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REFERENCES

1. **Hu D**, Hook EW III, Goldie SJ. Screening for Chlamydia trachomatis in women 15 to 29 years of age: a cost-effectiveness analysis. *Ann Intern Med* 2004;**141**:501–13.
2. **Boerma JT**, Mgalla Z. Introduction. In Women and infertility in sub-Saharan Africa: a multi-disciplinary perspective. Boerma JT, Mgalla Z, eds. KIT, Amsterdam, 2001:13–23.
3. **Larsen U**, Menken J. Measuring sterility from incomplete birth histories. *Demography* 1989;**26**:185–201.
4. **Brunham RC**, Cheang M, McMaster J, *et al.* Chlamydia trachomatis, infertility, and population growth in sub-Saharan Africa. *Sex Transm Dis* 1993;**20**:168–73.
5. **Roberts TE**, Robinson S, Barton P, *et al.* Screening for Chlamydia trachomatis: a systematic review of economic evaluations and modelling. *Sex Transm Infect* 2006;**82**:193–200.
6. **Low N**. Screening programmes for chlamydial infection: when will we ever learn? *BMJ* 2007;193–200.
7. **Wallace LA**, Scoular A, Hart G, *et al.* What is the excess risk of infertility in women following genital chlamydia infection? A systematic review of the evidence. *Sex Transm Infect* 2008;**84**:171–5.
8. **Wiesenfeld HC**, Cates W Jr. Sexually transmitted disease and infertility. In Sexually Transmitted Diseases (4th Edn). In: Holmes KK, Sparling FP, Stamm WE, *et al.*, eds. McGraw-Hill, 2008:1511–27.
9. **Brunham RC**, Pourbohloul B, MAK S, *et al.* The unexpected impact of a Chlamydia trachomatis infection control program on susceptibility to reinfection. *J Infect Dis* 2005;**192**:1836–44.

Prevention of HIV transmission in the UK: what is the role of male circumcision?

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These two scenarios raise the question of whether male circumcision should be used in the UK as a means to decrease HIV transmission. Lack of circumcision has long been recognised as a risk factor for HIV acquisition in heterosexual men.¹ This makes biological sense because superficial Langerhans cells, which express HIV-1 receptors, are more prevalent in the male foreskin than in the remainder of the penis.² In addition, decreased keratinisation of the foreskin increases susceptibility to minor trauma during intercourse, potentially aiding the passage of HIV.³ Finally, ulcerative sexually transmitted infections, found more commonly in uncircumcised men, are associated with increased rates of HIV transmission.⁴

The biological plausibility of adult male circumcision to reduce HIV transmission has now been shown to have clinical relevance in three recent large randomised controlled trials conducted in sub-Saharan

Africa, in which male circumcision reduced the rate of female-to-male HIV transmission by at least 50%.^{5–7} Overall, there was little evidence of increased risk-taking behaviour in circumcised men. Here we examine whether the results of the African trials are relevant in the UK.

THE ISSUE FOR THE UK

The yearly number of new HIV diagnoses in the UK has increased by 157% since 1997.⁸ Of the new diagnoses reported in 2006, 12% were in black African men, most of whom were thought to have been infected heterosexually in Africa, whereas 36% were in men who have sex with men (MSM). Indeed, 2006 saw the greatest yearly number of newly diagnosed HIV infections in MSM since the start of the epidemic. New prevention strategies for these groups are urgently needed.

SCENARIO 1

Based on the biological rationale and clinical trials data, the World Health Organization (WHO) has recently recommended that “countries with high prevalence, generalised heterosexual HIV

epidemics that currently have low rates of male circumcision consider urgently scaling up access to male circumcision services”.⁹ Although the hypothetical patient described in scenario 1 resides in the UK, we believe that the WHO guidelines would apply to him and argue that it would be appropriate to offer circumcision to him.

Current National Health Service (NHS) guidelines, however, appear to pertain mainly to paediatric circumcision and generally discourage the procedure. For example, NHS Choices notes that “In the UK, circumcision is only carried out on the NHS in cases where it is medically necessary. It is usually performed as a last resort when other types of treatment have been unsuccessful.”¹⁰ A similar stance is taken by the British United Provident Association, a major source of private healthcare insurance in the UK (personal communication with Actuarial and Medical Risk Analyst, British United Provident Association, 15 August 2007).

SCENARIO 2

In their guidelines, WHO made no specific recommendation regarding circumcision for MSM. The US Centers for Disease Control and Prevention also reached no conclusion on this issue for MSM in the United States.¹¹ Simple extrapolation from the data on African heterosexual men is clearly inappropriate because MSM engage in a variety of other sexual practices, including anal and oral intercourse, which can be either insertive or receptive. Of these practices, those involving anal sex, particularly receptive anal sex, are most likely to transmit HIV.

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Potentially, the circumcision of MSM who engage in insertive anal intercourse might protect them from infection from receptive partners or, if they are already infected, it might protect their receptive partners from infection. The results of studies on circumcision status and the risk of HIV in MSM are, however, very limited and conflicting. Both a cross-sectional and a prospective study of MSM in the United States found an increased prevalence of HIV in uncircumcised men.¹²⁻¹³ Neither study was designed to determine the specific sexual practice by which these men became infected. A more recent cross-sectional study of black and Latino MSM in three US cities found no evidence that circumcision was protective in men who engaged in unprotected insertive anal sex.¹⁴ A prospective Australian trial study also reported no relationship between circumcision status and HIV seroconversion in either the entire MSM study population or those men who denied practising receptive anal intercourse.¹⁵

Given the lack of data, we would also be unable to make a specific recommendation regarding the patient presented in scenario 2.

WHAT SHOULD BE DONE?

First, we believe the NHS and, potentially, private insurers, should examine their existing policies for adult male circumcision. For recent male émigrés from high HIV-prevalence countries in sub-Saharan Africa whose female partners are also from these countries, we believe circumcision should be made available in accordance with WHO recommendations.

Second, to determine definitively whether circumcision could play a role in HIV prevention for MSM, a clinical trial is needed. Major determinants of the feasibility of such a trial would be HIV incidence in the participants and current circumcision rates.

The Health Protection Agency estimated that among MSM attending sentinel genitourinary medicine clinics in London during 2006, HIV incidence was 2.6%,⁸ a rate generally similar to that seen in the non-circumcised control groups enrolled in the African clinical trials.⁵⁻⁷ Whereas these findings might suggest that an MSM trial could be performed with similar numbers of persons as in the African studies, an estimate of HIV incidence for men engaging in unprotected insertive anal sex would be needed for a sample size calculation.

The relatively low circumcision rates of UK men would also facilitate circumcision

Scenarios

Scenario 1: A 24-year-old HIV-uninfected heterosexual man who has recently emigrated from Kenya presents to a general practitioner surgery in the United Kingdom requesting a referral for circumcision, because several of his friends in Kenya have had the procedure to reduce their risk of acquiring HIV. His partners in the United Kingdom are also recent émigrés from sub-Saharan Africa. What should be the response?

Scenario 2: The presenting patient is an HIV-uninfected 24-year-old sexually active homosexual man from the United Kingdom, also requesting a referral for circumcision to reduce his risk of HIV infection. Should the response be different?

trials. The National Survey of Sexual Attitudes and Lifestyle of 2001 found that only 15.8% of UK men between the ages of 16 and 44 years had been circumcised.¹⁶

Of course, such trials would not be ethical without extensive risk-reduction counselling for participants and would not be possible without the willing participation of at-risk MSM. Pilot studies would be needed to determine the likely participation rates. A recent survey of uncircumcised MSM performed in the United States indicated that more than half would be willing to be circumcised if this were shown to reduce the risk of HIV infection.¹⁷

The primary study endpoint would be the effect of circumcision on HIV incidence for men engaging in unprotected insertive anal sex. We believe this study could be conducted in UK genitourinary medicine clinics. If additional trial sites were needed to recruit a sufficiently large number of participants, they could be identified in countries with a concentrated MSM HIV epidemic and low circumcision rates; eg, in Scandinavia or Latin America.

A second study might also be considered to examine the effect of circumcising previously infected MSM on the risk of infecting persons who are their receptive partners in unprotected anal sex. An analogous study of circumcision for HIV-infected Ugandan men to reduce the risk of infecting their female partners was, however, stopped prematurely.⁹ Preliminary data had shown no evidence of decreased transmission from circumcised men and suggested that men who resumed sexual activity before complete wound healing might pose an increased transmission risk to their partners. Further consultation would be needed to decide if this MSM study should be carried out in the UK.

The UK has the opportunity to lead in revising its male circumcision guidelines in accordance with new African data and

to develop data upon which to consider new circumcision strategies for MSM. We believe these opportunities are important and should not be missed.

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REFERENCES

1. **Moses S**, Bailey RC, Ronald AR. Male circumcision: assessment of health benefits and risks. *Sex Transm Infect* 1998;**74**:368–73.
2. **Patterson BK**, Landay A, Siegel JN, *et al.* Susceptibility to human immunodeficiency virus-1 of human foreskin and cervical tissue grown in explant culture. *Am J Pathol* 2002;**161**:867–73.
3. **McCombe SG**, Short RV. Potential HIV-1 target cells in the human penis. *AIDS* 2006;**20**:1491–5.
4. **Weiss HA**, Thomas SL, Munabi SK, *et al.* Male circumcision and risk of syphilis, chancroid and genital herpes: a systematic review and meta-analysis. *Sex Transm Infect* 2006;**82**:101–9.
5. **Auvert B**, Taljaard D, Lagarde E, *et al.* Randomized controlled intervention trial of male circumcision for reduction of HIV infection risk: the ANRS 1265 Trial. *PLoS Med* 2005;**2**:e298.
6. **Bailey RC**, Moses S, Parker CB, *et al.* Male circumcision for HIV prevention in young men is Kisumu, Kenya: a randomised controlled trial. *Lancet* 2007;**369**:643–56.
7. **Gray RH**, Kigozi G, Serwadda D, *et al.* Male circumcision for HIV prevention in men in Rakai, Uganda: a randomised trial. *Lancet* 2007;**369**:657–66.
8. **Health Protection Agency**. Testing times. HIV and other sexually transmitted infections in the UK: 2007. www.hpa.org.uk/infections/topics_az/hiv_and_sti/publications/AnnualReport2007/default.htm (accessed 8 Jan 2008).
9. **WHO and UNAIDS**. New data on male circumcision and HIV prevention: policy and programme implications. March 2007. www.who.int/hiv/mediacentre/MCRecommendations_en.pdf (accessed 8 Jan 2008).
10. **NHS Choices**. Circumcision. When should it be done? <http://www.nhs.uk/Conditions/Circumcision/Pages/When-should-it-be-done.aspx?url=Pages/What-is-it.aspx> (accessed 8 Jan 2008).
11. **Sullivan PS**, Kilmark PH, Peterman TA, *et al.* Male circumcision for prevention of HIV transmission: what the new data mean for HIV prevention in the United States. *PLoS Med* 2007;**4**:e223.

12. **Kreiss JK**, Hopkins SG. The association between circumcision status and human immunodeficiency virus infection among homosexual men. *J Infect Dis* 1993;**168**:1404–8.
13. **Buchbinder S**, Vittinghoff E, Heagerty P, *et al*. Sexual risk, nitrite inhalant use, and lack of circumcision associated with HIV seroconversion in men who have sex with men in the United States. *J Acquir Immune Def Syndr* 2005;**39**:82–9.
14. **Millett G**, Ding H, Lauby J, *et al*. Circumcision status and HIV infection among black and latino men who have sex with men in 3 US cities. *J Acquir Immune Defic Syndr* 2007;**46**:643–50.
15. **Templeton DJ**, Jin F, Prestage GP, *et al*. Circumcision status and risk of HIV seroconversion in the HIM cohort of homosexual men in Sydney. *4th IAS Conference on HIV Pathogenesis, Treatment and Prevention*. 22–25 July 2007, Sydney [Abstract WEAC 103].
16. **Dave SS**, Johnson AM, Fenton KA, *et al*. Male circumcision in Britain: findings from a national probability sample survey. *Sex Transm Infect* 2003;**79**:499–500.
17. **Begley E**, Jafa K, Voetsch A, *et al*. Willingness of men who have sex with men in the US to be circumcised as adults to reduce risk of HIV infection. *14th Conference on Retroviruses and Opportunistic Infections*; 25–28 February 2007, Los Angeles [Abstract 983].

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