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Systematic Review

Epidemiology and control of trachoma: systematic review

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Summary

Trachoma is the commonest infectious cause of blindness. Recurrent episodes of infection with serovars A–C of Chlamydia trachomatis cause conjunctival inflammation in children who go on to develop scarring and blindness as adults. It was estimated that in 2002 at least 1.3 million people were blind from trachoma, and currently 40 million people are thought to have active disease and 8.2 million to have trichiasis. The disease is largely found in poor, rural communities in developing countries, particularly in sub-Saharan Africa. The WHO promotes trachoma control through a multifaceted approach involving surgery, mass antibiotic distribution, encouraging facial cleanliness and environmental improvements. This has been associated with significant reductions in the prevalence of active disease over the past 20 years, but there remain a large number of people with trichiasis who are at risk of blindness.

keywords trachoma, review, Chlamydia trachomatis, epidemiology, control

Introduction

Trachoma is the leading infectious cause of blindness worldwide. It is caused by infection with Chlamydia trachomatis and is characterised by inflammatory changes in the conjunctiva in children with subsequent scarring, corneal opacity and blindness in adults. The World Health Organization (WHO) estimated in 2002 that 1.3 million people were blind from trachoma and 40 million people are thought to have active disease and 8.2 million to have trichiasis. The disease is largely found in poor, rural communities in developing countries, particularly in sub-Saharan Africa. The WHO promotes trachoma control through a multifaceted approach involving surgery, mass antibiotic distribution, encouraging facial cleanliness and environmental improvements. Where control measures have been implemented encouraging reductions in the prevalence of trachoma have been found.

Historical perspective

The earliest references to trachoma come from China in the 27th century BC (Al-Rifai 1988). Features of trachoma were also described in the Ebers papyrus from Egypt, 15th century BC, and epilation forceps discovered in tombs from the 19th century BC (Maccallan 1931, Hirschberg 1982). Trachoma became a major public health problem in Europe at the beginning of the 19th century, when the disease was believed to have been brought back by troops returning from the Napoleonic wars in Egypt. So great was the burden of the disease at that time that many of the major ophthalmic hospitals founded in the 19th century were established to treat trachoma, including Moorfields Eye Hospital and Massachusetts Eye and Ear Infirmary. By the end of the 19th century, immigrants to the United
States were routinely screened for trachoma and sent home if they had signs of the disease. Trachoma has now disappeared from developed countries (with the exception of Aboriginal communities in outback Australia (Tellis et al. 2007), probably as a result of general improvements in living and hygiene standards.

**Clinical features and natural history**

Trachoma is a chronic keratoconjunctivitis caused by recurrent infection with serovars A, B, Ba and C of *C. trachomatis*. Infection is most commonly found in children. With repeated reinfection, some people go on to develop scarring complications and blindness in later life. The clinical manifestations of trachoma are subdivided into those associated with ‘active’ disease, usually seen in childhood, and those associated the cicatricial or scarring complications, seen in late childhood and adults (Figure 1). Active disease is characterised by recurrent episodes of chronic, follicular conjunctivitis. Follicles are subepithelial collections of lymphoid cells and appear as small, yellow-white elevations on the conjunctiva of the everted upper lid. Papillary hypertrophy (engorgement of small vessels with surrounding oedema) also occurs and can obscure the deep tarsal vessels if severe enough. Vascular infiltration of the upper cornea (pannus) may also develop in active disease, but this rarely affects vision. Individuals are frequently asymptomatic or have only mild symptoms even if marked signs of inflammation are evident. If present, symptoms are similar to those associated with any chronic conjunctivitis: redness, discomfort, tearing, photophobia and scant muco-purulent discharge. Conjunctival follicles at the upper margin of the cornea leave shallow depressions after they resolve known as ‘Herbert’s pits’ which, unlike follicles and papillae, are a pathognomonic sign of trachoma.

Repeated and prolonged episodes of infection and inflammation can result in the scarring complications of trachoma. Initially, conjunctival scarring is seen in the subtarsal conjunctiva, which can range from a few linear or stellate scars to thick, distorting bands of fibrosis. Contraction of this scar tissue causes entropion (in-turning of the eyelids) and trichiasis (eyelashes touching the eyeball) which is often painful. Eventually, corneal opacification develops the blinding end-stage of the disease. This is probably a result of multiple insults to the cornea: mechanical trauma from lashes, secondary bacterial or fungal infection and a dry ocular surface.

Over the years, various grading systems for trachoma have been proposed. The one which is currently used by trachoma control programmes is the 1987 WHO simplified grading system (Table 1) (Thylefors et al. 1987).

The prevalence of active disease is highest in pre-school children and declines to low levels in adulthood (Dawson et al. 1976; West et al. 1991b; Dolin et al. 1998). This parallels the distribution of *C. trachomatis* infection, with up to half of the community bacterial load being found in children under the age of 1 year in some studies (Solomon et al. 2003; Melese et al. 2004b). Adult bacterial loads are usually lower than those of children, and the duration of infection and disease also declines with age, presumably as the result of an acquired immune response (Bailey et al. 1999; Grassly et al. 2008). This is in contrast to the scarring features of trachoma, the prevalence of which

![Figure 1](http://example.com/figure1.png)

**Figure 1** Clinical features of trachoma. (a) Active trachoma in a child, characterised by a mixed papillary (TI) and follicular response (TF). (b) Tarsal conjunctival scarring (TS). (c) Entropion and trichiasis (TT). (d) Blinding corneal opacification (CO) with entropion and trichiasis (TT).
increase with age, reflecting the cumulative nature of the damage. Where the prevalence of active disease is very high, cicatricial complications may be seen at an early age; trichiasis was reported in 2–3% of children under the age of 15 years in southern Sudan where the prevalence of active disease was 70–80% (Ngondi et al. 2006a; King et al. 2008).

Cohort studies in trachoma-endemic communities in The Gambia and Tanzania have looked at the progression of the scarring process:

- Worsening of conjunctival scarring was seen in nearly 50% of scarred subjects over 5 years (Tanzania) (Wolle et al. 2001).
- Progression from conjunctival scarring to trichiasis was seen in 10% after 7 years and 6% after 12 years (Tanzania and The Gambia) (Munoz et al. 1999; Bowman et al. 2001).
- Minor trichiasis (<5 lashes touching the eye) progressed to major trichiasis (five or more lashes touching the eye) in 33% after 1 year and in 37% after 4 years; and unilateral progressed to bilateral trichiasis in 46% after 1 year (The Gambia) (Bowman et al. 2002b; Burton et al. 2006).
- Trichiasis is associated with the development of corneal scarring: 8% of people with trichiasis developed incidental corneal scarring after 4 years, and there was worsening of established corneal scarring in 34% after 1 year (The Gambia) (Bowman et al. 2002b; Burton et al. 2006).

The first study from Tanzania had a standardised, prospective design but the others did not. There is considerable variation in the reported rates of progression, which may reflect both variation in progression rates in different populations and methodology. A key determinant of the rate of disease progression is probably the burden of *C. trachomatis* infection in a community over time, although the direct evidence for this is limited. Several studies found that the risk of developing scarring complications is greater in those with recurrent or persistent severe inflammatory trachoma (Dawson et al. 1990; Munoz et al. 1999; West et al. 2001; Burton et al. 2006).

**Table 1** 1987 WHO simplified trachoma grading (Thylefors et al. 1987)

<table>
<thead>
<tr>
<th>Grade</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Trachomatous inflammation – Follicular</td>
<td>TF The presence of five or more follicles (each &gt;0.5 mm in diameter) in the upper tarsal conjunctiva</td>
</tr>
<tr>
<td>Trachomatous inflammation – Intense</td>
<td>TI Pronounced inflammatory thickening of the tarsal conjunctiva that obscures more than half of the deep normal vessels</td>
</tr>
<tr>
<td>Trachomatous scarring</td>
<td>TS The presence of scarring in the tarsal conjunctiva</td>
</tr>
<tr>
<td>Trachomatous trichiasis</td>
<td>TT At least one lash rubs on the eyeball</td>
</tr>
<tr>
<td>Corneal opacity</td>
<td>CO Easily visible corneal opacity over the pupil</td>
</tr>
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Infection vs. disease

There is little doubt that *C. trachomatis* is the cause of trachoma; Koch’s postulates were largely fulfilled shortly after the first isolation of *C. trachomatis* in 1957 (Tang et al. 1957; Collier et al. 1958). However, *C. trachomatis* cannot be detected in all cases of active disease, even using highly sensitive nucleic acid amplification tests (NAAT) (Baral et al. 1999; Lietman et al. 2000; Burton et al. 2003; Miller et al. 2004b). In low prevalence communities, especially those that have received mass antibiotic treatment, *C. trachomatis* is only found in a minority of those with active disease. Those with intense trachomatous inflammation are more likely to be infected and have higher bacterial loads than those with follicular disease (Burton et al. 2003; Solomon et al. 2004b; Wright & Taylor 2005). In endemic communities infection is sometimes detected in those who do not fulfil the WHO criteria for active disease. Part of the explanation for this poor correlation is likely to be the kinetics of the disease with a short latent phase (infection before clinical signs with the incubation period for disease), a patent phase (infection and clinical signs) and a recovery phase (infection cleared but clinical signs persist, which can last for many months) (Bailey et al. 1994; Wright et al. 2008). The mismatch between the presence of infection and clinical findings is also partly explained by use of the simplified WHO grading system, which excludes those with fewer than five follicles in the subtarsal conjunctiva (Ward et al. 1990).

Transmission of *Chlamydia trachomatis* infection

*Chlamydia trachomatis* is probably transmitted between individuals by a variety of mechanisms, including:

- Direct spread from eye to eye during close contact such as during play or sleep.
- Spread of infected ocular or nasal secretions on fingers.
- Indirect spread by fomites such as infected face-cloths.
• Transmission by eye-seeking flies.
• Possible spread from nasopharyngeal infection by aerosol.

A combination of these and other transmission mechanisms probably operates in most environments, although their relative importance may vary. For example, in some environments eye-seeking flies probably contribute to the transmission of infection. *Chlamydia trachomatis* has been detected by polymerase chain reaction in around 20% of *Musca sorbens* caught on the faces of children in Ethiopia (Jones 1975; Miller et al. 2004a; Lee et al. 2007) and intervention trials to reduce fly density have been associated with a reduction in active trachoma in The Gambia (Emerson et al. 1999, 2004). However, in other locations, the density of eye-seeking flies is insignificant and does not appear to contribute towards transmission (Taylor et al. 1985). Genital strains of *C. trachomatis* do not cause endemic trachoma, although occasionally they cause a self-limiting conjunctivitis (Brunham et al. 1990).

Trachoma is a focal disease and has been found to cluster at the level of the community, the household and within bedrooms, reflecting the infectious nature of the disease and suggesting that prolonged intimate contact is necessary for the transmission of infection (Dawson et al. 1976; Katz et al. 1988; Bailey et al. 1989; West et al. 1991b; Burton et al. 2003). This is particularly important for trachoma control programmes, as it significantly increases the sample size necessary for estimating the prevalence within a region (Katz et al. 1988). Most transmission events occur within the household, and a failure to treat all infected household members during mass antibiotic distribution may result in rapid re-infection of that family followed by more gradual spread across the community (Blake et al. 2009).

No non-human reservoir of infection has been found, with flies only acting as passive vectors. The importance of extra-ocular sites of infection has been debated. *Chlamydia trachomatis* can be detected in secretions from the nasopharynx, and a recent study also showed that infected nasal discharge in children at baseline was associated with an increased risk of active disease and conjunctival infection 2 months after systemic treatment (Malaty et al. 1981; West et al. 1993; Gower et al. 2006). However, nasal swabs were taken only from children with visible discharge and were of the discharge rather than from nasal epithelium. Positive results may simply have been a reflection of severe ocular infection which was not cleared with one dose of antibiotic, with infected secretions passing through the nasolacrimal ducts. An earlier study using nasal swabs on all children showed that new ocular infection after treatment was not related to a positive or negative nasal specimen at baseline (West et al. 1993). In addition, genotyping of conjunctival and nasal samples from individuals with concurrent infection showed different genotypes to be present, suggesting that auto-infection was not an important factor (Andreasen et al. 2008).

**Prevalence and geographical distribution**

Trachoma is a major cause of blindness in many less-developed countries, especially in poor, rural areas. Blinding trachoma is believed to be endemic in over 50 countries, with the highest prevalence of active disease and trichiasis in Africa, predominantly in the savannah areas of East and Central Africa and the Sahel of West Africa (Figure 2). It is also endemic in a number of countries in the Middle East, Asia, Latin America and the Western Pacific (Polack et al. 2005). Current WHO estimates for the prevalence of active disease, trichiasis and blindness are significantly lower than previous ones and declines in the prevalence have been noted in several countries, but there is considerable uncertainty around these estimates, as little recent information is available from India and China.

About half of the global burden of active trachoma is concentrated in five countries: Ethiopia, India, Nigeria, Sudan and Guinea; while half of the global burden of trichiasis is concentrated in three countries: China, Ethiopia and Sudan (Mariotti et al. 2009). Recent studies from southern Sudan, previously inaccessible during the civil war, have shown very high levels of trachoma: up to 80% of children had active disease and one-fifth of adults had trichiasis (Ngondi et al. 2006a; King et al. 2008). Trachoma was shown to account for 35% of blindness, with 5% of the entire population (including children) suffering from low vision or blindness associated with trachoma (Ngondi et al. 2006b, 2007).

Some caution is required in the interpretation of global estimates of trachoma prevalence (Burton & Mabey 2009). These have generally been produced with models that have relied on the results of a limited number of surveys conducted in a few endemic countries. Various assumptions and extrapolations are then made, which have considerable potential for error, such as extrapolating data from a single survey within a district to give the district-level prevalence, and national averages being generated from available district prevalence data. The six million people estimated by the WHO to be blind from trachoma in the 1990s was probably a substantial overestimate as results were based on questionnaires reporting numbers of people who might become blind without treatment (Thylefors et al. 1995). More recent estimates have used more reliable survey data.
Notwithstanding the aforementioned limitations of the available data, there does appear to be a downward trend in the number of people affected by trachoma. Improved living standards in many countries probably account for at least part of this trend, as was the case with the disappearance of trachoma from industrialised countries a century ago (Dolin et al. 1997; Hoechsmann et al. 2001). The establishment of trachoma control programmes has probably played a major role, although this is difficult to quantify. Worryingly, the number of people estimated to have trichiasis has shown little decline since 1991, with a slight increase estimated between 2003 and 2008. This suggests that progressive conjunctival scarring can occur even when there has been a marked reduction in active disease and \textit{C. trachomatis} infection, which has long-term implications for control programmes.

The most recent estimate from the WHO places the burden of trachoma at 1.3 million disability-adjusted life years. This measures the gap between a normal, healthy population and the ‘cost’ of a disease from premature mortality and disability (WHO 2008). The economic cost of trachoma has been estimated at between US$ 3 billion – 8 billion in lost productivity (Frick et al. 2003a,b). Estimates of the global burden of trachoma, however, are faced with several problems including a lack of robust prevalence data and the decision over inclusion of different disease manifestations (Burton & Mabey 2009). Trichiasis without visual impairment, for example, causes a level of disability comparable to that caused by visual impairment from non-trachomatous causes, yet it has not always been included in disease burden calculations (Frick et al. 2001b).

**Risk factors for trachoma**

Many studies have examined potential risk factors for trachoma, which have been previously reviewed (Emerson et al. 2000; West 2004; Haylor 2008). Studies examining the relationship between trachoma and various environmental, socio-economic and behavioural factors are difficult to interpret as they often lack adequate controls and are potentially confounded with many factors being closely interrelated. For example, establishing what contribution a dirty face makes to trachoma, or vice versa, is difficult, as active disease may cause ocular/nasal discharge, but discharge may be an important route for transmission. In addition, variability in survey methodology and questionnaires may not allow reliable comparisons between studies (Emerson et al. 2000).

Trachoma is currently more common in dry areas, and the relationship between water and trachoma has been studied in several settings, with some conflicting results. It is plausible that better access to water would improve hygiene levels and reduce the transmission of infection. Several studies have indeed found an association between increased distance to water and the prevalence of active disease (Mathur & Sharma 1970; Tielsch et al. 1988; Taylor et al. 1989; West et al. 1989; Schemann et al. 2002). However, other studies have not supported this and the association appears to be absent when the distance to water is small (West et al. 1991b; Zerihun 1997; Kuper et al. 2003). This may be explained by the presence of a ‘water use plateau’ in which per capita water consumption between households often seems to be constant when the round trip to collect water is below a threshold of around 30 min (Cairncross & Feachem 1993). The quantity of water brought into a household may be more important than the distance to water. Indeed, one study found the quantity to be independent of distance and that children from households with a greater quantity of water had less active disease (Kupka et al. 1968). However, other studies have shown that after controlling for distance the total quantity of water used had no effect on the prevalence of disease (West et al. 1989; Bailey et al. 1991). The second
of these two studies may unlock the key issue with regard to water and trachoma: the authors actually measured how much water was brought into the house and also observed how the water was used. After controlling for family size, distance to water and other socio-economic factors, families with trachoma used less water for washing children than did control families without trachoma, regardless of the amount of water available for consumption (Bailey et al. 1991).

The association between frequent face washing and reduced trachoma has been reported in some, but not all, studies (Taylor et al. 1985; Tielsch et al. 1988; Bailey et al. 1991; Luna et al. 1992). Self-reporting may have compromised the results, as washing may be perceived as a desirable activity and hence over-reported. A large-scale randomised trial of an intensive educational intervention to encourage face washing in Tanzania showed that children with a clean face were less likely to have severe inflammatory trachoma (TI). However, there was no reduction in the overall prevalence of active trachoma and intensive behavioural intervention was required (West et al. 1991a, 1995; Schemann et al. 2002).

As discussed previously, flies are also a risk factor for trachoma by facilitating transmission. M. sorbens, the fly most commonly found in contact with eyes, preferentially breeds in human faeces. Latrine access is associated with a lower risk of trachoma. This has been attributed to the removal of faecal material from the environment leading to a smaller fly population (Emerson et al. 2004).

Crowding is probably a risk factor for trachoma, especially living in close proximity to children with active disease (Bailey et al. 1989; Sahl & Larson 1992). Women tend to have a higher rate of the scarring complications of trachoma and this is generally considered to be a result of their increased contact with young children, the main reservoir of infection (Turner et al. 1993). Migration between communities may also be important in the re-introduction of C. trachomatis (Burton et al. 2005b).

Assessing the burden of trachoma

Trachoma as a public health problem is defined by the WHO as a prevalence of TF of at least 10% in children aged 1–9, or a prevalence of TT of at least 1% in those aged 15 or more. Trachoma is no longer considered a public health problem when the TF prevalence in children falls below 5% and the prevalence of TT is <0.1% (WHO, 1997; Kuper et al. 2003). No specific guidelines are provided for areas where the prevalence falls between these thresholds.

Population-based prevalence surveys (PBPS)

To determine where trachoma is a public health problem, WHO recommends cluster random sampling (Ngondi et al. 2009b). Districts likely to be trachoma-endemic are identified using information from previous surveys, written reports, hospital eye surgery records and interviewing people with local experience. A list of all clusters within the districts identified is made. Clusters are preferably areas of approximately the same population size, so that the cluster selection is with probability of selection proportional to size. A random sample of clusters is then selected, which is sufficiently large such that the sample prevalence of TF in 1–9 year olds, or TT in those aged 15 or more, reflects the prevalence in the whole population (WHO, 2006). A two-stage design can be employed, whereby villages (clusters) are selected in the first stage, and households are selected in the second. If household lists are not available, other methods for selecting households are by random walk and compact segment sampling. Reports should present standardisation of the examiners’ grading, the sample size parameters, confidence intervals of the estimate, and adjustment for clustering (Ngondi et al. 2009b). As well as obtaining accurate estimates of TF and TT prevalence, surveys should collect data on the number of public access and surface water points in the district, and the proportion of households that have access to latrines and that are within 15 min walk of the nearest water source available during the dry season. These data allow planning, monitoring and evaluation of control interventions (WHO, 2006).

Population-based prevalence surveys provide comprehensive prevalence data and are rightly considered the ‘gold standard’ for trachoma surveys (Wright et al. 2005). Although they can be designed to provide precise prevalence estimates over wide areas, they generally do not give accurate estimates at the cluster level, and the sampling needs to incorporate large design effects (four or more) arising from the focal nature of active trachoma and use large numbers of clusters if they are not to overlook hyperendemic clusters of disease (WHO-ITI, 2004). Moreover, they are time consuming and expensive because of the large sample sizes needed. Two alternative methods have been proposed: trachoma rapid assessment (TRA) and acceptance sampling TRA (ASTRA).

Trachoma rapid assessment

Trachoma rapid assessment was designed to allow simple, fast and cost-effective assessment of active disease, trichiasis and environmental risk factors. Existing data are first used to identify areas that are likely to be
trachoma-endemic. The burden of trichiasis, active disease and associated risk factors is then assessed in these areas (Negrel & Mariotti 1999). At least three, but no more than seven, villages are selected per district, with priority given to those areas ‘deemed most socio-economically disadvantaged’ (Wright et al. 2005). In these communities, individuals with TT are identified, leading to a crude estimate of TT prevalence. Fifty children aged 1–9 from at least 15 households that ‘appear to have the lowest socio-economic status’ are then assessed for TF and/or TI. Finally, a survey is performed to determine household level trachoma risk factors.

Trachoma rapid assessment provides rankings rather than prevalence estimates, and the method of selection of areas, communities and households outlined previously will generally be subjective. This may lead to overestimated and/or inconsistent prevalence data, with the possible extrapolation of biased data to the whole village and district (Negrel et al. 2001; Myatt et al. 2003; Solomon et al. 2004b). Evaluations of TRA rankings in comparison to PBPS in Tanzania and China found comparable ranking of communities, but TRA performed worse in low prevalence settings (Paxton 2001; Liu et al. 2002). However, PBPS does not itself provide reliable estimates or rankings for individual clusters, so these comparisons are flawed (Ngondi et al. 2009b). In The Gambia, a study comparing two TRA surveys found that active disease prevalence estimates and rankings were inconsistent, indicating that it is not a reliable method (Limburg et al. 2001).

Acceptance sampling TRA (ASTRA)

Acceptance sampling TRA, based on the principle of sequential sampling methods, such as lot quality assurance sampling (LQAS), has been proposed as an alternative to TRA. A maximum sample size and an acceptable number of TF cases are set and sampling stops when one of these is met. Villages are classified as high prevalence if sampling is stopped because the set number of TF cases was exceeded, or as low prevalence where sampling is stopped because the maximum sample size was reached (Myatt et al. 2003). Thus, there is no fixed sample size. ASTRA was evaluated in Malawi (Myatt et al. 2003) and Vietnam (Myatt et al. 2005) and found to be more reliable than TRA for the prioritisation of communities with active disease.

The advantages of ASTRA are its speed and low cost as a result of smaller sample sizes than are required for PBPS. Sample sizes may, however, become large if the option of continuing sampling in a lot until the maximum sample size is met is taken, rather than stopping when the expected number of TF cases is met (Ngondi et al. 2009b). ASTRA may provide reliable TF prevalence estimates in individual communities so long as the sample size is not too small and, if combined with Centric Systematic Area Sampling, may provide prevalence estimates over wide areas and basic mapping of TF prevalence (WHO-ITI, 2004). However, LQAS sampling works best when the distribution of cases is homogeneous (Anker et al. 1998), and when village populations do not vary too much. Because trachoma clusters both within communities and districts, an optimal rapid and affordable strategy to take clustering into account when choosing households and communities to sample is still a challenge. In the mean time, PBPS remain the only reliable source of prevalence data for trachoma, and have generally been used to prepare national control plans, and to forecast ultimate intervention goals for surgery and antibiotic treatment.

Clinical signs versus infection

An additional concern with all of these survey methods is their reliance on clinical signs as a measure of trachoma prevalence. As mentioned previously, clinical signs are sometimes poorly correlated with ocular C. trachomatis infection, especially in low prevalence communities and those that have received mass treatment. As three of the four components of the WHO endorsed SAFE strategy for the control of trachoma aim to interrupt transmission of the bacteria, logic dictates that control measures should be directed to areas with most infection. It has been suggested that NAAT testing should be used to assess the prevalence of ocular C. trachomatis infection in areas where the prevalence of TF is <10% or between 10% and 20% (Lansingh & Carter 2007). NAAT testing is not considered necessary in higher prevalence areas as the correlation between disease and infection is more reliable (Lansingh & Carter 2007). However, NAAT testing is beyond the budget of most trachoma control programmes, although cost savings can be made by pooling samples from low prevalence communities (Diamant et al. 2001). A simple point of care test for C. trachomatis showed promise when evaluated in trachoma-endemic communities in Tanzania (Michel et al. 2006) but it is not yet commercially available. Its sensitivity and specificity were lower when evaluated in subsequent, larger studies in The Gambia and Senegal (article in preparation).

Controlling trachoma: the SAFE strategy

Blindness from trachoma is essentially irreversible, but it can be prevented. The Alliance for the Global Elimination of Blinding Trachoma by the year 2020 (GET 2020) was established by the WHO in 1997 and recommends the SAFE strategy for trachoma control: Surgery for trichiasis;
Antibiotics to treat *C. trachomatis* infection; Facial cleanliness through personal hygiene; Environmental improvement with education and improved local economy.

### Surgery for trichiasis

The aim of surgery for trichiasis is to reduce the progression to corneal opacity and blindness as a result of lashes abrading the cornea. Surgery has been shown to improve comfort, reduce ocular discharge and improve visual acuity in major trichiasis cases (Reacher et al. 1992; Bowman et al. 2000a; Burton et al. 2005a). While trichiasis surgery has not been directly shown to reduce the progression to corneal opacity (Bowman et al. 2001, 2002b), the consensus view is that there is some protective effect. The WHO recommends regular surgical sessions at fixed sites once a week, with periodic outreach stations held in trachoma-endemic communities, and should be offered to anyone with trichiasis, regardless of the number of in-turned eyelashes (WHO, 2006).

#### What type of surgery?

Several procedures are in routine use by trachoma control programmes. These generally involve a full thickness incision through the tarsal plate combined with several everting sutures to turn the distal part of the eyelid outwards (Yorston et al. 2006). In a randomised controlled trial, the bilamellar tarsal rotation (BLTR), which also includes incision of the skin, was found to give the best results of the procedures that were compared and therefore WHO recommends this method (Reacher et al. 1992, 1993). The main alternatives in regular use are variations of the posterior lamellar tarsal rotation (PLTR), including the Trabut procedure. In the only study comparing recurrence rates, no significant difference between BLTR or PLTR was found (Adamu & Alemayehu 2002).

#### What are the challenges for surgery?

One of the major problems is high post-surgery trichiasis recurrence rates, ranging from about 20% in the first 2 years (Reacher et al. 1992; Bog et al. 1993; Zhang et al. 2004a; Merbs et al. 2005; El Toukhy et al. 2006) to 60% after 3 years (Reacher et al. 1993; Bowman et al. 2000a). Several factors may contribute to recurrent trichiasis such as the type of procedure used, the surgeon’s experience, the severity of pre-operative disease (severe scarring and entropion are associated with increased recurrence), suture type and infection status (Reacher et al. 1992; Alemayehu et al. 2004; Burton et al. 2005c; Merbs et al. 2005; El Toukhy et al. 2006). The presence of conjunctival inflammation, which may reflect ongoing inflammatory-cicatricial responses, has been observed in patients with trichiasis and conjunctival scarring and may be important in the process of recurrent trichiasis. It is unclear what is driving this process as infection with *C. trachomatis* is relatively uncommon and has not been associated with recurrent trichiasis (Burton et al. 2005c; West et al. 2006c). Other bacteria (non-chlamydial) are commonly associated with trichiasis and so may contribute to inflammation in the late stages of the disease (Burton et al. 2005c). To explore whether controlling infection improved results, three randomised trials of post-operative azithromycin have been conducted. These have given different results. No effect was found in a low-prevalence Gambian setting (Burton et al. 2005c), reduced recurrence was observed in a high-prevalence Ethiopian settings (West et al. 2007a), and reduced recurrence was observed for major trichiasis, but increased recurrence for minor trichiasis, in a medium-prevalence area of Nepal (Zhang et al. 2006).

In many settings, the up-take of surgical services by patients has been relatively low. Patient barriers include cost, fear of surgery, transport difficulties, need for an escort, lack of awareness about the need for treatment or how to access care (Courtright 1994; West et al. 1994; Bowman et al. 2002a; Melese et al. 2004a; Habte et al. 2008). It has been shown that community-based surgery has greater attendance rates (66%) than health centre-based surgery (44%) (Bowman et al. 2000c). Surgery is therefore most successful when performed within the community by a trained nurse, with little or no cost to the patient (Mabey et al. 2003). Provider-level barriers include lack of training, auditing, availability of sterilised equipment and supplies and surgeons. To increase the number of surgeons, ophthalmic nurses can be successfully trained (Alemayehu et al. 2004). Case finding is of crucial importance and is facilitated by having individuals living in endemic communities trained to recognise trichiasis and refer cases (WHO, 2006).

### Non-surgical alternatives

In many trachoma-endemic regions, epilation of the eyelashes is commonly practised with home-made equipment. For mild trichiasis with a few peripheral lashes in the absence of significant entropion, this may be a reasonable alternative to surgery; however, this has not yet been formally tested. In a cross-sectional analysis before surgery, epilation was associated with a reduced risk of corneal opacification in people with more severe entropion but made no difference for mild disease (West et al. 2006a). A retrospective study showed that epilation neither helped
nor hindered the progression process, although when combined with hot ash there was more corneal damage (Bowman et al. 2002b). Eyelid-taping has also been proposed as a non-surgical intervention, but this is generally a short-term measure prior to surgery (Yorston et al. 2006). Nevertheless, eyelid-taping alone is more effective than a single episode of epilation at keeping lashes off the eye at 3 months (Graz et al. 1999).

**Antibiotics**

The demonstration that a single oral dose of azithromycin was as effective as 6 weeks of daily tetracycline ointment in the treatment of active disease was a major advance (Bailey et al. 1993) and led directly to the launching of the global elimination initiative. Mass treatment of whole districts or communities is recommended, as this is more effective in preventing reinfection than the treatment of individual cases (Schachter et al. 1999). The WHO criteria for deciding whether or not to treat are shown in Table 2. A district is defined as a geographical area containing between 100 000 and 150 000 people.

**Which antibiotic?**

The WHO recommends two antibiotic treatment regimes: either 1% tetracycline eye ointment twice daily for 6 weeks or a single oral dose of azithromycin. Randomised controlled trials comparing these two treatments demonstrated that they are equally efficacious (Bailey et al. 1993; Tabbara et al. 1996; Dawson et al. 1997; Schachter et al. 1999) but that azithromycin is more effective in operational use (Bowman et al. 2000b). Tetracycline is almost universally available but suffers from poor compliance because of the length of administration, being difficult and unpleasant to apply, and side-effects such as stinging and blurred vision (West 1999; Kuper et al. 2003). Azithromycin is well tolerated by both adults and children, has good compliance, and has fewer side-effects than tetracycline (Schachter et al. 1999; West 1999). It is also active against extra-ocular *C. trachomatis*. A recent cluster-randomised trial in Ethiopia showed that at 12 months, there was a 50% reduction in childhood mortality in communities where children had been treated with oral azithromycin compared to those where they had not (Porco et al. 2009). Pfizer has donated 135 million doses of azithromycin for use in control programmes, distributed by the ITI. The ITI is active in 18 trachoma-endemic countries. Azithromycin dosage is based on weight for children (20 mg/kg body weight), with adults receiving 1 g. As weighing scales need daily calibration, are cumbersome to carry, and the cooperation of young children can be hard to obtain, height as a surrogate for weight has been suggested and proved successful for dosing (Munoz et al. 2003).

Azithromycin for trachoma control is not currently recommended for children under 6 months or pregnant women, and therefore tetracycline ointment is the treatment of choice for these groups. However, azithromycin is recommended by the Centre for Disease Control in infants under 1 month for pertussis prophylaxis (Tiwari et al. 2005) and is also recommended for the treatment of genital chlamydial infection in pregnant women (Gray et al. 2001; Pitsouni et al. 2007). The treatment of infants is important as infants under 1 year have the highest bacterial load, as discussed previously.

While oral azithromycin would seem to be a safe option, a potential alternative is azithromycin eye drops. A clinical trial of short duration azithromycin eye drops found that at 2 months, the cure rate and safety of topical 1.5% azithromycin was non-inferior to oral azithromycin (Cochereau et al. 2007), and mass treatment with the eye drops of a district in Cameroon saw the prevalence of active disease fall from 31.5% before treatment to 6.3% 1 year after treatment (Huguet et al. 2010).

As with any antibiotic, there are concerns that widespread use might lead to drug resistance. Azithromycin resistance has not yet been observed in *C. trachomatis* (Solomon et al. 2005; Hong et al. 2009), but resistance in other bacteria, such as *Streptococcus pneumoniae*, has been documented, especially after multiple rounds of mass treatment (Leach et al. 1997; Chern et al. 1999; Gaynor et al. 2005) although this disappeared within 12 months of treatment (Fry et al. 2002; Gaynor et al. 2003a). The clinical relevance of this resistance has yet to be determined. It has been argued that in communities where

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**Table 2 WHO criteria for mass antibiotic treatment distribution (WHO-ITI 2004)**

<table>
<thead>
<tr>
<th>Prevalence of TF in children</th>
<th>Recommendation</th>
</tr>
</thead>
<tbody>
<tr>
<td>District level</td>
<td></td>
</tr>
<tr>
<td>≥10%</td>
<td>Mass treat whole district annually for 3 years, then re-assess the prevalence in the district</td>
</tr>
<tr>
<td>&lt;10%</td>
<td>Do community-level assessment</td>
</tr>
<tr>
<td>Community level</td>
<td></td>
</tr>
<tr>
<td>≥10%</td>
<td>Mass treat whole community annually for 3 years, then re-assess the prevalence in the community</td>
</tr>
<tr>
<td>≥5% but &lt;10%</td>
<td>Target treatment to affected children and the household they live in</td>
</tr>
<tr>
<td>&lt;5%</td>
<td>Antibiotic treatment not recommended</td>
</tr>
</tbody>
</table>
macrolide resistance is rare, mass treatment with azithromycin is unlikely to increase the prevalence of resistant 
*S. pneumoniae* (Batt et al. 2003). Nasopharyngeal 
*S. pneumoniae* resistance to topical tetracycline has also 
been detected (Gaynor et al. 2005). The risk of drug 
resistance highlights the need for sensitive diagnostic tests, 
where treatment can be targeted to limit the over-use of 
antibiotics within the mass treatment policy (Mabey et al. 
2003).

To whom should treatment be given?
The optimal strategy of mass antibiotic treatment is subject 
to some debate and probably varies depending on the 
prevalence. Alternative treatment target groups have been 
proposed:

- All children under 10 years old, because children are 
  the main reservoir of infection (Holm et al. 2001; 
  Solomon et al. 2003, 2004a). Frequent mass treat-
  ment of all children under 11 years has shown herd 
  protection in the entire community in high-prevalence 
  settings (House et al. 2009).
- All people living in a household containing an 
  individual with active disease (Holm et al. 2001; 
  Burton et al. 2003; Blake et al. 2009).
- All people living in a community (e.g. village), where 
  the prevalence of active disease rises above a specific 
  threshold (Burton et al. 2003).
- Children with active disease and other children 
  residing with the TF/TI child more than 50% of the 
  time (Laming et al. 2000).
- All TI individuals, as they have the highest number of 
  chlamydial DNA copies per swab (Solomon et al. 
  2003).
- Only infected individuals, as this would remove the 
  source of infection from the community, but this 
  requires the means to detect infection in the field 
  (Lietman et al. 1999).

In a study from a low-prevalence setting in The Gambia 
in which the residents of 14 villages were examined and 
tested for *C. trachomatis* infection, the theoretical effec-
tiveness of several strategies in delivering antibiotic to 
infected individuals was compared (Burton et al. 2003). If 
only the individuals with active trachoma were treated, 
then only 24% of infected people would receive antibi-
otic. If treatment was targeted to all the residents of a 
household where at least one case of active trachoma was 
found, then 96% of infections would be treated. How-
ever, the number of people needed to be treated for each 
infection case was 9.6. Finally, if all villages with more 
than 15% active disease in children were treated, then 
90% of infections would have been treated and the 
number needed to treat was 5.4. Thus, for a low-to-
medium prevalence setting, community level treatment, 
determined by the prevalence of active disease in the 
children, appears to be a relatively efficient approach.

In a high-prevalence village in Tanzania, it was estimated 
that if only children under 10 years were treated, only 69% 
of those with high loads would be treated. If all members of 
households with children aged <10 years were treated, 
90% of the entire population would be treated because 
most people live in households with children. Treating only 
those with clinical signs would miss 23% of those with 
high loads. The authors concluded that it is therefore more 
practical and effective to treat the entire community, so 
long as coverage is high enough (West et al. 2005a).

Mass treatment is considered the most cost-effective 
strategy, especially in high-prevalence areas (Frick et al. 
2001a; Holm et al. 2001). Targeted treatment strategies 
require all children to be examined, which can be expensive 
and time consuming. In addition, re-infection is more likely 
to occur, as those treated may be re-infected by untreated 
individuals (West et al. 1993). Mass treatment has the 
advantage that all infected individuals are captured 
(Schachter et al. 1999; Solomon et al. 2004a; West et al. 
2005a). This is important, as clinical signs are not reliable 
as a basis for targeted treatment (Baral et al. 1999), 
asymptomatic individuals act as a reservoir of infection 
(Burton et al. 2003; West et al. 2005a) and adults can also 
act as an important reservoir of high load infection (West 
et al. 2005a).

What treatment coverage should be achieved?
WHO recommends that treatment coverage should be 
between 80% and 90% (WHO, 2004). Mathematical 
modelling assuming 80% treatment coverage and 3 years of 
anual treatment demonstrated elimination of infection in 
95% of communities (Ray et al. 2009). In contrast, data 
from Tanzania demonstrated that despite overall treatment 
coverage of 86%, ocular *C. trachomatis* infection remained 
in the community for up to 18 months after treatment, 
albeit at a low level (Burton et al. 2005b). Much of the near 
elimination of ocular *C. trachomatis* infection at 2 years 
after a single round of azithromycin mass treatment in a 
Tanzanian community has been attributed to the 97.8% 
treatment coverage (Solomon et al. 2004a), However, 
treatment coverage is not the sole key to success. In a 
low-prevalence setting in The Gambia, significant 
re-infection in two villages post-treatment was observed, 
despite treatment coverage being 86% and 92%. This was 
attributed to a mass migration event where virtually the 
entire population of these two communities attended a
religious festival in Senegal shortly after being treated (Burton et al. 2005b). In one Ethiopian study, an overall treatment coverage of 91.9% was achieved, but was followed by a 12.3% exponential rate of return of infection (Melese et al. 2004b). Also in Ethiopia, it was demonstrated that although treatment coverage was important in determining the prevalence of ocular C. trachomatis infection at 2 months post-treatment, coverage was no longer a predictor of infection at 6 months (Lakew et al. 2009a).

Factors found to affect the acceptability of azithromycin are local prevention norms (for example, believing that injections are better than oral medicine), perceptions of the distribution team’s expertise, witnessing adverse effects in others, and the timing, quality and quantity of information provided. Therefore, to maximise coverage, it is important to understand the community’s perceptions, conduct a pre-distribution assessment and community education, provide advance notice of the distribution, build a good relationship with the community, create and follow standardised distribution guidelines, and improve distributor training (Desmond et al. 2005).

How often should mass treatment be given?

It has been argued that a single round of mass treatment, with high coverage, may reduce the prevalence of infection to below a threshold at which it cannot persist, and from which it cannot return. This is known as the Allee effect (Chidambaram et al. 2005). Alternatively, mass treatment may eliminate some strains of C. trachomatis from the community, reducing the antigenic diversity which may enable the bacteria to evade the human immune system, and this less diverse population may never re-attract a high prevalence (Zhang et al. 2004b; Burton et al. 2005b; Chidambaram et al. 2006; Andreasen et al. 2008). Factors affecting the success of a single round of mass treatment are the baseline prevalence, treatment coverage, treatment efficacy in the individual, whether ‘F’ and ‘E’ component measures are in place, and the amount of in- and out-migration (Lietman et al. 1999; Gaynor et al. 2003b; Burton et al. 2005b; Chidambaram et al. 2006).

A single round of mass azithromycin treatment was successful in reducing the prevalence of ocular C. trachomatis infection from 9.5% at baseline to 0.1% at 2 years in a Tanzanian community (Solomon et al. 2004a). After a second round of mass treatment at 2 years, infection was eliminated by 5 years post-baseline (Solomon et al. 2008). This demonstrates that antibiotic treatment alone can result in elimination, as no ‘F’ or ‘E’ interventions were introduced. However, the baseline treatment coverage of 97.8% far exceeds that which would normally be achieved under operational conditions. The decline in trachoma prevalence may not have been a result solely of the mass azithromycin treatment as tetracycline eye ointment was distributed at the 6, 12 and 18 month follow-ups to individuals with active disease. However, 15–100% of the community ocular C. trachomatis load at each of the final three follow-ups was found in participants who had received tetracycline in the previous follow-up, indicating tetracycline treatment did not play a major role in the observed prevalence decline. Alternatively, random fluctuation, seasonal effects, secular trend and regression to the mean may have contributed to the outcome. In contrast, two rounds of mass azithromycin treatment (at baseline and 18 months) in a different Tanzanian village did not eliminate active disease or C. trachomatis infection 5 years post-baseline (West et al. 2007b). In fact, although the infection rate declined between baseline and 18 months, the prevalence of infection was higher at the 5-year follow-up than at 18 months in all age groups.

Several studies have reported moderate success of one round of antibiotic treatment, with infection initially falling immediately post-treatment, but increasing (albeit to a lower prevalence than at baseline) within 12 months of treatment. In Egypt, The Gambia and Tanzania, it was observed that the prevalence of infection at 1 year after mass treatment was substantially lower than at baseline, but was higher than the prevalence at the 3-month follow-up (Schachter et al. 1999). Similar results of an initial reduction in infection with re-emergence approximately 1 year after treatment, but which does not return to pretreatment levels by 2 years, have been reported by others in high-prevalence settings (Melese et al. 2004b; West et al. 2005b; Chidambaram et al. 2006; Lakew et al. 2009b). A study of 14 Gambian villages demonstrated that in low or medium prevalence areas, a single round of mass azithromycin treatment could lead to long-term control of infection, but that monitoring is required because of re-infection (Burton et al. 2003, 2005b). To help overcome the risk of re-infection from migration, it has been recommended that broader geographical areas should be treated, and people who have immigrated into the village should be treated after the initial mass treatment (Schachter et al. 1999; Burton et al. 2003b).

The aforementioned studies demonstrate that the effect of a single round of mass azithromycin treatment is heterogeneous, with some communities experiencing elimination of infection whereas others observe rapid re-emergence. In fact, more frequent treatment distributions could be beneficial in high-prevalence settings. Mathematical modelling has shown that where the prevalence of active disease is >50% in children, bi-annual treatment could eliminate disease (Lietman et al. 1999). Where disease prevalence is <35%, treatment annually or
of face washing compared mass tetracycline treatment with mass tetracycline treatment combined with a face-washing programme (West et al. 1995). Screening was performed at baseline, 6 and 12 months post-baseline. Children who received both the face-washing programme and treatment were more likely to have sustained clean faces than those who only received treatment, although the difference was not significant (OR 1.61, 95% CI 0.94–2.74). However, 65% of children in the intervention group still had a dirty face at two or more follow-ups. The risk of having severe trachoma (defined as the presence of ≥15 follicles or the presence of inflammation that obscured all tarsal plate vessels) in the face-washing group after 1 year was significantly lower than the treatment-only group (OR 0.62, 0.40–0.94), as mentioned previously. However, there was no difference in the overall prevalence of active disease between the two arms. The programme was labour intensive and expensive.

Environmental improvement

The ‘E’ component of the SAFE strategy aims to reduce transmission of C. trachomatis by promoting better personal and environmental hygiene. The elimination of trachoma from Europe and North America in the 19th century in the absence of any specific intervention, demonstrates the importance of environmental improvement components of the SAFE strategy (Mabey et al. 2003). Through increasing water supply and quality, improving access to latrines, decreasing fly density, reduced crowding and providing health education, transmission of trachoma should be interrupted (Kuper et al. 2003).

Resnikoff et al. compared health education alone, mass tetracycline treatment alone, health education combined with tetracycline treatment, and a control group (no intervention), assigning only one village to each arm (Resnikoff et al. 1995). They found that at 6 months, the incidence of active disease was lower in the health education only group compared with the control group. However, there was no value in the addition of health education to mass treatment, with mass treatment alone producing the best results in terms of cure rate and lower incidence.

There was little evidence for the fly control component of the SAFE strategy until Emerson et al. demonstrated that insecticide spraying in The Gambia led to an overall and significant 61% lower community prevalence of active disease, a reduction of 75% in the M. sorbens fly population, and a 96% reduction in fly-eye contacts in the intervention villages at 3 months (Emerson et al. 1999). However, the study only compared two pairs of villages and was open to bias. In 2004, Emerson et al. compared seven clusters that received spraying with seven that did
Insecticide spraying led to an 88% decrease in fly-eye contacts and a significant 55.8% reduction in the prevalence of active disease in the intervention clusters (Emerson et al. 2004). West et al. randomised 16 Tanzanian communities to receive a single round of mass azithromycin treatment (control group) or to receive azithromycin and frequent rounds of insecticide spraying (intervention group) (West et al. 2006b). In contrast to the previous studies, they found no difference in the prevalence of active disease at either 6 or 12 months post-baseline, or of *C. trachomatis* infection at 6 months, between intervention and control communities, despite the mean prevalence of flies being significantly lower in the intervention group. Furthermore, spraying is labour intensive, expensive and not sustainable (Rabiu et al. 2007).

Only one randomised controlled trial examining latrine use exists. Emerson et al. compared seven clusters that received latrines, with seven that did not (Emerson et al. 2004). Latrine provision resulted in a 30% decrease in *M. sorbens*-eye contacts, and an associated 29.5% reduction in trachoma prevalence, which did not reach statistical significance, despite latrine use reported to be 98%.

Latrines will only improve environmental sanitation if they are used consistently by a large proportion of the community. Therefore, latrine provision should be in accordance with what already exists and what is acceptable in the community (WHO, 2006).

**The SAFE Strategy: putting the pieces together**

Although the individual components of the SAFE strategy have demonstrated success in controlling trachoma, it is through the implementation of all four elements together that this control strategy is expected to have most success. Some studies have evaluated the combined effect of multiple components of the SAFE strategy. A cross-sectional analysis of implementation of the A, F and E components in Ethiopia demonstrated that receiving three rounds of azithromycin treatment, having a clean face, and increased face-washing frequency, were independently associated with a reduced prevalence of active disease in children (Ngondi et al. 2008). Thus, implementation of the different SAFE components would have an additive effect in trachoma control.

Implementation of the entire SAFE strategy in five Ethiopian districts showed that uptake of all components was high by the 3-year evaluation time-point (Ngondi et al. 2009a). The declines in TF, TI and unclean face prevalence in children aged 1–9 were statistically significant, and the prevalence of TT significantly decreased in three of the districts. The overall prevalence of ocular *C. trachomatis* infection at 3 years was 3.1%, but was higher in districts last treated over a year ago (4.3%) and lower in those treated recently (1.4%), suggesting on-going transmission. In Zambia, introduction of SAFE measures led to a reduction in the prevalence of total trachoma in children under 10 years from 55% at baseline to 10.6% at 2 years. The prevalence of TF fell from 24.9% to 4.5% in children, and the prevalence of TT in adults fell from 0.6% to 0.3% (Astle et al. 2006). However, in the absence of control groups, secular trend cannot be excluded as an explanation. In addition, without investigating these interventions through a randomised controlled trial, the relative impact of each component cannot be elucidated.

In Sudan, a control area was included with which to compare four areas in which the SAFE strategy had been implemented. The evaluation at 3 years showed heterogeneous uptake of interventions and results. All four intervention areas experienced declines in the prevalence of TF, TI and unclean faces. This was substantial in two of the areas, moderate in one, and non-significant in one, compared with the control. The decline in active disease was most likely attributable to antibiotic treatment and improved facial cleanliness was a result of hygiene health education combined with water provision, as the greatest trachoma declines were achieved where uptake of these activities was highest (Ngondi et al. 2006c). In Ethiopia, Cumberland et al. conducted a trial where 40 communities were randomised to either health promotion by national radio only (control); mass azithromycin treatment and radio; mass treatment, radio and information, education and communication (IEC) materials; or mass treatment, radio, IEC and community video and drama shows (Cumberland et al. 2008). The exact allocation schedule was unable to be followed, but the 3 year evaluation demonstrated a significantly reduced risk of active disease in communities given antibiotics combined with IEC (OR 0.35, 95% CI 0.13–0.89), and in communities additionally receiving video health messages (OR 0.31, 0.11–0.89). Similarly, the risk of having ocular *C. trachomatis* infection was significantly lower in these two intervention groups. Although antibiotic treatment was identified as being the most active component for all outcomes, the addition of health education was beneficial.

**Conclusion**

Blinding trachoma disappeared from Western Europe and North America at the beginning of the 20th century, yet it continues to cause an enormous burden of disease in poor rural communities in the developing world. There have been encouraging reductions in the prevalence of active disease in many countries in the past 20 years. However, a large backlog of unoperated trichiasis cases which remains
in many countries will have to be addressed by national eye care programmes if blinding trachoma is to be eliminated by 2020.

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