia screening programme. *Sex Transm Infect* 2004;**80**:362–70.
- 6 **Department of Health**. *The national chlamydia screening programme in England, Programme overview, core requirements and data collection*. London: DoH, April 2004;(2).
- 7 **Health Protection Agency**, SCIEH, ISD, National Public Health Service for Wales, CDSC Northern Ireland, the UASSG. *Renewing the focus. HIV and other sexually transmitted infections in the United Kingdom in 2002*. London: Health Protection Agency, November 2003.
- 8 Additional information on the epidemiology of genital chlamydia infections diagnosed in GUM clinics in the United Kingdom is available at www.hpa.org.uk/infections/topics_az/hiv_and_sti/sti-chlamydia/epidemiology/epidemiology.htm.
- 9 **Morre SA**, van den Brule AJC, Rozendaal L, *et al*. The natural course of asymptomatic Chlamydia

- trachomatis infections: 45% clearances and no development of clinical PID after one-year follow-up. *Int J STD AIDS* 2002;
13(Suppl 2):12–18.
- 10 **Cates W**, Wasserheit JN. Genital chlamydial infections: epidemiology and reproductive sequelae. *Am J Obstet Gynecol* 1991;**164**(6 Pt 2):1771–81.
- 11 **Honey E**, Aungood C, Templeton A, *et al*. Cost effectiveness of screening for Chlamydia trachomatis: a review of published studies. *Sex Transm Infect* 2002;**78**:406–12.
- 12 **Chief Medical Officer's Expert Advisory Group**. *Main report of the CMO's expert advisory group on Chlamydia trachomatis*. London: Department of Health, 1998.
- 13 **Department of Health**. *Chlamydia Screening Pilot: Report of 1999–2000 study*. London: DoH, 2002.
- 14 **Pimenta JM**, Catchpole M, Rogers PA, *et al*. Opportunistic screening for genital chlamydial infection I: Acceptability of urine testing in primary and secondary healthcare settings. *Sex Transm Infect* 2003;**79**:16–21.
- 15 **Pimenta JM**, Catchpole M, Rogers PA, *et al*. Opportunistic screening for genital chlamydial infection II: Prevalence among health care attenders, outcome and evaluation of positive cases. *Sex Transm Infect* 2003;**79**:22–7.
- 16 **Fenton K**, Korovessis C, Johnson AM, *et al*. Sexual behaviour in Britain: reported sexually transmitted infections and prevalence genital Chlamydia trachomatis infection. *Lancet* 2001;**358**:1851–4.
- 17 **Department of Health**. *The national strategy for sexual health and HIV*. London: DoH, July 2001.
- 18 **Department of Health**. *The national strategy for sexual health and HIV—implementation action plan*. London: DoH, June 2002.

syndrome (GUS).⁵ In another study in Indonesia, the positive predictive value (PPV) of a syndromic approach for gonococcal and/or chlamydial urethritis was between 75% and 97%, resulting in a low cost per real case treated.⁶ In addition, the cure rate for urethral discharge with the syndromic approach was 99%.⁶ In order to decrease the number of women who would be treated unnecessarily for cervical infections, a risk assessment was incorporated into the syndromic approach to vaginal discharge. As a result, a woman with a complaint of vaginal discharge is treated systematically for vaginal infections, but only if her risk assessment is positive will she receive treatment for gonococcal and chlamydial infection as well. Using a risk score assessment in Tanzania, the overtreatment rate for cervical infections decreased from 92% to 17% in pregnant women and from 89% to 36% in non-pregnant women with vaginal discharge.⁷ By the late 1990s, the syndromic approach was largely promoted and used worldwide, and not only in developing countries.

There is enough evidence now that the syndromic approach is effective and has had an impact on the STI epidemic. Dramatic declines in STI rates have been observed following control strategies based on the syndromic approach, such as in sex workers in Côte d'Ivoire, Senegal and South Africa, and in STI

Key messages

- The syndromic approach has been a major step forward in rationalising and improving management of STI
- The performance of genital ulcer syndrome (GUS) treatment flow charts depends on the aetiological patterns of GUS in different settings
- The risk score approach should not be used as an STI screening tool or diagnostic test in asymptomatic or poorly symptomatic women
- Simple and rapid point of care tests may contribute to improve STI care for women in the near future

clinics in Kenya and in Burkina Faso.⁸⁻¹⁰ The studies in Mwanza (Tanzania) and Masaka (Uganda) demonstrated the impact of syndromic management beyond the STI clinic attendees they targeted by decreasing STI prevalences in the general population: serological syphilis by 20% and male urethritis by 50% in Mwanza, and gonorrhoea by 70% in Masaka.^{11, 12} The declining prevalence of bacterial infections in some of the key syndromes in parts of Africa is a testimony to the success of widespread syndromic management use.⁹

In this issue of *STI* (p 392, Wolday *et al*) describe the results of a study on risk factors associated with the failure of syndromic management of STIs among women seeking treatment in a primary healthcare centre in Addis Ababa. Syndromic treatment did not result in clinical improvement in 30% of the women, and the GUS was significantly associated with treatment failure. The authors argue that the treatment failure is probably a result of the high proportion of ulcers caused by herpes simplex type 2 virus (HSV-2) in this high HIV prevalence setting. The performance of syndromic treatment flow charts depends on the aetiological patterns of the syndrome, and herpes is not addressed by the former WHO algorithms.¹³ The syndromic approach became victim of its own success; because of the improved control of chancroid and syphilis in some regions it has become apparent that the GUS, particularly in the sub-Saharan countries, is more frequently caused by HSV-2 infections. The WHO is currently

recommending including the treatment for HSV-2 in the management of genital ulcers, especially in settings where HSV-2 prevalence is 30% or higher.¹³ Adding aciclovir to the syndromic treatment of ulcers, however, will not necessarily lead to higher cure rates.

Another area of concern is the use of the syndromic management in low STI prevalence settings, especially when the approach is used as a screening tool.^{14, 15} It should be stressed that the syndromic approach was developed as a diagnostic tool in symptomatic patients, it was never meant to be a screening tool. Traditionally, screening tools are used to minimise the number of (more expensive) standard diagnostic tests by identifying a group of people with a higher than average prevalence of infection. In the absence of such a test, the risk score approach should not be used as a substitute for standard diagnosis because of its poor discriminative ability. The current picture may change, however, when simple, cheap, and rapid diagnostic tests for *Neisseria gonorrhoeae* and *Chlamydia trachomatis* are available in developing countries. The development of such tests is considered by STI control programme managers and STI specialists to be an absolute priority in STI research. Major progress has recently been made in this field. A rapid (25 minute), cheap (\$US 0.85) dipstick for chlamydial infection "Firstburst" has been developed recently and is awaiting FDA approval. Another duplex (*N gonorrhoeae* and *C trachomatis*) test is undergoing evaluation.¹⁶ These tests may represent an important breakthrough for STI control in symptomatic and asymptomatic women in developing countries.

In conclusion, the syndromic approach has been a major step forward in rationalising and improving management of STI, and its impact on the STI epidemic has been observed in various settings. However, syndromic algorithms have some shortcomings, and they should be periodically revised and adapted to the epidemiological patterns of STI in a given setting. Simple and rapid point of care tests might help the screening of asymptomatic and low symptomatic women and the diagnosis of STI in symptomatic women. Finally, we should not forget that many other factors play a part in the successful control of STIs, including availability of effective and affordable drugs, accessible and acceptable health services, training and supervision of healthcare workers, and behavioural interventions to prevent new infections by promoting safer sex.^{17, 18}

Sex Transm Infect 2004;**80**:333-334.
doi: 10.1136/sti.2004.009407

Correspondence to: Dr Bea Vuylsteke, Institute of Tropical Medicine, Antwerp, Belgium; [bvuykste@itg.be](mailto:bvuylsteke@itg.be)

REFERENCES

- 1 Meheus AZ. Practical approaches in developing nations. In: Holmes KK, Mardh P-A, Sparling PF, *et al*. *Sexually transmitted diseases*. 1st ed. New York: McGraw-Hill, 1984:998-1008.
- 2 World Health Organization. *Guidelines for the management of sexually transmitted infections*. Geneva: WHO, WHO/HIV_AIDS/2001, 01.
- 3 Lush L, Walt G, Ogden J. Transferring policies for treating sexually transmitted infections: what's wrong with global guidelines? *Health Policy and Planning* 2003;**18**:18-30.
- 4 STI. Syndromic approach to STD management. *Sex Transm Infect* 1998;**74**(Suppl 1).
- 5 Hun Y, Morse SA, Dangor Y, *et al*. Comparison of clinically directed, disease specific, and syndromic protocols for the management of genital ulcer disease in Lesotho. *Sex Transm Infect* 1998;**74**(Suppl 1):S23-8.
- 6 Djajakusumah T, Sudigdoadi S, Keersmaekers K, *et al*. Evaluation of syndromic patient management algorithms for urethral discharge. *Sex Transm Infect* 1998;**74**(Suppl 1):S29-33.
- 7 Mayaud P, ka-Gina G, Cornelissen J, *et al*. Validation of a WHO algorithm with risk assessment for the clinical management of vaginal discharge in Mwanza, Tanzania. *Sex Transm Infect* 1998;**74**(Suppl 1):S77-84.
- 8 Ghys PD, Diallo MO, Etienne-Traore V, *et al*. Increase in condom use and decline in HIV and sexually transmitted diseases among female sex workers in Abidjan, Cote d'Ivoire, 1991-1998. *AIDS* 2002;**16**:251-8.
- 9 Steen R. Eradicating chancroid. *Bull World Health Organ* 2001;**79**:818-26.
- 10 Nagot N, Meda N, Ouangre A, *et al*. Review of STI and HIV epidemiological data from 1990 to 2001 in urban Burkina Faso: implications for STI and HIV control. *Sex Transm Infect* 2004;**80**:124-9.
- 11 Mayaud P, Moshia F, Todd J, *et al*. Improved treatment services significantly reduce the prevalence of sexually transmitted diseases in rural Tanzania: results of a randomized controlled trial. *AIDS* 1997;**11**:1873-80.
- 12 Kamali A, Quigley M, Kakiyungi J, *et al*. Syndromic management of sexually-transmitted infections and behaviour change interventions on transmission of HIV-1 in rural Uganda: a community randomised trial. *Lancet* 2003;**361**:645-52.
- 13 World Health Organization. *Report of the consultation meeting on improving management of sexually transmitted diseases*. Geneva: WHO, 28-30 Nov, 2001.
- 14 Hawkes SJ, Morison L, Foster S, *et al*. Reproductive tract infections in women in low-income, low-prevalence situations: assessment of syndromic management in Matlab, Bangladesh. *Lancet* 1999;**354**:1776-81.
- 15 Kapiga SH, Vuylsteke B, Lyamya EF, *et al*. Evaluation of sexually transmitted diseases diagnostic algorithms among family planning clients in Dar es Salaam, Tanzania. *Sex Transm Infect* 1998;**74**(Suppl 1):S132-8.
- 16 Peeling R. *Sexually transmitted diseases diagnostics initiative (SDI)*. Presented at the consultation meeting on improving management of sexually transmitted diseases. Geneva: WHO, 28-30 Nov, 2001.
- 17 Buvé A, Changalucha J, Mayaud P, *et al*. How many patients with a sexually transmitted infection are cured by health services? A study from Mwanza region, Tanzania. *Trop Med Int Health* 2001;**6**:971-9.
- 18 UNAIDS/WHO. *Sexually transmitted diseases: policies and principles for prevention and care*. Geneva: UNAIDS, 1999, UNAIDS/01.11E.