

Sexually transmitted diseases among randomly selected attenders at an antenatal clinic in The Gambia

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SUMMARY One hundred randomly selected women attending a free government antenatal clinic in the town of Bakau, The Gambia, were examined. Vaginal swabs were taken for microscopical examination for *Trichomonas vaginalis* and for culture on Sabouraud's medium. Cervical swabs were taken for culture of *Neisseria gonorrhoeae* and *Chlamydia trachomatis* and, in 50 cases, *Herpesvirus hominis*; in addition, urethral swabs were taken for culture of *N gonorrhoeae*. Serum samples were tested for antibodies to *Treponema pallidum* by the Venereal Diseases Research Laboratory (VDRL) test and *T pallidum* haemagglutination assay (TPHA), and to *C trachomatis* and *H hominis* by microimmunofluorescence. The prevalence of infection with *Candida albicans* was found to be 35%, *T vaginalis* 32%, *C trachomatis* 6.9%, *N gonorrhoeae* 6.7%, *T pallidum* 1%, and *H hominis* 0%. IgG antibodies at a titre of at least 1/16 to *C trachomatis* serotypes D-K were found in 29.4%, and to serotypes A-C in a further 10.6%. IgG antibodies at a titre of at least 1/16 to *H hominis* type I were found in 94%, and to type II in 53%, although a proportion of the latter probably represent cross reacting antibodies to type I.

Introduction

Those who have worked in the specialty generally agree that sexually transmitted diseases (STD) are a major problem in developing countries.¹⁻³ As they are not notifiable in most of these countries, however, and as STD clinics are few and far between, the exact extent of the problem remains unknown. In a few African countries an attempt has been made to calculate the incidence of gonorrhoea by dividing the number of patients seen in a hospital in a given period by the population that it is thought to serve.^{4,5} In this way the yearly incidence of gonorrhoea in Swaziland and in Uganda has been estimated to be between 3000 and 10 000 per 100 000 total population, though clearly this method may underestimate the true incidence. An ingenious attempt has been made to calculate the incidence of syphilis in Swaziland from the rate at which positive results to serology tests increased with age,⁶ but in many African countries this is not possible because ages of

patients are not accurately known; moreover, positive results to serology tests in older patients may be due to endemic treponematoses rather than venereal syphilis.

An alternative method of estimating the importance of STD in a community is to measure its prevalence in a representative sample of the population. The problem here lies in the selection of the sample to be studied. Apart from the work of Arya *et al* in rural Uganda,⁷ most prevalence studies so far reported from Africa have been based on patients attending either family planning or antenatal clinics in urban areas.⁸⁻¹² How representative these are of the general population depends on the proportion of the population which attends the clinic, and how this proportion is selected. This is difficult to ascertain and varies according to the cost of treatment at each clinic as well as its location and reputation, which may account for some of the widely divergent results that have been reported by different authors.

In The Gambia free antenatal care is provided by the government, and it is estimated that at least 90% of pregnant women living in the town of Bakau, in which this study was carried out, attend the clinic on at least one occasion in each pregnancy (MGM

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Rowland, unpublished observation). We undertook this study to discover the prevalence of a variety of STDs among antenatal clinic attenders in this community.

Patients and methods

The Gambia is a small country on the west coast of Africa between latitudes 13° and 14° N. The population (between half a million and one million) is heterogeneous, consisting of four major tribes and several smaller minorities. The people are mainly Moslems, apart from the Jola tribe (most of whom are Roman Catholics) and the westernised professional classes in the capital (who are mainly Christians of various denominations). Polygamy is practised, and bride prices are high in relation to average earnings. Women are not strictly confined as in some Moslem countries, however, and many enjoy a considerable degree of independence.

This study was carried out between November 1981 and February 1982 on randomly selected patients in the third trimester of pregnancy attending the antenatal clinic in the town of Bakau (population 10 000), which is eight miles from the capital, Banjul. Every tenth patient was examined until 100 had been seen; only one of those asked to participate in the study declined to do so. The age, tribal origin, and obstetric history of each patient were recorded and they were specifically asked about the presence of lower abdominal pains, vaginal discharge, and pain on passing urine. A vaginal speculum was passed and the following specimens were taken: (1) a swab from the posterior fornix (examined for *Trichomonas vaginalis* in a wet preparation); (2) a swab from the posterior fornix or any other site where discharge had collected (cultured for *Candida albicans* on Sabouraud's medium); (3) cervical and urethral swabs for the isolation of *Neisseria gonorrhoeae* (plated direct on to modified Thayer-Martin medium and transferred to a 5% carbon dioxide incubator jar within two hours); (4) an endocervical swab for isolation of *Chlamydia trachomatis* (expressed immediately into sucrose phosphate transport medium containing 10% fetal calf serum and stored at -70°C); (5) an additional endocervical swab was taken from 50 patients for the isolation of *Herpesvirus hominis* (expressed immediately into viral transport medium containing 10% fetal calf serum and stored at -70°C); and (6) from all patients, 3 ml of venous blood (separated the same day; serum was stored at -20°C).

IDENTIFICATION OF PATHOGENS

The swabs were tested as follows:

Candida albicans—Inoculated plates of Sabouraud's

medium were incubated at 37°C for 48 hours, and colonies were examined as a wet preparation; yeasts were identified as *C. albicans* by the germ tube test.

Neisseria gonorrhoeae—After incubation at 37°C in 5% carbon dioxide for 48 hours, isolates were confirmed as *N. gonorrhoeae* by colonial appearance, Gram stain, and oxidase test.

Chlamydia trachomatis—Specimens were inoculated by centrifugation (2500 × g at 33°C for 1 hour) on to McCoy cells, which were subsequently treated with cycloheximide 2 mg/l in minimum essential medium containing glucose and 10% fetal calf serum, as described by Ripa and Mårdh.¹³ Monolayers were stained with Giemsa at 48 hours and examined for inclusions by dark field microscopy.

Herpesvirus hominis—Specimens were inoculated on to Vero cells in minimum essential medium with 5% fetal calf serum and examined daily for five days for cytopathic effect. A final examination was made after 14 days before specimens were discarded as negative.

SEROLOGICAL ANALYSIS

Serum samples were analysed as follows: the Venereal Diseases Research Laboratory (VDRL) test was performed according to standard techniques using undiluted serum and Wellcome reagent VD02-03; the *Treponema pallidum* haemagglutination assay (TPHA) was performed according to standard techniques using serum at dilutions of 1/80 and 1/160 with reagent obtained from Fujizoki pharmaceutical company, Tokyo; and tests for antibody to *C. trachomatis* and *Herpesvirus hominis* were carried out at the Institute of Ophthalmology, London.

Serum was examined for type specific antibodies at starting dilutions of 1/16 for IgG and 1/8 for IgM. A modified microimmunofluorescence test,¹⁴ using pooled, egg grown preparations of *C. trachomatis* serotypes A-C (trachoma types), D-K (oculogenital types), LGV 1 to LGV 3 (lymphogranuloma venereum types), and of *C. psittaci* was used to detect antibodies to chlamydiae. Antibodies to herpes virus were detected by a microimmunofluorescence test¹⁵ using cell culture grown antigens of herpes simplex types I and II.

TABLE 1 Tribal origins of 100 antenatal patients

Tribe	No
Jola	44
Mandinka	21
Wollof	11
Fula	13
Other	11

Results

CLINICAL FINDINGS

Table I shows the tribal origins of the 100 women examined (average age 23.6 (range 16-40) years).

Table II shows the symptoms described by these women on direct questioning. An abnormal vaginal discharge was noted on examination in 67, only 24 of whom had complained of this. There were 10 patients who complained of discharge in whom no abnormality was noted.

TABLE II Prevalence of symptoms described by 100 antenatal patients on direct questioning

Symptom	%
Lower abdominal pain	55
Vaginal discharge	34
Dysuria	27
No symptoms	34

PATHOGENS ISOLATED

Table III shows the prevalence of the various infective agents in this population. As was expected in view of the small numbers involved, no appreciable association was found between the isolation of *N gonorrhoeae* or *C trachomatis* and any symptom or sign; all six patients harbouring *N gonorrhoeae* and five of the six with *C trachomatis* had at least one symptom. Dysuria was complained of significantly more frequently in those infected with *C albicans* (15 out of 35) than in those not infected (12 out of 65) ($\chi^2 = 5.3$, $p < 0.025$). A vaginal discharge was seen in significantly more of those harbouring *T vaginalis* (28 out of 32) than in those who were not (39 out of 68) ($\chi^2 = 7.9$, $p < 0.005$).

TABLE III Isolation of pathogens from 100 antenatal patients

Pathogen	No tested	No (%) positive
<i>Candida albicans</i>	100	35 (35)
<i>Trichomonas vaginalis</i>	100	32 (32)
<i>Neisseria gonorrhoeae</i>	90	6 (6.7)
<i>Chlamydia trachomatis</i>	87	6 (6.9)
<i>Herpesvirus hominis</i>	50	0 (0)

NB 10 specimens for isolation of *N gonorrhoeae* and *C trachomatis* were lost due to electricity failure; three specimens for isolation of *C trachomatis* were contaminated.

SEROLOGY

Table IV shows the results of serological tests. Although the VDRL test was positive in nine cases, the TPHA was only positive in one. The remaining eight are presumably biological false positives due to pregnancy or a disease such as malaria, which is

TABLE IV Prevalence of antibodies to various pathogens in 100 antenatal patients

Pathogen	Test used	No tested	No (%) positive
<i>T pallidum</i>	VDRL	100	9 (9)
<i>T pallidum</i>	TPHA	100	1 (1)
<i>C trachomatis</i> A-C	MIF (IgG)	85	9 (10.6)
<i>C trachomatis</i> D-K	MIF (IgG)	85	25 (29.4)
<i>H hominis</i> type I	MIF (IgG)	85	80 (94.1)
<i>H hominis</i> type II	MIF (IgG)	85	45 (52.9)*

VDRL = Venereal Diseases Research Laboratory test; TPHA = *T pallidum* haemagglutination assay; MIF = Microimmunofluorescence test (a titre of 1/16 or more was considered positive).

*In all but 7% the titre to *H hominis* type I was higher than to type II.

NB 15 serum samples were lost in transit between The Gambia and the Institute of Ophthalmology.

prevalent in this area at the season when the study was undertaken.

IgG antibodies to *C trachomatis* were found at a titre of at least 1/16 in 34 (40.0%) specimens tested. In 25 (29.4%) patients, these antibodies were to *C trachomatis* serotypes D-K. In two of these women (whose swabs had not yielded chlamydia on culture) IgM against these serotypes was also detected at titres of 1/8 and 1/16. Nine (10.6%) patients had IgG antibodies specific for *C trachomatis* serotypes A-C. No patient had antibodies specific for lymphogranuloma venereum types of *C trachomatis* or *C psittaci*. The figure shows the distribution of titres of IgG antibodies to *C trachomatis* serotypes D-K among 85 patients tested. Their average age was 24.3 years, which is not appreciably different from the average age of the whole study population (23.6 years). Table V shows the prevalence of antibodies to *C trachomatis* by tribal group; although it was higher

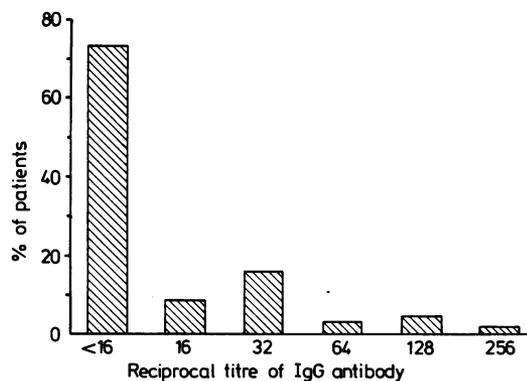


FIGURE Distribution of titres of IgG antibodies to *Chlamydia trachomatis* serotypes D-K in 85 antenatal patients.

TABLE V *Chlamydial antibody prevalence by tribe (serotypes D-K)*

Tribe	No tested	No (%) positive
Jola	37	14 (37.8)*
Mandinka	19	4 (21.1)
Fula	11	3 (27.3)
Wolof	11	3 (27.3)

*Jolas v others: 14 out of 37 v 10 out of 41 ($\chi^2 = 1.65$; $p > 0.2$).

among Jola women than others, the difference was not significant. It is, however, remarkable that all six women from whom *C trachomatis* was isolated belonged to this tribe. Patients yielding positive isolation results tended to be among the younger women examined (average 20.5 (range 18-24) years).

IgG antibodies to *H hominis* type I were present at a titre of at least 1/16 in 94% of women tested, and to type II in 52.9%. Many of the latter represent cross reacting type I antibodies, however, as the titre of antibodies to type I was higher in all but 7%. No patient had IgM antibodies to *H hominis*, and herpes virus was not isolated from any of the 50 women from whom swabs were tested.

Discussion

The prevalence of infection with *T vaginalis* (32%) found in this study is similar to that reported among randomly selected women in other antenatal clinics in Africa. In Ibadan it was found to be 21%,¹⁰ in Swaziland 23%,¹¹ in Zambia 38%,¹⁶ and in Durban 49%.¹⁷ By contrast, a study in Britain showed a prevalence of 4%.¹⁸ While 17 (53%) of our patients with this infection did not complain of vaginal discharge, there was an appreciable association between infection with *T vaginalis* and an abnormal discharge seen on examination. Of the 67 women in whom an abnormal discharge was seen, however, 39 (58%) were not infected with *T vaginalis*, and 27 of these were not infected with *C albicans* either. Presumably other pathogens such as *Gardnerella vaginalis* are also prevalent in this community and were responsible for the vaginal discharge seen in these patients. It is interesting to note that *G vaginalis* was isolated from 75 of 100 patients complaining of vaginal discharge in Nairobi.¹⁹

The prevalence of infection with *C albicans* (35%) seen in this community is also similar to that found in random surveys in other African antenatal clinics (23% in Durban,¹⁷ 33% in Ibadan,¹⁰ 37% in Swaziland¹¹). We found that isolation of this organism was associated with dysuria. Apart from the discomfort it may cause mothers, *C albicans* may infect infants in

up to 50% of cases,²⁰ and a high incidence of oral thrush has been seen in neonates in this community (MGM Rowland, unpublished observation.) This is occasionally so severe as to interfere with breast feeding.

The prevalence of gonorrhoea in African antenatal clinics has been found to vary from 0% in a small study population in Nairobi⁹ to 14% in Yaoundé.¹² It was 6.7% in this study and in most published studies it has been between 3% and 10%.^{10 11 21} That the prevalence may be even higher in rural areas is suggested by the findings of Arya *et al*, who showed 40% of pregnant women in the Teso district of Uganda to be infected,⁷ and those of Widy-Wirski and D'Costa who found 27.5% of all women in one rural village in the Central African Republic had gonorrhoea.²²

Apart from the present study, in which *C trachomatis* was isolated from 6.9% of patients, we are aware of only two other attempts to isolate this organism from the cervixes of women in Africa. Nsanze *et al* isolated *C trachomatis* from three (6%) of 54 antenatal patients in Nairobi,⁹ and in Johannesburg, Ballard *et al* isolated it from 13% of women attending an STD clinic and 16% of those attending a family planning clinic.²³ We found circulating IgM antibody to *C trachomatis* types D-K, which is strongly suggestive of active infection, in a further two patients. IgG antibodies to *C trachomatis* types D-K were found in 29.4% of our patients at a titre of at least 1/16. A further 10.6% had IgG antibodies to serotypes A-C, which may reflect exposure to genital infection or trachoma. High titres of IgG antibodies to *C trachomatis* have been found in the sexually active population in Nairobi⁹ and Johannesburg²³ as well as among STD clinic attenders in Ethiopia.²⁴ Among antenatal patients in Ibadan, Nigeria, the prevalence of IgG antibodies to types D-K was 8.4%.²⁵ Ballard has suggested that, by analogy with the chronic chlamydial eye disease, trachoma, the likelihood of isolating *C trachomatis* is reduced in patients with chronic cervical infections and high titres of circulating antibody. This would suggest that the true prevalence of genital *C trachomatis* infection is higher than isolation figures suggest in communities with a high prevalence of circulating antibody, which is a debatable hypothesis.

Heyman estimated that 30% of babies born to women with cervical gonorrhoea develop gonococcal ophthalmia neonatorum.²⁶ This potentially blinding condition is therefore likely to occur in about 2% of babies born in Bakau. A previous study showed that 30% of infants born in the government hospital in Banjul developed conjunctivitis in the first three days of life.²⁷ Although most of these cases were neither severe nor of gonococcal origin, it would seem that prophylaxis with silver nitrate may play a useful part

in preventing serious eye disease in Gambian infants.

The role of *C trachomatis* in the aetiology of ophthalmia neonatorum in Africa has been studied little, and was first described in The Gambia in 1965 by the MRC Trachoma Unit.²⁸ We described a series of 37 patients with ophthalmia neonatorum in whom *C trachomatis* was isolated from 35% and *N gonorrhoeae* from 24%.²⁹ Méheus *et al* found a 9.7% incidence of ophthalmia neonatorum in Bangui, 26% of cases being due to *N gonorrhoeae* and 19% to *C trachomatis*.³⁰ The incidence of other diseases due to *C trachomatis* among neonates, such as pneumonitis, is not known and is currently under investigation.

The most important complications of gonorrhoea are pelvic inflammatory disease (PID) and its sequelae, notably ectopic pregnancy and infertility. Westrom *et al* have shown that in Sweden the incidence of ectopic pregnancy is increased sevenfold following an attack of PID.³¹ Figures for the incidence of ectopic pregnancy in Africa are scarce, but several workers have drawn attention to the high incidence of PID in Africa,^{32,33} and most have linked it with gonococcal infection.³³⁻³⁵ *C trachomatis* is the major cause of PID in certain industrialised countries.³⁶ The results of this study and those from Nairobi and Johannesburg suggest that *C trachomatis* may be at least as important as *N gonorrhoeae* in causing PID in Africa.

At least 12% of women in Sweden are rendered infertile after a single attack of PID, even with optimum antibiotic treatment,³⁷ and the figure rises to over 30% after two attacks. In Africa, where suitable antibiotic treatment is not obtainable in many areas, it seems likely that a higher proportion of women become infertile following an attack of PID. Many workers have drawn attention to the high rate of infertility in certain parts of Africa.^{32,38-40} In The Gambia, a careful longitudinal study of births and deaths over 25 years in two rural villages has shown that 3% to 5% of women suffer from primary infertility, and 13% to 19% from secondary infertility (defined as failure to bear children after the age of 30).⁴¹ Exact figures are not available for other areas, but over 50% of gynaecological consultations in the government hospital in Banjul are for infertility, and most women investigated by hysterosalpingography or at operation have bilateral tubal occlusions (G Ogbasellasse, unpublished observation). Ballard *et al* have shown appreciably higher titres of circulating chlamydial antibody in women in Johannesburg with tubal infertility than in controls with normal fallopian tubes, suggesting that chlamydial salpingitis is an important cause of infertility in that community.²³ Similar studies are needed elsewhere in Africa if a rational policy is to be suggested for the control of PID and its sequelae.

It is sometimes stated that venereal syphilis in Africa is more common in urban areas.⁴²⁻⁴⁴ This has not been our experience in The Gambia. Only one of 100 urban women in this study had serological evidence of syphilis, whereas at a mission hospital 60 miles from the capital 11 of 100 randomly selected antenatal patients had positive results to VDRL and TPHA tests (DCW Mabey, unpublished observation). Endemic treponematoses was prevalent in The Gambia in the early 1950s,⁴⁵ before the World Health Organisation's yaws eradication programme, but has not been seen here subsequently. It therefore seems likely that all positive serological tests in patients aged under 30 are due to venereal syphilis.

Congenital syphilis is diagnosed fairly frequently at the government hospital in Banjul and at the MRC hospital in Bakau (about 30 cases a year in all), but many other cases are probably missed as serological tests are not performed elsewhere in the country. Late manifestations of congenital syphilis are rarely seen in The Gambia. For example, only one case of interstitial keratitis has been seen at the government eye clinic in the past eight years (S Sowa, unpublished observation). This suggests either that there is a very high mortality among those with congenital syphilis, or that the disease is becoming more common. If the latter is the case, it may well be that we are experiencing an epidemic of venereal syphilis in a population previously rendered immune by endemic treponematoses in childhood. This hypothesis is supported by the relative rarity of tertiary syphilis.

It is difficult to interpret the results of serological tests for antibody to *H hominis*, as there is cross reaction between antibodies to types I and II with the microimmunofluorescence test used.¹⁵ It is clear that at least 90% of patients studied had IgG antibody to type I. Although antibodies to type II were detected in 53% of patients, in all but 7% of them the titre of type I antibodies was higher, suggesting the possibility of cross reacting type I antibodies. Studies of the aetiology of genital ulceration in Africa suggest that *H hominis* is responsible for a much smaller proportion of ulcers than in Europe and North America.^{46,47} In a pilot study carried out in The Gambia in 1981 *H hominis* was isolated from only one of 37 men with genital ulceration (DCW Mabey and HC Whittle, unpublished observation). This may simply reflect the higher prevalence of genital ulceration due to other pathogens in Africa, although it is tempting to postulate that Africans are protected from genital herpes by antibodies acquired in childhood as a result of non-venereal infections. Neutralising antibodies to *H hominis* type II have been found in young Nigerian children,⁴⁸ and further studies are needed in other areas to elucidate this issue.

References

1. Anonymous *Annual medical and sanitary report for 1921*. Entebbe, Uganda: Government Printer, 1921; Appendix III: 69.
2. Arya OP, Osoba AO, Bennett FJ. *Tropical Venereology*. 1st ed. Edinburgh, London, and New York: Churchill Livingstone, 1980.
3. Osoba AO. Sexually transmitted diseases in tropical Africa: a review of the present situation. *Br J Vener Dis* 1981; **57**:89-94.
4. Arya OP. Changing patterns in the organisation of the venereal diseases and treponematoses service in Uganda. *Br J Vener Dis* 1973; **49**:134-8.
5. Méheus A, Ballard R, Dlamini M, Urusi JP, van Dyck E, Piot P. Epidemiology and aetiology of urethritis in Swaziland. *Int J Epidemiol* 1980; **9**:239-45.
6. Urusi JP, van Dyck E, van Houtte C, et al. Syphilis in Swaziland: a serological study. *Br J Vener Dis* 1981; **57**:95-9.
7. Arya OP, Nsanzumuhire H, Taber SR. Clinical, cultural and demographic aspects of gonorrhoea in a rural community in Uganda. *Bull WHO* 1973; **49**:587-95.
8. Hopcraft M, Verhagen AR, Ngigi S, Haga ACA. Genital infections in developing countries: experience in a family planning clinic. *Bull WHO* 1973; **48**:581-6.
9. Nsanze H, Waigwa SRN, Mirza N, Plummer F, Roelants P, Piot P. Chlamydial infections in selected populations in Kenya. In: Mårdh P-A, et al, eds. *Chlamydial infections*. Amsterdam: Elsevier Biomedical Press, 1982;421-4.
10. Osoba AO, Onifade A. Venereal diseases among pregnant women in Nigeria. *West Afr Med J* 1973; **22**:23-5.
11. Méheus A, Friedman F, van Dyck E, Guyver T. Genital infections in prenatal and family planning attendants in Swaziland. *East Afr Med J* 1980; **57**:212-7.
12. Nasah BT, Nguematcha R, Eyong M, Godwin S. Gonorrhoea, trichomonas and candida among gravid and non-gravid women in Cameroon. *Int J Gynaecol Obstet* 1980; **18**:48-52.
13. Ripa KL, Mårdh P-A. New simplified culture techniques for *Chlamydia trachomatis*. In: Hobson D, Holmes KK, eds. *Non-gonococcal urethritis and related infections*. Washington DC: American Society for Microbiology, 1977;323-7.
14. Treharne JD, Darougar S, Jones BR. Modification of the microimmunofluorescence test to provide a routine serodiagnostic test for chlamydial infection. *J Clin Pathol* 1977; **30**:510-7.
15. Forsey T, Darougar S. Indirect immunofluorescence test for detecting type-specific antibodies to herpes simplex virus. *J Clin Pathol* 1980; **33**:171-6.
16. Hira PR. Observations on *Trichomonas vaginalis* infections in Zambia. *J Hyg Epidemiol Microbiol Immunol (Praha)* 1977; **21**:215-24.
17. Hoosen AA, Ross SM, Mulla MJ, Patel M. The incidence of selected vaginal infections among pregnant urban blacks. *S Afr Med J* 1981; **59**:827-9.
18. Bramley M. Study of female babies of women entering confinement with vaginal trichomoniasis. *Br J Vener Dis* 1976; **52**:58-62.
19. Mirza NB, Nsanze H, D'Costa LJ, Piot P. Microbiology of vaginal discharge in Nairobi, Kenya. *Br J Vener Dis* 1983; **59**:186-8.
20. Kozinn PJ, Taschdjian CL, Wiener H. Incidence and pathogenesis of neonatal candidiasis. *Pediatrics* 1958; **21**:421-9.
21. Finlayson MH, Gibbs B, Brede HD. Diagnosis and incidence of *Neisseria gonorrhoeae* in cape coloured females in the Western Cape. *S Afr Med J* 1974; **48**:259-60.
22. Widy-Wirski R, D'Costa LJ. Maladies transmises par voie sexuelle dans une population rurale en Centrafrique. In: *Rapport final, 13th Conference Technique, Yaoundé*. OCEAC 1980:651.
23. Ballard RC, Fehler HG, Duncan MO, Van det Wat IJ. Urethritis and associated infections in Johannesburg—the role of *Chlamydia trachomatis*. *Southern African Journal of Sexually Transmitted Diseases* 1981; **1**:24-6.
24. Forsey T, Darougar S, Dines RJ, Wright DJM, Friedmann PS. Chlamydial genital infection in Addis Ababa, Ethiopia. *Br J Vener Dis* 1982; **58**:370-3.
25. Darougar S, Forsey T, Osoba AO, Dines RJ, Adelusi B, Coker GO. Chlamydial genital infection in Ibadan, Nigeria: a sero-epidemiological survey. *Br J Vener Dis* 1982; **58**:366-9.
26. Heyman DL. Etudes sur la gonococcie réalisées par des étudiants en médecine au Cameroun. In: *Rapport final, 13th Conference Technique, Yaoundé*. OCEAC. 1980:661.
27. Sowa S, Sowa J, Collier LH. Investigation of neonatal conjunctivitis in The Gambia. *Lancet* 1966; **ii**:243-7.
28. Sowa S, Sowa J, Collier LH, Blyth W. Trachoma and allied infections in a Gambian village. *MRC Special Report Series* 1965; No 308.
29. Mabey DCW, Whittle HC. Genital and neonatal chlamydial infections in a trachoma endemic area. *Lancet* 1982; **ii**:300-1.
30. Méheus A, Delgadillo R, Widy-Wirski R, Piot P. Chlamydial ophthalmia neonatorum in Central Africa. *Lancet* 1982; **ii**:882.
31. Weström L, Bengtsson LP, Mårdh P-A. Incidence, trends, and risks of ectopic pregnancy in a population of women. *Br Med J* 1981; **282**:15-8.
32. Muir DG, Belsey MA. Pelvic inflammatory disease and its consequences in the developing world. *Am J Obstet Gynecol* 1980; **138**:913-28.
33. Grech ES, Everrett JV, Mukasa F. Epidemiological aspects of acute pelvic inflammatory disease in Uganda. *Trop Doct* 1973; **3**:213-217.
34. Carty MJ, Nzioki JM, Verhagen AR. The role of the gonococcus in acute pelvic inflammatory disease in Nairobi. *East Afr Med J* 1972; **49**:376-9.
35. Ratnam AV, Din SN, Chatterjee TK. Gonococcal infection in women with pelvic inflammatory disease in Lusaka, Zambia. *Am J Obstet Gynecol* 1980; **138**:965-8.
36. Mårdh P-A. An overview of infectious agents of salpingitis, their biology, and recent advances in methods of detection. *Am J Obstet Gynecol* 1980; **138**:933-51.
37. Weström L. The effect of pelvic inflammatory disease on fertility. *Am J Obstet Gynecol* 1975; **121**:707-13.
38. Arya OP, Taber SR, Nsanze H. Gonorrhoea and female infertility in rural Uganda. *Am J Obstet Gynecol* 1980; **138**:929-32.
39. Armagnac C, Retel-Laurentin A. Relations between fertility, birth intervals, foetal mortality and maternal health in Upper Volta. *Population Studies* 1981; **35**:217-34.
40. Griffith HB. Gonorrhoea and fertility in Uganda. *Eugenics Review* 1963; **55**:103-8.
41. Billewicz WZ, McGregor IA. The demography of two West African (Gambian) villages, 1951-1975. *J Biosoc Sci* 1981; **13**:219-40.
42. Rampen F. Venereal syphilis in tropical Africa. *Br J Vener Dis* 1978; **54**:364-8.
43. Idsoe O, Kiraly K, Causse G. Venereal disease and treponematoses—the epidemiological situation and WHO's control programme. *WHO Chron* 1973; **27**:410-7.
44. Dogliotti M. The incidence of syphilis in the Bantu: survey of 587 cases from Baragwanath hospital. *S Afr Med J* 1971; **45**:8-10.
45. McFadzean JA, McCourt JF, Wilkison AE. Treponematoses in Gambia, West Africa. *Trans R Soc Trop Med Hyg* 1956; **51**:169-81.
46. Nsanze H, Fast MV, D'Costa LJ, Tukei P, Curran J, Ronald A. Genital ulcers in Kenya: clinical and laboratory study. *Br J Vener Dis* 1981; **57**:378-81.
47. Méheus A, van Dyck E, Urusi JP, Ballard RC, Piot P. Etiology of genital ulcerations in Swaziland. *Sex Transm Dis* 1983; **10**:33-5.
48. Sogbetun AO, Montefiore D, Anong CN. Herpes virus hominis antibodies among children and young adults in Ibadan. *Br J Vener Dis* 1976; **55**:44-7.