Progress to Eliminate Trachoma as a Public Health Problem in Amhara National Regional State, Ethiopia: Results of 152 Population-Based Surveys

Aisha E. P. Stewart,1 Mulat Zerihun,2 Demelash Gesesse,2 Berhanu Melak,2 Esheetu Sata,3 Andrew W. Nute,1 Tigest Astale,3 Tekola Endeshaw,3 Tesfaye Teferi,4 Zerihun Tadesse,3 Elizabeth Kelly Callahan,1 Melsew Chanyalew,5 Birhan Gaudie,6 Paul M. Emerson,7 Jonathan D. King,4 and Scott D. Nash1*

1The Carter Center, Atlanta, Georgia; 2The Carter Center, Bahir Dar, Ethiopia; 3The Carter Center, Addis Ababa, Ethiopia; 4International Trachoma Initiative, Addis Ababa, Ethiopia; 5Amhara National Regional Health Bureau, Bahir Dar, Ethiopia; 6Dr. Abdul Higher Eye Clinic, Bahir Dar, Ethiopia; 7World Health Organization, Geneva, Switzerland

Abstract. At baseline in 2006, Amhara National Regional State, Ethiopia, was the most trachoma-endemic region in the country. Trachoma impact surveys (TIS) were conducted in all districts between 2010 and 2015, following 3–5 years of intervention with the WHO-recommended SAFE (surgery, antibiotics, facial cleanliness, and environmental improvement) strategy. A multistage cluster random sampling design was used to estimate the district-level prevalence of trachoma. In total, 1,887 clusters in 152 districts were surveyed, from which 208,265 individuals from 66,089 households were examined for clinical signs of trachoma. The regional prevalence of trachomatous inflammation-follicular (TF) and trachomatous inflammation-intense among children aged 1–9 years was 25.9% (95% CI: 24.9–26.9) and 5.5% (95% CI: 5.2–6.0), respectively. The prevalence of trachomatous scarring and trachomatous trichiasis among adults aged ≥15 years was 12.9% (95% CI: 12.2–13.6) and 3.9% (95% CI: 3.7–4.1), respectively. Among children aged 1–9 years, 76.5% (95% CI: 75.3–77.7) presented with a clean face; 66.2% (95% CI: 64.1–68.2) of households had access to water within 30 minutes round-trip, 48.1% (95% CI: 45.5–50.6) used an improved water source, and 46.2% (95% CI: 44.8–47.5) had evidence of a used latrine. Nine districts had a prevalence of TF below the elimination threshold of 5%. In hyperendemic areas, 3–5 years of implementation of SAFE is insufficient to achieve trachoma elimination as a public health problem; additional years of SAFE and several rounds of TIS will be required before trachoma is eliminated.

INTRODUCTION

National trachoma programs conduct impact and surveillance surveys to assess the prevalence of clinical signs of trachoma and progress toward elimination as a public health problem. For trachoma, the targets for elimination as a public health problem include a prevalence of trachomatous inflammation-follicular (TF) among children aged 1–9 years of less than 5% at the health district level and a prevalence of trachomatous trichiasis (TT) unknown to the health system among the total population of less than one case per 1,000 at the health district level.1 Trachoma impact surveys (TIS) are presently conducted following 1–7 years of implementation of the WHO-endorsed surgery, antibiotics, facial cleanliness, and environmental improvement (SAFE) strategy;2 however, previous guidance, followed until roughly 2017, called for TIS following 3–5 years of SAFE implementation.3

The National Survey on Blindness, Low Vision, and Trachoma, conducted in 2006, suggested Ethiopia to be the most trachoma-endemic country, among countries with available data. Within Ethiopia, the survey demonstrated that the Amhara Region harbored the highest regional prevalence of active trachoma (TF and/or trachomatous inflammation-intense [TTI]) among children aged 1–9 years, 62.6%.4 Although trachoma interventions started in 2001 in Amhara, they were established in only four districts.5 Following the 2006 National Survey, further baseline data were collected during a 2007 zonal-level survey, which provided needed evidence to determine the zones (administrative unit below a region with about 2,000,000 population and made up of districts) that warranted SAFE intervention.

The 2007 baseline survey demonstrated that trachoma was endemic in all 10 zones in the region, each requiring full implementation of the SAFE strategy. Resultingly, SAFE interventions were gradually scaled up to all 152 districts between 2007 and 2010. Following 3–5 years of SAFE implementation, in accordance with global recommendations at the time, TIS were conducted in all districts between 2010 and 2015 to assess progress toward elimination.6,7 This article presents these aggregate TIS results, providing regional-, zonal-, and district-level summaries of progress toward trachoma elimination targets.

MATERIALS AND METHODS

Ethical considerations. Survey methodology was approved by the Ethical Review Committee of the Amhara Public Health Research Institute, Ethiopia, and Emory University Institutional Review Board under protocol 079-2006. Permission to obtain verbal informed consent and assent was granted by the review boards because of the high rate of illiteracy among the study population. Verbal informed consent to conduct a household interview was obtained from heads of households or a representative 18 years or older. Verbal informed consent and assent were obtained from all individuals screened for trachoma and recorded on the household questionnaire. Permission to conduct the survey in the selected clusters was obtained from zonal and district health officials, and the cluster, or village, leader.

Study site and time frame. The trachoma program began in Amhara in 2001 in four districts and expanded to cover 19 districts in 2003 (Figure 1). Following the 2007 zonal-level baseline survey, SAFE interventions were gradually scaled up to all 152 districts between 2007 and 2010 (Table 1). This phased approach was used for logistical reasons, as providing trachoma services to an estimated 20 million residents8 in the
region required substantial planning and resources. Areas with high zonal-level prevalence were prioritized for the early phase of scale-up, as it was assumed these areas would take longer to reach elimination thresholds. However, factors including geographic location, road access, and needs from the Regional Health Bureau also influenced the order in which SAFE interventions were rolled out to districts. Districts became eligible for TIS during different years between 2010 and 2015 because of the phased scale-up and varying baseline TF prevalence. In adherence with global guidelines, all TIS took place between 7 and 9 months after the last mass drug administration (MDA), with TIS occurring 8 months after MDA in 90 (59.2%) districts.

**Sampling methodology.** The TIS methodology used in Amhara between 2010 and 2015 evolved during that period to reflect best practices and innovation within the global trachoma program. As a result, three multistage, cluster random sampling methodologies were used. All surveys used the following: in the first stage, clusters, defined by a single gott (administrative unit comparable to a village), were selected from a geographically ordered list of all gotts using probability proportional to estimated population size. In the second stage, segments of households were selected once the field team arrived in the cluster. With the gott leader, or representative, the field team enumerated a list of development teams (official segments of about 40 households within gotts) and asked the gott leader to blindly pick the name of a segment from a hat. The gott leader or designated representative familiar with the village accompanied the field team to the randomly selected segment and during the fieldwork. All households and household members within a selected segment were eligible to participate. If a household member older than 18 years was not present or if no one was at home when the field team arrived, the team made one attempt to return to the household to conduct an interview, and, if unsuccessful, the household was skipped without replacement. Children aged 1–9 years who were absent were added to a separate, electronic list in Swift Insights, the electronic data collection system, so that the survey team could return to the household once more for grading.

**FIGURE 1.** Scale-up of SAFE interventions by district, Amhara, Ethiopia, 2001–2010. This figure appears in color at www.ajtmh.org.
| Activity                                                                 | Indicator                                                                 | 2007 | 2008 | 2009 | 2010 | 2011 | 2012 | 2013 | 2014 | 2015 | Total  |
|-------------------------------------------------------------------------|----------------------------------------------------------------------------|------|------|------|------|------|------|------|------|------|--------|
| Regional scale-up                                                       | Number of districts with SAFE                                            | 66   | 113  | 134  | 152  | 152  | 152  | 152  | 152  | 152  |        |
| Surgery: Awareness raising on the cause of TT and surgical approach to TT management; provision of TT surgical services through outreach camps and at health centers | Number of TT patients operated                                           | 45,271 | 31,561 | 35,681 | 33,021 | 39,076 | 66,766 | 44,867 | 40,450 | 71,460 | 408,153 |
| Antibiotics: Annual dose of azithromycin or tetracycline eye ointment to all eligible community members | Number of doses of antibiotic distributed*                                | 6,568,335 | 12,984,025 | 13,720,673 | 15,141,608 | 15,231,371 | 12,931,668 | 15,677,633 | 16,875,459 | 15,394,959 | 124,525,731 |
|                                                                         | % coverage                                                               | 59.4% | 116.0% | 91.9% | 93.7% | 89.1% | 95.4% | 93.0% | 100.1% | 94.4% |
| Facial cleanliness: Face washing and hygiene promotion in communities; school trachoma health curriculum | Number of villages with health education                                  | 1,447 | 2,898 | 3,432 | 3,428 | 3,427 | 3,449 | 3,459 | 3,459 | 3,459 |
|                                                                        | Number of schools with health education                                  | No school education. | 6,181 | 6,922 | 6,935 | 7,884 | 8,374 | 8,374 | 8,374 | 2,553 |
| Environmental improvement: Promotion of household latrine construction and use | Number of household latrines constructed                                 | 466,359 | 373,677 | 544,205 | 590,119 | 284,423 | 520,885 | 282,078 | 104,777 | 305,511 | 3,472,034 |

TT = trachomatous trichiasis.

* Includes azithromycin and tetracycline eye ointment.
2012–2015 TIS; however, the same number of clusters was chosen from each district, and, on average, 30 households per development team were assumed as evidence from previous surveys indicated these were rarely found to be as large as 40 households.

Among the 25 districts surveyed from 2010 to 2011, a sample size of 600 children aged 1–9 years was required to detect a prevalence of TF of 3% ± 2%, assuming 95% confidence, a design effect of two, five people per household with children aged 1–9 years representing 30% of the population, and allowing for an 8% nonresponse rate among children aged 1–9 years; 10 clusters per subdistrict with 40 households each were randomly selected. Survey weights were used to aggregate data to the district and zonal levels.19

For 120 districts surveyed from 2012 to 2015, the sample size was calculated to detect a reduction in TF among children aged 1–9 years of at least 20% from baseline. Therefore, the greatest number of participants required was calculated using an assumed prevalence of TF among children aged 1–9 years of 50% and that children aged 1–9 years comprised 35% of the population. For an effect of at least 20% with a 95% level of significance, a design effect of five,20 300 children aged 1–9 years per district were needed, requiring a sample of about 900 people of all ages per district.

For the final seven districts surveyed in 2015, a prevalence of 20% was assumed based on data from previous TIS in Amhara, where 137 of 145 districts maintained a TF prevalence greater than the elimination threshold of 5%. To detect a prevalence of 20% ± 10% from baseline with a 95% level of significance, a sample of approximately 215 children aged 1–9 years per district was estimated. This required a sample of about 615 people of all ages per district. This assumed a design effect of 3.04 and a 15% nonresponse rate, both of which had been calculated from previous TIS results from 145 districts in Amhara.

**Trachoma grading.** Before each round of TIS, trachoma graders were trained for 5–7 days, including those who had previously participated in TIS teams. Graders practiced grading at primary schools and at TT surgery camps to ensure exposure to all clinical signs of trachoma. Graders were required to pass both a written examination, identifying all the trachoma clinical signs based on the WHO-simplified grading system21 (Table 2) using a set of 50 slides, and a field-reliability examination that included TF, TI, and trachomatous scarring (TS). Interobserver reliability for the field examination was assessed against concordant assessments from three master trainers. Graders achieving ≥ 84% agreement with TF and a κ ≥ 0.7 were eligible to participate in a survey team. Graders used 2.5 × binocular magnification loupes and appropriate light for screening.

**Questionnaires and trachoma examination.** Three questionnaires were used, including 1) gott-level interview with a community leader or representative, 2) household-level interview with a household representative older than 18 years, and 3) individual-level interview and trachoma screening. Each questionnaire was translated from English to Amharic then back-translated from Amharic to English. All surveys, except the first conducted in 2010 which was paper based, collected data electronically in Amharic using Swift Insights.22 Amharic questionnaires were used rather than English versions to ensure standardization among the way questions were asked, as the data recorders spoke limited English.

**Data recorders** received 5–7 days of training to learn to use the electronic data collection system and Android tablets, and to administer questionnaires following a standardized protocol. Data recorders pilot-tested questionnaires and use of the tablets in nonsurvey gotts to ensure quality and standardization of interviews and translation, as well as the functionality and usability of individual tablets and Swift Insights. Data recorders were required to pass an examination before the start of each survey to participate.

The gott-level interview collected information about the village, including the distance the survey team had to walk to reach the cluster from where the vehicle dropped them, and the presence of a paved road, electricity, mobile phone coverage, and a health facility in the community. The global positioning system coordinates of each cluster were taken using the tablet. The household interview was conducted at each selected and consenting household to gather information on socioeconomic status, water, and sanitation, as well as hygiene practices. Latrine presence and use were verified by observation by the data recorders.

All household members were enumerated and their name, age, and gender listed, regardless of whether they were present. Trained trachoma graders screened both eyes of all present household members for all clinical signs of trachoma.21 Trachoma graders assessed children aged 1–9 years for clean face, described as the absence of nasal and ocular discharge. Multiple teams worked in each district to reduce grader bias so that no one team surveyed all clusters in a district. Individuals presenting signs of active trachoma were offered treatment with 1% tetracycline eye ointment. Individuals with TT were referred to the nearest health center for surgical services. Questions to determine whether individuals with TT were known or unknown to the health system based on
new recommendations from the WHO\textsuperscript{23} were included in the questionnaire for the final 37 districts surveyed. If an individual with unoperated TT was detected, then that person was asked if he/she had been offered surgery, with a response of “yes” indicating that the person was known to the health-care system.

Survey teams were supervised by dedicated TIS supervisors, all of whom worked for the regional trachoma program or the Carter Center. Each supervisor supported two to four teams per survey and accompanied the teams for the duration of the survey. Supervisors were trained before the start of each survey on the protocol. Supervisors’ primary role was to ensure survey teams conducted the survey according to the standardized protocol and to address issues as they arose.

Data management and analysis. Data were stored on the internal memory and on removable storage cards on the tablets and downloaded to password-protected computers by field supervisors every 2–5 days. Data from each enumerated individual were linked to the corresponding household-level data using a household serial number automatically generated by Swift Insights.

Statistical analysis was conducted using Stata version 14.0 (StataCorp, College Station, TX). Zonal and regional estimates were calculated, in some instances by aggregating data from multiple survey rounds. All estimates were calculated using the complex survey commands in Stata (svy) with weights created using the inverse of the sampling probability of both stages of sampling. Sampling weights allowed for district-level data to be aggregated to the zonal and regional levels. Robust CIs were estimated using Taylor-series linearization, which accounted for the clustering at household and cluster levels, and a finite population correction was specified for both stages of sampling.

RESULTS

Characteristics of study communities, households, and participants. One thousand eight hundred eighty-seven clusters in 152 districts were surveyed. Among districts for which gott-level data were available (1,801), 30% of the sampled communities were located along a paved road, 11.5% had any community-level electricity supply, 66.2% had mobile phone coverage at the time the survey team visited, and 23.8% had a health facility. Survey teams recorded the time taken to walk from the vehicle to the cluster: 53.9% required less than a 1-hour walk, 25.8% required a 1- to 3-hour walk, and 20.4% required greater than a 3-hour walk.

Sixty-six thousand eighty-nine households were included in the survey. The prevalence of a functional household radio was 18.3% (95% CI: 17.4–19.2), a functioning TV 4.0% (95% CI: 3.4–4.7), working electricity in the home 11.4% (95% CI: 9.9–13.0), and a functioning mobile phone 20.2% (95% CI: 19.1–21.3). Corrugated iron or metal (54.5%) was the primary construction material making up the roof of the house, followed by thatch (38.9%).

Within the study clusters, 276,068 people were enumerated, and, among those, 208,265 people were examined for clinical signs of trachoma (Table 3), representing a 75.4% response rate. The average household size was 4.2 people. Children aged 1–9 years made up 29.0% of survey participants, 57.5% were adults aged ≥15 years, and 50.6% were female.

Water, sanitation, and hygiene characteristics. The round-trip time taken to collect water, including travel to the water source, queuing, collecting water, and returning home, was less than 30 minutes for 66.2% (95% CI: 64.1–68.2; Table 4) of households. This figure reflects a 10.7% decline from 74.1% at baseline in 2007.\textsuperscript{7} Households collecting water from an improved or a safe source increased to 48.1% (95% CI: 45.5–50.6) from 34.4% at baseline\textsuperscript{7}; 50.2% (95% CI: 48.8–51.5) of households had a latrine present, a 106.6% increase from 24.3% at baseline\textsuperscript{7}; and 46.2% (95% CI: 44.8–47.5) of households had evidence of latrine use. A handwashing station was present in 13.9% (95% CI: 12.7–15.2) of households. Among children aged 1–9 years, 76.5% (95% CI: 75.3–77.7) had a clean face free of nasal and ocular discharge.

Trachoma prevalence. The regional prevalence of TF among children aged 1–9 years was 25.9% (95% CI: 24.9–26.9; Table 5), a decrease of 33.8% from the National Survey in 2006\textsuperscript{5} and of 20.8% from the zonal baseline survey in 2007.\textsuperscript{7} The prevalence of TI among children aged 1–9 years saw the greatest decline among all trachoma clinical signs since baseline, decreasing 86.8% from 41.7% in 2006\textsuperscript{6} to 5.5% (95% CI: 5.2–6.0). The prevalence of active trachoma was 28.3% (95% CI: 27.3–29.3), a decrease of 54.8% from 62.6% in 2006.\textsuperscript{6} A lower zonal TF point estimate was observed at TIS than at baseline in seven of the 10 zones.

Trachomatous scarring prevalence among the total population was 7.7% (95% CI: 7.3–8.2) and 12.9% (95% CI: 12.2–13.6) among adults aged ≥15 years. Trachomatous trichiasis prevalence among the total population was 2.1% (95% CI: 2.0–2.2). Among adults aged ≥15 years, 3.9% (95% CI: 3.7–4.1) presented with TT, a decrease of 37.1% from 6.2% at baseline in 2007\textsuperscript{7} (Table 6). Zonal-level estimates of TT unknown to the health system could not be calculated, as these data were collected only for the final 37 districts surveyed, in accordance with recommendations from the WHO released in 2015,\textsuperscript{23} and estimates for all districts in any one zone were not available; however, the range by district was 0.0% to 5.0%. Eight of the 10 zonal TT point estimates were lower at TIS than at baseline.

Substantial differences between zones were observed (Figures 2 and 3). North Gonder Zone presented the lowest prevalence of TF among children aged 1–9 years of 16.8% (95% CI: 14.3–19.7). Both North Gonder and Awil zones had the lowest prevalence of TI among children aged 1–9 years of 3.4%. Waghemra Zone had the highest prevalence of TF and TI among children aged 1–9 years of 54.7% (95% CI: 49.7–59.6) and 13.7% (95% CI: 11.5–16.1), respectively. South Wollo Zone evidenced the lowest prevalence of both TS and TT among adults aged ≥15 years of 7.2% (95% CI: 6.3–8.3) and 2.6% (95% CI: 2.3–3.0), respectively. East Gojam Zone had the highest prevalence of TT among adults aged ≥15 years of 5.9% (95% CI: 5.2–6.6), whereas South Gonder Zone harbored the highest prevalence of TS among adults aged ≥15 years of 23.0% (95% CI: 20.8–25.4).

Overall, nine districts achieved the TF target for elimination as a public health problem, whereas four districts achieved the target for TT (Figures 2 and 3, Supplemental Table 1). Among those districts, only two achieved both TF and TT targets. Seventy districts had an estimated prevalence of TF among children aged 1–9 years between 10% and 29.9%, whereas TF was hyperendemic (≥30%) in 56 districts. One hundred two districts presented a TT prevalence among adults aged ≥15
year between 1.0% and 4.9%, whereas 39 demonstrated a prevalence of at least 5.0%.

**DISCUSSION**

Between 2010 and 2015, all 152 districts in the Amhara Region were surveyed to evaluate the SAFE strategy and assess progress toward elimination targets. Compared with baseline data, overall, these TIS demonstrated substantial uptake of SAFE interventions and successful reductions in the clinical signs of TF, TI, and TT. As a result of this work, Amhara became the first region in Ethiopia to complete TIS in every district.

After the full recommended 3–5 years of SAFE was implemented throughout the region with high coverage, only nine districts met the elimination thresholds for TF, four met the TT targets, and two of these met the targets for both TF and TT. Substantial heterogeneity of trachoma prevalence among districts, even districts within the same zone, persists. Notably, 56 (37%) districts are hyperendemic, despite the SAFE activities. The results of these surveys make it clear that a combination of all S, A, F, and E activities continue to be warranted in 150 of 152 districts in Amhara, and that guidelines for SAFE interventions may need to be revisited for trachoma hyperendemic areas.

The TIS demonstrated an increase in the use of an improved water source from baseline, as well as a substantial increase in the presence of household latrines—both indications of some improvement in hygiene and sanitation developments that impact trachoma. Still, less than half of households reported collecting water from an improved water source and about 50% had an observed household latrine, highlighting the continued need for water, sanitation, and hygiene infrastructure. Continued health education, supported through delivery at schools and in the community, can also encourage the uptake of F and E components of the SAFE strategy.

Despite improvements in water and sanitation, the results from these surveys are consistent with those reported in other hyperendemic settings, in randomized trials, and in mathematical models. Unlike the success seen in hypendemic and mesoendemic areas, evidence from hyperendemic settings highlight that 3–5 years of MDA may not be sufficient to reduce the prevalence of TF below elimination thresholds. Trachomatous inflammation-follicular has been shown to stabilize or return in high-burden areas despite annual or biannual MDA. Although evidence from clinical trials of biannual MDA did not show a statistically significant difference in efficacy to a single-round MDA, intensified MDA shows promise for accelerating the speed of decline of trachoma. Other treatment regimens, with
either more intense, or better targeted MDA strategies should be tested.28,32,33

Although TF and TT are the indicators by which elimination as a public health problem is defined, evidence suggests that TI may be a better indicator of infection with Chlamydia trachomatis, the bacterium that causes trachoma. Trachomatous inflammation-follicular has also been shown to persist in areas despite low levels or no infection with C. trachomatis.6,34,35 Purporting that TF may overestimate the presence of chlamydial infection in communities. Instead, studies have demonstrated a highly correlated relationship between TI and C. trachomatis.8,34,36 Future surveys should grade individuals for TI, as this clinical sign can be used as a proxy indicator for trachoma infection.

The TIS survey schedule recommended by the global program has evolved, shifting from 3–5 years following SAFE implementation, as established in 2010,37,38 to the current standard of 1–7 years, adopted in 2017. The exact number of years is dependent on the baseline prevalence of TF in the health district. The results from TIS in Amhara support the decision by the International Trachoma Initiative Trachoma Expert Committee to extend the number of years of SAFE implementation, including annual MDA in areas with baseline TF prevalence ≥ 50%, from 5 to 7 years,39 and align with previous evidence suggesting 7–10 years may be more appropriate.40 However, given the sheer scale of work required to complete TIS, enormous resources and time will be required for national programs to adhere to the presently recommended frequency of surveys. For example, almost one-half of the 1,887 clusters surveyed during the TIS described here required survey teams to walk more than 1 hour each way to reach the selected clusters.

Because multiple TIS and surveillance surveys are required in each health district, future survey designs must not only be scientifically sound but also be feasible to complete in a timely manner to also allow sufficient time during the year for SAFE strategy implementation. As the requirements for surveys increase as districts approach elimination targets, there is a risk that trachoma elimination programs become trachoma surveillance programs, which will do nothing to eliminate the disease.

These TIS provided robust prevalence estimates following full-scale implementation of the SAFE strategy. Compared with the 2007 zonal-level baseline, three zones evidenced a higher prevalence of TF among children aged 1–9 years, whereas one zone presented a higher prevalence of TT among adults aged ≥ 15 years. These apparent higher estimates of TF and TT may be due to methodological differences in the surveys, particularly because the magnitude of SAFE interventions delivered to these zones would make an increase in chlamydial infection unlikely. Interpreting different prevalence levels from baseline to TIS, both increases and decreases, will be a challenge that many trachoma programs will face globally as baseline surveys are often conducted using large

| Zone            | No examined | % (95% CI) | No examined | % (95% CI) | % (95% CI) | % (95% CI) |
|-----------------|-------------|------------|-------------|------------|------------|------------|
| Awi             | 893         | 5.4 (4.0–7.3) | 5,242       | 11.5 (9.0–14.5) | 3.5 (2.9–4.2) |
| East Gojam      | 881         | 7.1 (5.4–9.4) | 11,165      | 11.3 (9.3–13.6) | 5.9 (5.2–6.6) |
| North Gondar    | 730         | 4.3 (2.8–6.6) | 11,979      | 13.3 (11.4–15.3) | 3.2 (2.7–3.7) |
| North Shoa      | 943         | 9.0 (6.7–11.9) | 12,625      | 10.7 (9.6–11.9) | 4.2 (3.8–4.7) |
| North Wollo     | 971         | 9.4 (7.2–12.1) | 9,566       | 11.9 (10.1–14.0) | 4.0 (3.4–4.6) |
| Oromia          | 904         | 3.8 (2.5–5.7) | 18,067      | 23.0 (20.8–25.4) | 3.8 (3.4–4.3) |
| South Gondar    | 931         | 3.2 (2.2–4.6) | 26,522      | 7.2 (6.3–8.3) | 2.6 (2.3–3.0) |
| South Wollo     | 1,030       | 6.3 (3.9–9.9) | 3,562       | 19.4 (16.3–23.0) | 5.8 (4.8–7.1) |
| Waghemra        | 874         | 10.0 (6.3–15.6) | 8,326       | 14.7 (12.3–17.4) | 2.9 (2.4–3.6) |
| Region          | 9,121       | 6.2 (5.3–7.4) | 110,211     | 12.9 (12.2–13.6) | 3.9 (3.7–4.1) |
enumeration units. Future district-level TIS in Amhara will allow for clearer assessments of progress toward trachoma elimination targets and uptake of SAFE interventions.

From 2010 to 2015, 152 districts, including 1,887 clusters throughout the Amhara Region, were surveyed to assess trachoma prevalence and the uptake of SAFE interventions. Two districts have met the trachoma elimination targets, whereas continued interventions are required in the remaining 150 districts. Trachoma impact surveys and then trachoma surveillance surveys have continued in eligible districts in Amhara. In areas

**Figure 2.** District counts of trachomatous inflammation-follicular prevalence among children aged 1–9 years, Amhara, Ethiopia, 2010–2015.

**Figure 3.** District counts of trachomatous trichiasis prevalence among adults aged ≥ 15 years, Amhara, Ethiopia, 2010–2015.
with a high trachoma burden at baseline, like Amhara, trachoma interventions are likely to be required for a consider-
able period before the elimination targets are achieved.

Received June 13, 2019. Accepted for publication July 29, 2019.
Published online September 23, 2019.

Note: Supplemental table appears at www.ajtmh.org.

Acknowledgments: We are grateful to all study participants who vol-
unteered their time to participate in the study. We appreciate and thank the dedicated field teams, drivers, and supervisors who per-
severed through difficult conditions to complete the data collection process, many of whom participated in all surveys.

Financial support: Funding for the survey in summer 2013 was pro-
vided by USAID’s ENVISION project led by RTI International.

Authors’ addresses: Aisha E. P. Stewart, Andrew W. Nute, Elizabeth
Kelly Callahan, and Scott D. Nash, The Carter Center, Atlanta, GA, E-mails: aisha.stewart@cartercenter.org, andrew.nute@cartercenter.org, kelly.
callahan@cartercenter.org, and scott.nash@cartercenter.org. Mulat
Zerihun, Demelash Gessese, and Berhanu Melak, The Carter Center, Bahir
Dar, Ethiopia, E-mails: mulat.zerihun@cartercenter.org, demelash.
gessese@cartercenter.org, and berhanu.melak@cartercenter.org. Esheu
Sata, Tigist Astale, Tekolak Endeshaw, and Zerihun Tadesse, The Carter
Center, Addis Ababa, Ethiopia, E-mails: esheu.sata@cartercenter.org,
tigist.astale@cartercenter.org, tekolak.endeshaw@cartercenter.org,
and zerihun.tadesse@cartercenter.org. Tesfaye Teferi, International Tra-
choma Initiative, Addis Ababa, Ethiopia, E-mail: tteferi@taskforce.org.
Melsew Chanyalew, Amhara National Regional Health Bureau, Bahir
Dar, Ethiopia, E-mail: yeshework97@yahoo.com. Birhan Gaudie, Dr.
Abdu Higher Eye Clinic, Bahir Dar, Ethiopia, E-mail: birhanayabi@gmail.
com. Paul M. Emerson, International Trachoma Initiative, Atlanta, GA,
E-mail: pemerson@taskforce.org. Jonathan D. King, World Health Or-
ganization, Geneva, Switzerland, E-mail: kingj@who.int.

This is an open-access article distributed under the terms of the
Creative Commons Attribution License, which permits unrestricted
use, distribution, and reproduction in any medium, provided
the original author and source are credited.

REFERENCES
1. WHO, 2016. Validation of the Elimination of Trachoma as a Pub-
licity Health Problem. Geneva, Switzerland: World Health
Organization.
2. WHO, 1997. Future Approaches to Trachoma Control. Geneva,
Switzerland: World Health Organization.
3. WHO, 2010. Report of the Third Global Scientific Meeting on
Trachoma Elimination. Geneva, Switzerland: World Health
Organization.
4. Berhane Y, Worku A, Bejiga A, 2006. National Survey on Blind-
ness, Low Vision and Trachoma in Ethiopia. Addis Ababa, Ethiopia: Federal Ministry of Health.
5. Ngondi J et al., 2009. Evaluation of three years of the safe strategy
(surgery, antibiotics, facial cleanliness and environmental
improvement) for trachoma control in five districts of Ethiopia
hyperendemic for trachoma. Trans R Soc Trop Med Hyg 103:
1001–1010.
6. Nash SD et al., 2018. Ocular Chlamydia trachomatis infection
under the surgery, antibiotics, facial cleanliness, and environ-
mental improvement strategy in amhara, Ethiopia, 2011–2015.
Clin Infect Dis 67: 1840–1846.
7. Emerson PM et al., 2008. Integrating an NTD with one of “the
big three”: combined malaria and trachoma survey in Amhara
region of Ethiopia. PLoS Negl Trop Dis 2: e197.
8. BRFED, 2006. Population Estimates by Ethiopia Bureau of Fi-
nance and Economic Development for the Year 2006/2007.
Addis Ababa, Ethiopia: Ethiopia Bureau of Finance and Eco-
nomic Development.
9. The Carter Center, 2016. Summary Proceedings: All Eyes on
2020. 17th Annual Trachoma Program Review, March 7–9,
2016, Atlanta, GA.
10. The Carter Center, 2008. Summary Proceedings: Ensuring
Implementation of the Full Safe Strategy. Ninth Annual
Trachoma Program Review, February 11–13, 2008, Atlanta,
GA.
11. The Carter Center, 2015. Summary Proceedings: Looking Back,
Moving Forward. 16th Annual Trachoma Program Review,
March 2–4, 2015, Atlanta, GA.
12. The Carter Center, 2014. Summary Proceedings: Focus on Im-
 pact. 15th Annual Trachoma Program Review, February 25–27,
2014, Atlanta, GA.
13. The Carter Center, 2013. Summary Proceedings: Efficient Pro-
gram Delivery-Doing More, Better, Faster. 14th Annual
Trachoma Control Program Review, March 11–13, 2013, Atlanta,
GA.
14. The Carter Center, 2012. Summary Proceedings: Shaping Pro-
grams to Fit the Need-The Relevance of Prevalence. 13th An-
nual Trachoma Control Program Review, February 27–29,
2012, Atlanta, GA.
15. The Carter Center, 2011. Summary Proceedings: Achieving
Elimination Targets. 12th Annual Trachoma Control Program
Review, February 22–24, 2011, Atlanta, GA.
16. The Carter Center, 2010. Summary Proceedings: Planning for
Trachoma Elimination, District by District. 11th Annual
Trachoma Control Program Review, March 29–31, 2010, Atlanta,
GA.
17. The Carter Center, 2009. Summary Proceedings: From Control to
Elimination. 10th Annual Trachoma Control Program Review,
February 11–13, 2009, Atlanta, GA.
18. Turner AG, Magnani RJ, Shuaib M, 1996. A not quite as quick but
much cleaner alternative to the expanded programme on im-
munization (epi) cluster survey design. Int J Epidemiol 25:
198–203.
19. King JD et al., 2014. Prevalence of trachoma at sub-district level in
Ethiopia: determining when to stop mass azithromycin distri-
bution. PLoS Negl Trop Dis 8: e2732.
20. Minassian D, 1997. Epidemiology in practice: sample size cal-
culation for eye surveys: a simple method. J Comm Eye Health
10: 42–44.
21. Thylefors B, Dawson CR, Jones BR, West SK, Taylor HR, 1987. A
simple system for the assessment of trachoma and its com-
plications. Bull World Health Organ 65: 477–483.
22. King JD et al., 2013. A novel electronic data collection system for
large-scale surveys of neglected tropical diseases. PLoS One
8: e74570.
23. WHO, Department of Control of Neglected Tropical Diseases,
2015. Technical Consultation on Trachoma Surveillance. Ge-
neva, Switzerland: World Health Organization.
24. Solomon AW et al., 2015. The global trachoma mapping project:
methodology of a 34-country population-based study. Oph-
thalmic Epidemiol 22: 214–225.
25. Barnani S, King JD, Dembele M, Coulibaly F, Sankara D,
Kamissoko Y, Ting J, Rotondo LA, Emerson PM, 2010. Where
do we go from here? Prevalence of trachoma three years after
stopping mass distribution of antibiotics in the regions of Kayes
and Koulikoro, Mali. PLoS Negl Trop Dis 4: e734.
26. West SK, Munoz B, Mkocha H, Gaydos CA, Quinn TC, 2011.
Number of years of annual mass treatment with azithromycin
needed to control trachoma in hyper-endemic communities in
Tanzania. J Infect Dis 204: 268–273.
27. Keenan JD et al., 2018. Mass azithromycin distribution for hy-
perendemic trachoma following a cluster-randomized trial: a
continuation study of randomly reassigned subclusters (TANA
II). PLoS Med 15: e1002633.
28. Gebre T et al., 2012. Comparison of annual versus twice-yearly
mass azithromycin treatment for hyperendemic trachoma in
Ethiopia: a cluster-randomised trial. Lancet 379: 143–151.
29. Lietman T, Porco T, Dawson C, BLOWER S, 1999. Global elimina-
tion of trachoma: how frequently should we administer mass az-
ithromycin therapy? Nat Med 5: 572–576.
30. Ray KJ et al., 2007. A rationale for continuing mass antibiotic
distributions for trachoma. BMC Infect Dis 7: 91.
31. Lakew T et al., 2009. Reduction and return of infectious trachoma
in severely affected communities in Ethiopia. PLoS Negl Trop
Dis 3: e376.
32. Lietman TM, Pinsent A, Liu F, Deiner M, Hollingsworth TD, Porco TC, 2018. Models of trachoma transmission and their policy implications: from control to elimination. *Clin Infect Dis* 66: S275–S280.

33. Melese M et al., 2008. Comparison of annual and biannual mass antibiotic administration for elimination of infectious trachoma. *JAMA* 299: 778–784.

34. Ramadhani AM, Derrick T, Macleod D, Holland MJ, Burton MJ, 2016. The relationship between active trachoma and ocular *Chlamydia trachomatis* infection before and after mass antibiotic treatment. *PLoS Negl Trop Dis* 10: e0005080.

35. West SK, Munoz B, Mkocha H, Holland MJ, Aguirre A, Solomon AW, Foster A, Bailey RL, Mabey DC, 2005. Infection with *Chlamydia trachomatis* after mass treatment of a trachoma hyperendemic community in Tanzania: a longitudinal study. *Lancet* 366: 1296–1300.

36. Keenan JD et al., 2010. Clinical activity and polymerase chain reaction evidence of chlamydial infection after repeated mass antibiotic treatments for trachoma. *Am J Trop Med Hyg* 82: 482–487.

37. WHO, 2004. *Report of the Eighth Meeting of the Who Alliance for the Global Elimination of Blinding Trachoma*. Geneva, Switzerland: World Health Organization.

38. Solomon AW, Zondervan M, Kuper H, Buchan J, Mabey D, Foster A, 2006. *Trachoma Control: A Guide for Programme Managers*. Geneva, Switzerland: World Health Organization.

39. International Trachoma Initiative, 2019. *Zithromax Management Guide*. Decatur, GA: International Trachoma Initiative.

40. Adera TH et al., 2016. Prevalence of and risk factors for trachoma in southern nations, nationalities, and peoples’ region, Ethiopia: results of 40 population-based prevalence surveys carried out with the global trachoma mapping project. *Ophthalmic Epidemiol* 23: 84–93.

41. Bero B et al., 2016. Prevalence of and risk factors for trachoma in Oromia regional state of Ethiopia: results of 79 population-based prevalence surveys conducted with the global trachoma mapping project. *Ophthalmic Epidemiol* 23: 392–405.