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KATABARWA AND OTHERS

TRANSMISSION OF ONCHOCERCIASIS IN NORTHWESTERN UGANDA

Transmission of *Onchocerca volvulus* Continues in Nyagak-Bondo Focus of Northwestern Uganda after 18 Years of a Single Dose of Annual Treatment with Ivermectin

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Abstract.

The objective of the study was to determine whether annual ivermectin treatment in the Nyagak-Bondo onchocerciasis focus could safely be withdrawn. Baseline skin snip microfilariae (mf) and nodule prevalence data from six communities were compared with data collected in the 2011 follow-up in seven communities. Follow-up mf data in 607 adults and 145 children were compared with baseline (300 adults and 58 children). Flies collected in 2011were dissected, and poolscreen analysis was applied to ascertain transmission. Nodule prevalence in adults dropped from 81.7% to 11.0% (P < 0.0001), and mf prevalence dropped from 97.0% to 23.2% (P < 0.0001). In children, mf prevalence decreased from 79.3% to 14.1% (P < 0.0001). Parous and infection rates of 401 flies that were dissected were 52.9% and 1.5%, respectively, whereas the infective rate on flies examination by polymerase chain reaction (PCR) was 1.92% and annual transmission potential was 26.9. Stopping ivermectin treatment may result in onchocerciasis recrudescence.

INTRODUCTION

Onchocerciasis, a leading cause of blindness, is caused by infection with *Onchocerca volvulus*, a filarial nematode parasite. The female worms that live in the nodules produce microfilariae (mf), which inflame the skin. The mf may also enter the eyes, giving rise to inflammatory lesions that may eventually result in partial or complete blindness.¹ The parasite is transmitted by black flies of the genus *Simulium*, which breed in fast-flowing rivers and streams (hence, the name river blindness for the disease). The mf are picked up by female *Simulium* flies from an infected person during a blood meal, and within the fly, they develop into larval (L1, L2, and L3) stages. The L3 (infective) stage is passed on to other persons during subsequent bites.

Ivermectin (Mectizan), a safe and effective microfilaricide, has been donated by Merck & Co Inc. since 1987. Merck pledged to provide as much ivermectin as required for as long as necessary for mass treatment of onchocerciasis. Ivermectin kills the mf and reduces the risk of developing eye disease and severe skin lesions associated with the infection. Ivermectin also reduces the fecundity of adult worms and shortens their

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lifespan, although treatment must still be given for an undetermined length of time when taken one time in a year.^{2,3}

The African Program for Onchocerciasis Control (APOC) was set up to establish, promote, and support sustainable community-directed treatment with ivermectin (CDTI) projects for the control of onchocerciasis in meso- and hyperendemic areas that had onchocercal nodule rates of $\geq 20\%$ and $\geq 40\%$, respectively.^{4–6} The goal was to target the most highly onchocerciasis-endemic communities with a single annual dose of ivermectin through mass treatment using the CDTI strategy; therefore, the disease would no longer be a public health problem.^{7–9} This goal was not defined but logically taken to be when prevalence is driven below the original baseline threshold required to launch the mass ivermectin treatment program, which is an onchocerciasis as a public health problem defined at these levels did not necessarily indicate interruption of transmission. In the case that transmission has not been interrupted, halting mass treatment could result in disease recrudescence.^{2,3}

Despite this information, a recent study in Mali and Senegal suggested that 15–17 years of annual treatment was sufficient to eliminate *O. volvulus* transmission.¹⁰ This study called for additional studies in other endemic areas of Africa that have distributed single annual doses of ivermectin for onchocerciasis control over similar time frames to determine whether treatment could safely be withdrawn without the risk of disease recrudescence. It is possible that the amount of time necessary to eliminate transmission could vary considerably among foci in Africa depending on the initial prevalence of infection and the intensity of transmission.¹¹

Epidemiological assessments done in 1993 had shown that the Nyagak-Bondo area of northwestern Uganda (Zombo district and parts of Arua and Nebbi districts) was endemic for onchocerciasis. The vectors in Nyagak-Bondo focus are members of the S. neavei complex, with larvae and pupae that live in a phoretic association on freshwater crabs of species Potamonautes aloysiisabaudiae and P. niloticus.¹² The River Blindness Foundation (RBF) -assisted community-based treatment with ivermectin (CBTI) for onchocerciasis control activities was carried out from 1993 to 1996 in the then Nebbi district (now Nebbi and Zombo districts). Subsequently, RBF operations were taken over by The Carter Center in 1996, which continued to assist the program to date. The Kuluva Missionary Hospital, funded by Christoffel Blinden Mission (CBM), assisted mass treatment activities in neighboring Arua district from 1993 to 1995. In 1996, APOC was established and subsequently provided 5 years of financial assistance for implementation and establishment of mechanisms to sustain the CDTI program. APOC continued providing financial support to the program for 3 additional years to replace capital items and provide some training and advocacy. Subsequently, the CDTI project was expected to sustain annual mass treatment using only national resources. However, the national government did not take over the funding of CDTI activities.¹³ Therefore, APOC continued to provide some assistance for specific activities alongside regular CDTI support from The Carter Center in Nebbi and Zombo districts and recently, Arua district by USAID funds for Neglected Tropical Diseases through RTI International. Overall, with the support of these agencies, annual ivermectin treatment has been provided to the at-risk population of Nyagak-Bondo for a period of 18 years. The objective of the present study was to determine whether annual mass treatment with ivermectin for 18 years could be withdrawn in Nyagak-Bondo onchocerciasis focus like in Mali and Senegal without the risk of disease recrudescence

MATERIALS AND METHODS

Study sites.

The study was carried out in Nyagak-Bondo focus of onchocerciasis, which covers areas of Arua, Nebbi, and Zombo districts of northwestern Uganda (Figure 1). The main river systems responsible for vector breeding are the Agoi, Nyagak, and Ora.¹⁴ The Nyagak-Bondo focus covers an area of about 1,550 km², with a total of about 510,600 people.

Annual mass treatment with ivermectin for onchocerciasis control began in 1993 after six communities (Abilambe, Agweci, Jupa Ngali Upper, Nyadima, Patek-Athele, and Ukongo) were selected for sentinel evaluations, and assessed for microfilardermia and nodule prevalence. Three of these communities (Abilambe, Agweci, and Patek-Athele) were assessed again in 2011, and the results were compared with those results obtained in 1993. It was not possible to locate the other sentinel communities, because their structure and names had changed because of administrative changes since 1993. However, the Health Department wanted to know the situation in and outside the sentinel communities, and therefore, additional communities (Aguru, Kairo, Pachen, and Oloamura) were included in the follow-up survey.

Parasitological (nodule and mf prevalence) surveys.

Baseline parasitological surveys carried out in 1993 before mass treatment were followed up in 2011 (11 months after mass treatment).

Nodule assessments.

Every participant was examined in a well-lit private room. Trained district and Ministry of Health workers performed a palpation examination on the partially undressed participant, paying attention to bony prominences of the torso, iliac crests, and upper trochanter of the femurs. Onchocercal nodules were identified clinically as being firm, painless, and mobile.^{15,16} In 1993, 180 adults ages 20 years and over who had lived in their respective communities for at least 10 years were assessed for nodule prevalence. In the seven communities in 2011, 607 adults were assessed, and the results were recorded on the study registration form as positive or negative. Nodule prevalence was expressed as a percentage, and follow-up data were compared with baseline data.

Mf assessments.

At baseline, 300 adults ages 20 years and over who had lived in their respective communities for at least 10 years and 58 children at 5 years of age from six baseline communities were assessed for mf prevalence. In the 2011 follow-up study, 607 resident adults and 145 resident children less than 10 years of age (born after mass treatment in all seven communities had started) were skin snipped. The procedure for skin snipping involved cleaning the site with an antiseptic; then, a piece of skin raised with the help of a

disposable sterile dermal hook was carefully removed with a sterilized surgical blade. Two skin snip samples were taken from the posterior iliac crests of every selected person. Every person that was skin snipped had his or her own dermal hook and surgical blade, which were carefully disposed of after use.

The skin samples were placed immediately in wells of microtiter plates containing a sterile normal saline solution; they were kept at room temperature for 12–24 hours and examined microscopically for mf.^{17–19} The results were expressed for each individual as positive or negative and recorded in the study registration form. Mf prevalence was expressed as a percentage.⁴ The follow-up results were compared with baseline mf prevalence data of 1993 from adults and children.

ENTOMOLOGICAL ASSESSMENT

Crab infestation.

Crab trapping was conducted in the months of July and December of 2010 and continued through the months of January, March to June, and September to December of 2011 at a number of sites on the Nyagak, Agoi, and Wariki river systems. The crabs carrying larval and pupal stages of *S. neavei* were counted, and infestation rate (number of crabs positive for young stages of the fly) was expressed as a percent of the total number of crabs captured.²⁰

Fly collection and analysis.

Two *Simulium* fly collection sites were set up at selected breeding points on the Agoi and Wariki rivers. Organized full-day catches were conducted for 7 consecutive days, 1 month in June, and September to December of 2010. Human landing collections were carried out from 07:00 to 18:00 daily.¹⁹ A total of 401 flies collected during the period were dissected to determine parity, infection, and infective rates.^{21,22} Examination of the larval stages of *O. volvulus* involved dissection of the abdomens, thoraces, and heads of the collected *Simulium* flies. The numbers of flies with larval stages (L1, L2, and L3) were counted, and monthly infection and infective rates were calculated. The result was expressed as a percentage of the total number of flies dissected × 100.⁴

Additional *Simulium* flies were caught from routine fly collections in 2011 from two previous collection sites on the Agoi and Wariki rivers, with an additional site on the Nyagak river system over a period of 9 months (2 days per week every month). Human landing collections were carried out from 07:00 to 18:00 daily as previously described.¹⁹ A total of 1,100 individual flies divided into 22 pools of 50 flies each was analyzed using polymerase chain reaction (PCR). Only the heads of these flies were subjected to PCR analysis to calculate the infective rate and annual transmission potential (ATP).^{23–25} The ATP was estimated based on the number of flies collected and the estimated 7,766 exposure hours (i.e., daylight hours when flies could bite) at the site. This information implies 12 hours of daylight over 365 days in the year. Collection times in estimating the monthly and annual biting rate were based on 40 collection hours per month over the 9-month collection period.

History of mass treatment with ivermectin.

Annual mass treatment with ivermectin commenced in 1993, and it was carried out annually through 2011 in all three districts. The objective was to sustain annual treatment of at least 90% of the ultimate treatment goal (UTG). The UTG is the sum of all eligible persons for treatment (minus children < 5 years old) among the total number of people at risk living in all at-risk communities in the onchocerciasis-endemic area that the program ultimately has to treat.²⁶ Annual mass ivermectin treatment had been provided for 18 years in Arua, Nebbi, and Zombo districts. The reports since inception of the program were available, and from these reports, UTG treatment coverage for each district was computed.

Data analysis.

Parasitological and entomological data were entered and analyzed graphically in Microsoft Excel and Epi Info, Centers for Disease Control and Prevention, Atlanta GA, for χ^2 test of independence. The prevalence of flies carrying infective larvae and an associated 95% confidence surrounding this point estimate were calculated using Poolscreen v2.0 software.²⁷

Ethical approval.

Collections of data from the baseline to the follow-up studies were considered routine program evaluation by the Government of Uganda and the Emory University Institutional Review Board (eIRB 11 438). Therefore, this study was considered non-research and routine program evaluation. Individuals assessed were educated and informed of the option to opt out without any repercussions.

RESULTS

Mass treatment.

Treatment coverage of the eligible population in Nebbi and Zombo districts was better than in the Arua district. It reached the desired level of 90% in Nebbi and Zombo districts in 1998 and remained above 90% for the next 13 years, whereas in Arua district, it was above 90% for 10 years during the 18-year period of CDTI implementation (Table 1).

Parasitology.

Nodule prevalence.

There was a highly significant reduction in baseline nodule prevalence among adults from 81.7% in 180 persons assessed (range = 73.3-96.7%) to 11.0% in 607 persons (range = 2.3-20%) in the follow-up survey (P < 0.0001) (Table 2). However, nodule prevalence in Kairo community remained at 20%, the threshold where mass treatment is recommended. All other communities were still positive for nodules.

Mf prevalence.

Baseline mf prevalence in 1993 was 97% in 300 adult skin snips, with a range of 90–100% (Table 3). In the follow-up survey of 2011, mf prevalence had been reduced to 23.2% in 607 persons examined, with a range of 3.4-40% (P < 0.0001). However, mf prevalence in six of seven communities in 2011 was at least 19%. In children, mf prevalence had reduced from 79.3% (with a range of 36.4-100% in 1993) to 14.1% (with a range of 0% to 36.8%, P < 0.0001) (Table 4).

Entomology.

In the Nyagak-Bondo onchocerciasis focus, crab infestation rate was 21.4% (N = 3,245), with a monthly range of 8.3–43.9% in an intermittent 11-month survey from July of 2010 to December of 2011 (Figure 2). The *S. neavei* parous rate investigated from June and September to December (a period of 5 months) was 52.9% (N = 401), with a monthly range of 0–67.8%. The infection rate (L1, L2, and L3 larval stages) was 1.50%, with a monthly range of 0–3.85%, whereas the infective rate was zero (Table 5).

Pool screening of 22 head pools (1,100 flies) resulted in a single confirmed positive pool, which revealed a prevalence of infective flies of 1.92% (95% upper limit = 7.4/2,000 flies). In addition, the ATP was estimated at 26.9 (95% confidence interval = 0–66) third-stage (L3) larvae per person per year.

DISCUSSION

The results showed that, after 18 years of annual mass treatment with ivermectin, there was a significant reduction in infection. However, the children born well after ivermectin distribution commenced were still getting infected with *O. volvulus*, and a substantial proportion of adults were still positive for mf. This finding was not surprising, because breeding of *S. neavei* was ongoing, with mean crab infestation at 21.4%, a fly parous rate of 52.9%, infection rate of 1.50%, and poolscreen infective rate of 1.92% with the ATP of 26.9.

Nodule and mf rates.

The reduction in nodule rate was significant in most communities; however, one community remained at 20%, which under the APOC policy for controlling onchocerciasis as a public health problem, was at the threshold for mass treatment. It is, however, possible that, at low nodule prevalence, there are likely confounding factors, such as ganglia and *Taenia solium*-related nodules, that may artificially increase the calculated nodule rate.⁶ Despite this information, the data showed consistency of high microfildermia prevalences, implying a continuing high force of transmission. These data suggest that, if annual mass treatment was halted, it is likely that recrudescence of onchocerciasis would occur.²⁸

It is interesting to note that, in the Patek-Athele community, the mf rate was brought to zero from 100%, whereas in other communities, it was not. Because the vector of onchocerciasis is *S. neavei*, it is possible that there could have been ecological changes resulting into unfavorable breeding conditions of the vector in the vicinity of Patek-Athele. Such conditions could result in interruption or suppression of transmission.^{29,30}

However, no *Simulium* fly collection and crab trapping were done in the vicinity of Patek-Athele. We recommend entomological investigations in the streams close to this community to understand the reason for the dramatic change. Alternatively, given that skin snip (microscopy) has low sensitivity at low endemicity levels, the results obtained may not reflect the actual endemicity levels.³¹ If this result is true, undetected microfiladermic individuals could still pose serious threat to disease recrudescence.^{31,32} Additional studies are needed to understand the roles of ecological changes, low endemicity in disease transmission, and possible recrudescence.

Continuing transmission.

Interruption of transmission is considered to have been attained if the calculated upper 95% confidence interval of the ATP is less than 5–20 L3/person or the prevalence of flies carrying infective larvae is less than $\frac{1}{2}$,000 in the overall fly population (< 0.05%).³³ The point estimate of the ATP in Nyagak-Bondo was still 26.9, with a point estimate of the prevalence of infective flies of 1.92/2.000 flies. The upper bound of the 95% confidence interval for both metrics was considerably higher than the accepted cutoffs indicative of transmission suppression. These findings are consistent with those findings obtained from northern Cameroon, which showed that 17 years of annual mass treatment did result in a significant decrease of skin mf and nodules but did not interrupt transmission.¹¹ When contrasted with recently published studies from Senegal, Mali, and Nigeria, which show successful interruption of transmission after annual ivermectin treatment, this study points out the importance of local variations in the ecology of transmission of onchocerciasis in determining the ultimate success of an elimination program.¹¹ Local variations in vector density, biting rate, and vector capacity will all affect the potential for the success of an elimination program and by extension, the strategic plan adopted by the program to interrupt transmission. For example, S. neavei, although present in relatively low densities, is known to be a highly anthropophilic and competent vector for *O. volvulus*.³⁴ Annual treatment may not be sufficient to interrupt transmission in areas where such an efficient vector is present. In such areas, it may be necessary to increase the frequency of ivermectin treatments to two times per year.

Efficiency of diagnostic methods in entomology.

During dissection of freshly collected flies, larval stages L1 and L2 were observed, but none of the infective larval (L3) stages were observed. However, during poolscreen, L3 stages were observed. However, only 401 flies were dissected, which contrasts the 1,100 flies that were poolscreened. Dissections of flies may not be necessarily less efficient with well-trained and experienced personnel: the rate of infected flies and numbers of larvae (L1, L2, and L3) can rapidly be assessed, and immediate conclusions on the fly populations and *O. volvulus* can be made. However, when interruption of transmission is the objective, poolscreen should be applied, because it is 100% sensitive and 100% specific in detecting *O. volvulus* DNA.^{35,36} Therefore, we recommend poolscreen in flies where interruption of transmission is launched without vector elimination.

Risk of emergence of resistance.

Anguru and Kairo communities had mf rates of 38% and 40%, respectively. The difference in infection between them and the rest of the sentinel communities was highly significant (P < 0.0001). This result could be because of either low treatment coverage or the possibility of risk of drug resistance caused by long-term mass drug administration.^{11,37} However, there are validated reports of consistent and good annual treatment coverage with a significant reduction of mf rates from baseline to follow-up survey. Therefore, it is doubtful that treatment coverage could be the reason for high follow-up mf rate; alternatively, the high force of infection may be responsible.³ However, studies are required to show whether resistance to ivermectin could be a threat to the long-term success of elimination programs in this region.

CONCLUSION

Although an annual dose of ivermectin given over a period of 18 years has reduced onchocerciasis infection significantly, it has not interrupted the transmission of *O. volvulus*. Although the reports from Senegal and Mali show that, in some areas, onchocerciasis can be eliminated with 15–17 years of annual treatment, the Nyagak-Bondo focus of northwestern Uganda showed that it was not possible. This finding alludes to the fact that local variations in the ecology of transmission of onchocerciasis could determine the ultimate success of an elimination program. Hence, it is imperative to focus on new innovative and flexible approaches to meet the needs of varying ecological conditions that determine the level of onchocerciasis transmission if onchocerciasis elimination becomes the goal throughout Africa.

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REFERENCES

- <jrn>1. Enk CD, Gardlo K, Ruzicka T, BenEzra D, 2003. Onchocerciasis. *Hautarzt 54:* 513–517.
- <jrn>2. Taylor HR, Pacque M, Munoz B, Greene BM, 1990. Impact of mass treatment of onchocerciasis with ivermectin on the transmission of infection. *Science 250:* 116– 118.
- <jrn>3. Borsboom GJ, Boatin BA, Nagelkerke NJ, Agoua H, Akpoboua KL, Alley EW, Bissan Y, Renz A, Yameogo L, Remme JH, Habbema JD, 2003. Impact of ivermectin on onchocerciasis transmission: assessing the empirical evidence that repeated ivermectin mass treatments may lead to elimination/eradication in West-Africa. *Filaria J 2:* 8.
- <bok>4. World Health Organisation, 1991. Report of the World Health Organization: Strategies for Ivermectin Distribution through Primary Health Care System. Geneva: World Health Organization.</bok>
- <jrn>5. Noma M, Nwoke BE, Nutall I, Tambala PA, Enyong P, Namsenmo A, Remme J, Amazigo UV, Kale OO, Sékétéli A, 2002. Rapid epidemiological mapping of onchocerciasis (REMO): its application by the African Programme for Onchocerciasis Control (APOC). *Ann Trop Med Parasitol 96 (Suppl 1):* S29–S39.
- <jrn>6. Katabarwa M, Eyamba A, Habomugisha P, Lakwo T, Ekobo S, Kamgno J, Kuete T, Ndyomugyenyi R, Onapa A, Salifou M, Ntep M, Richards FO, 2008. After a decade of annual dose mass ivermectin treatment in Cameroon and Uganda, onchocerciasis transmission continues. *Trop Med Int Health 13*: 1196–1203.
- <jrn>7. Amazigo UV, Brieger WR, Katabarwa M, Akogun O, Ntep M, Boatin B, N'Doyo J, Noma M, Sékétéli A, 2002. The challenges of community-directed treatment with ivermectin (CDTI) within the African Programme for Onchocerciasis Control (APOC). *Ann Trop Med Parasitol 96 (Suppl 1):* S41–S58.
- <jrn>8. Sékétéli A, Adeoye G, Eyamba A, Nnoruka E, Drameh P, Amazigo UV, Noma M, Agboton F, Aholou Y, Kale OO, Dadzie KY, 2002. The achievements and challenges of the African Programme for Onchocerciasis Control (APOC). *Ann Trop Med Parasitol 96 (Suppl 1):* S15–S28.
- <jrn>9. Brieger WR, Okeibunor JC, Abiose AO, Ndyomugyenyi R, Kisoka W, Wanji S, Elhassan E, Amazigo UV, 2007. Feasibility of measuring compliance to annual ivermectin treatment in the African Programme for Onchocerciasis Control. *Trop Med Int Health 12*: 260–268.
- <jrn>10. Diawara L, Traoré MO, Badji A, Bissan Y, Doumbia K, Goita SF, Konaté L, Mounkoro K, Sarr MD, Seck AF, Toé L, Tourée S, Remme JH, 2009. Feasibility of onchocerciasis elimination with ivermectin treatment in endemic foci in Africa: first evidence from studies in Mali and Senegal. *PLoS Negl Trop Dis 3:* e497.

- <jrn>11. Katabarwa MN, Eyamba A, Nwane P, Enyong P, Yaya S, Baldiagaï J, Madi TK, Yougouda A, Andze GO, Richards FO, 2011. Seventeen years of annual distribution of ivermectin has not interrupted onchocerciasis transmission in North Region, Cameroon. *Am J Trop Med Hyg* 85: 1041–1049.
- <jrn>12. Nelson GS, 1958. Onchocerciasis in the West Nile District of Uganda. Trans R Soc Trop Med Hyg 52: 368–376.
- <jrn>13. Hopkins DR, Richards FO, Katabarwa M, 2005. Whither onchocerciasis control in Africa. Am J Trop Med Hyg 72: 1–2.
- <jrn>14. Katabarwa M, Onapa AW, Nakileza B, 1999. Rapid epidemiological mapping of onchocerciasis in areas of Uganda where *Simulium neavei* is the vector. *East Afr Med J 76:* 440–446.
- <jrn>15. Albiez EJ, Buttner DW, Duke BO, 1988. Diagnosis and extirpation of nodules in human onchocerciasis. *Trop Med Parasitol 39 (Suppl 4):* 331–346.
- <jrn>16. Ngoumou P, Walsh JF, Mace JM, 1994. A rapid mapping technique for the prevalence and distribution of onchocerciasis: a Cameroon case study. *Ann Trop Med Parasitol 88:* 463–474.
- <jrn>17. Prost A, Prod'hon J, 1978. Le diagnostique parasitologique de l'onchocercose. Revue critique des methods en usage. *Medicine Tropicale 38:* 519–532.
- <jrn>18. Schulz-Key H, 1978. A simple technique to assess the total number of Onchocerca volvulus microfilariae in skin snips. Tropenmed Parasitol 29: 51– 54.
- <bok>19. World Health Organization, 1995. Onchocerciasis and Its Control. Report of a WHO Expert Committee on Onchocerciasis Control. Technical Report Series 852. Geneva: World Health Organization .</bok>
- <jrn>20. Garms R, Lakwo TL, Ndyomugyenyi R, Kipp W, Rubaale T, Tukesiga E, Katamanywa J, Post RJ, Amazigo UV, 2009. The elimination of the vector *Simulium neavei* from the Itwara onchocerciasis focus in Uganda by ground larviciding. *Acta Trop 111*: 203–210.
- <jrn>21. Garms R, Cheke RA, 1985. Infections with Onchocerca volvulus in different members of the Simulium damnosum complex in Togo and Benin. Zoology (Jena) 72: 479–495.
- <jrn>22. Jacobi CA, Enyong P, Renz A, 2010. Individual exposure to *Simulium* bites and intensity of *Onchocerca volvulus* infection. *Parasit Vectors 3:* 53.
- <jrn>23. Toé L, Back C, Adjami AG, Tang JM, Unnasch TR, 1997. *Onchocerca volvulus*: comparison of field collection methods for the preservation of parasite and vector samples for PCR analysis. *Bull World Health Organ 75:* 443–447.</jrn>
- <jrn>24. Yamèogo L, Toè L, Hougard JM, Boatin BA, Unnasch TR, 1999. Pool screen polymerase chain reaction for estimating the prevalence of Onchocerca volvulus infection in *Simulium damnosum* sensu lato: results of a field trial in an area subject to successful vector control. *Am J Trop Med Hyg 60:* 124–128.</jrn>

- <jrn>25. Rodríguez-Pérez MA, Danis-Lozano R, Rodríguez MH, Unnasch TR, Bradley JE, 1999. Detection of *Onchocerca volvulus* infection in *Simulium ochraceum* sensu lato: comparison of a PCR assay and fly dissection in a Mexican hypoendemic community. *Parasitology 119:* 613–619.
- <jrn>26. Richards FO Jr, Miri ES, Katabarwa M, Eyamba A, Sauerbrey M, Zea-Flores G, Korve K, Mathai W, Homeida MA, Mueller I, Hilyer E, Hopkins DR, 2001. The Carter Center's assistance to river blindness control programs: establishing treatment objectives and goals for monitoring ivermectin delivery systems on two continents. *Am J Trop Med Hyg 65:* 108–114.
- <jrn>27. Katholi CR, Toe L, Merriweather A, Unnasch TR, 1995. Determining the prevalence of *Onchocerca volvulus* infection in vector populations by polymerase chain reaction screening of pools of black flies. *J Infect Dis* 172: 1414–1417.
- <jrn>28. Bottomley C, Isham V, Collins RC, Basáñez MG, 2008. Rates of microfilarial production by *Onchocerca volvulus* are not cumulatively reduced by multiple ivermectin treatments. *Parasitology 135*: 1571–1581.
- <jrn>29. Raybould JN, White GB, 1979. The distribution, bionomics and control of onchocerciasis vectors (Diptera: Simuliidae) in Eastern Africa and the Yemen. *Tropenmed Parasitol 30:* 505–547.
- <jrn>30. Katabarwa MN, Walsh F, Habomugisha P, Lakwo TL, Agunyo S, Oguttu DW, Unnasch TR, Unoba D, Byamukama E, Tukesiga E, Ndyomugyenyi R, Richards FO, 2012. Transmission of onchocerciasis in Wadelai focus of northwestern Uganda has been interrupted and the disease eliminated. *J Parasitol Res 2012*: 748540. doi:10.1155/2012/748540 [Epub 2012 Aug 26].
- <jrn>31. Boatin BA, Toé L, Alley ES, Nagelkerke NJ, Borsboom G, Habbema JD, 2002. Detection of *Onchocerca volvulus* infection in low prevalence areas: a comparison of three diagnostic methods. *Parasitology* 125: 545–552.
- <jrn>32. Duerr HP, Eichner M, 2010. Epidemiology and control of onchocerciasis: the threshold biting rate of savannah onchocerciasis in Africa. *Int J Parasitol 40:* 641– 650.
- <box>33. World Health Organization, 2001. Certification of elimination of human onchocerciasis: criteria and procedures. *Criteria for Certification of Interruption of Transmission/Elimination of Human Onchocerciasis*. Geneva: World Health Organization.</box/
- <jrn>34. Fischer P, Yocha J, Rubaale T, Garms R, 1997. PCR and DNA hybridization indicate the absence of animal filariae from vectors of *Onchocerca volvulus* in Uganda. *J Parasitol 83*: 1030–1034.
- <edb>35. Unnasch TR, Meredith SEO, 1996. The use of degenerate primers in conjunction with strain and species oligonucleotides to classify *Onchocerca volvulus*. Clapp JP, ed. *Species Diagnostic Protocols: PCR and Other Nucleic Acid Methods*. Totowa, NJ: Humana Press, 293–303.</edb>

- <jrn>36. Fischer P, Rubaale T, Meredith SEO, Büttner DW, 1996. Sensitivity of a polymerase chain reaction-based assay to detect *Onchocerca volvulus* DNA in skin biopsies. *Parasitol Res* 82: 395–401.
- <jrn>37. Osei-Atweneboana MY, Eng JK, Boakye DA, Gyapong JO, Prichard RK, 2007. Prevalence and intensity of *Onchocerca volvulus* infection and efficacy of ivermectin in endemic communities in Ghana: a two-phase epidemiological study. *Lancet 369:* 2021–2029.

FIGURE 1. Nyagak-Bondo onchocerciasis focus of northwest Uganda.

FIGURE 2. Percent monthly infestation of crabs (N = 3,245) captured in Nyagak-Bondo onchocerciasis focus in 2010 and 2011.

	Population and treatment coverage of the eligible population from 1993 to 2010																	
	1993	1994	1995	1996	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010
Nebbi																		
Total population	26,849	33,329	40,296	44,915	50,579	58,992	69,920	81,855	79,981	79,340	81,434	83,407	85,904	87,389	89,574	93,636	95,434	101,360
population	25,256	31,389	38,133	42,581	48,003	52,134	59,048	68,334	65,305	67,573	69,223	70,950	70,035	69,912	72,018	78,905	78,723	84,186
Treated	21,034	24,364	31,636	32,125	35,763	48,592	56,648	65,439	63,536	64,989	66,717	68,468	69,026	68,983	70,843	75,346	77,819	82,226
Covered (%)	83	78	83	75	75	93	96	96	97	96	96	97	99	99	98	95	99	98
Zombo Total																		
population Eligible	96,976	105,873	123,791	132,600	158,075	139,177	141,628	169,411	175,208	181,237	188,561	193,180	208,479	213,552	218,890	213,591	221,715	232,807
population	90,994	99,256	116,858	135,656	149,120	121,866	116,426	140,046	145,346	153,011	160,402	163,749	171,177	170,873	175,988	179,400	183,346	192,498
Treated	80,847	80,469	96,745	103,856	116,001	117,295	111,339	135,426	141,163	147,029	154,752	158,084	167,630	168,678	173,610	171,490	180,361	188,692
Covered (%)	89	81	83	77	78	96	96	97	97	96	96	97	98	99	99	96	98	98
Portion of Arua in																		
Nyak- Bondo																		
Total population	137,987	145,980	149,786	152,890	156,890	168,651	168,651	173,710	178,922	183,432	189,856	194,911	200,416	212,328	217,676	220,744	229,781	231,655
Eligible population	126,852	130,826	132,705	138,206	140,261	143,353	143,353	147,654	152,083	157,914	179,983	185,382	190,943	196,371	202,262	208,329	214,578	221,015
Treated		8,781	10,384	18,856	80,664	80,664	137,795	126,328	128,935	153,296	168,974	181,779	188,295	193,758	201,688	204,232	200,759	212,078
Covered (%)		7	8	14	58	56	96	86	85	97	94	98	99	99	100	98	94	96

TABLE 1

|--|

	con	Baseline (100	3)	ie prevalence of	1999 With a	aano				
District	Community	Examined (r	o.) Positive (no	.) Positive (%)	District	Community	Examined (no) () Positive (no	.) Positive (%)	P value
Zombo	Patek Athele	30	29	96.7	Zombo	Patek Athele	89	9	10.1	< 0.0001
Zombo	Abilambe	30	18	60.0	Zombo	Abilambe	100	14	14.0	< 0.0001
Nebbi	Agweci	30	25	83.3	Nebbi	Agweci	87	2	2.3	< 0.0001
Zombo	Jupa Ngali	30	27	90.0	Arua	Kairo*	75	15	20.0	
Zombo	Nyandima	30	26	86.7	Zombo	Anguru*	89	13	14.6	
Zombo	Ukongo	30	22	73.3	Zombo	Pachen*	101	11	10.9	
					Nebbi	Oloamura*	66	3	4.5	
Overall		180	147	81.7			607	67	11.0	< 0.0001

Comparing data on baseline nodule prevalence of 1993 with data from the follow-up survey of 2011 among adults

* No baseline data.

TABLE 3

		Compar	ing data on base	eline mf from	1993 with d	ata from the f	ollow-up su	urvey of 2011 amo	ong adults	
		1993)				P value for x^2 test of association				
District	Community	Examined (n	o.) Positive (no	.) Mf positive	(%) District	Community E	$\frac{(\%)}{(\%)}$ F value for χ test of association			
Zombo	Patek Athele	50	50	100.0	Zombo	Patek Athele	89	3	3.4	< 0.0001
Zombo	Abilambe	50	48	96.0	Zombo	Abilambe	100	19	19.0	< 0.0001
Nebbi	Agweci	50	48	96.0	Nebbi	Agweci	87	17	19.5	< 0.0001
Zombo	Nyandima	50	50	100.0	Zombo	Anguru*	89	34	38.2	
Zombo	Ukongo	50	50	100.0	Zombo	Pachen*	101	24	23.8	
Zombo J	upa Ngali Uppe	r 50	45	90.0	Nebbi	Oloamura*	66	14	21.2	
					Arua	Kairo*	75	30	40	
Overall		300	291	97.0			607	141	23.2	< 0.0001

* No baseline data.

TABLE 4

		Compa	ring data on bas	eline mf of 19	93 with data	a from the foll	ow-up surv	ey of 2011 amo	ong children				
		(1993)				\mathbf{P} and $\mathbf{h} = \mathbf{h} + \mathbf{h}^2$ test of equation							
District	District Community Examined (no.) Positive (no.) Mf pos					positive (%) District Community Examined (no.) Positive (no.) Mf positive (%)							
Zombo	Patek Athele	11	9	81.8	Zombo	Patek Athele	19	0	0.0	< 0.0001			
Zombo	Abilambe	11	4	36.4	Zombo	Abilambe	25	2	8.0	< 0.041			
Nebbi	Agweci	9	7	77.8	Nebbi	Agweci	24	3	12.5	< 0.0001			
Zombo	Nyandima	4	3	75.0	Zombo	Anguru*	19	7	36.8				
Zombo	Ukongo	11	11	100.0	Zombo	Pachen*	20	2	10.0				
Zombo J	upa Ngali Upper	r 12	12	100.0	Nebbi	Oloamura*	38	6	15.8				
					Arua	Kairo*	25	4	16.0				
Overall		58	46	79.3			170	24	14.1	< 0.0001			

* No baseline data.

TABLE 5

Monuny parous and infection (incroscopy) rate of 3. neaver spp. in Nyagak-Bondo onchocerclasis rocus during 2010											
Month	Total collected and freshly	Parous flies	Parous rate (%)	No. positive for any of the	Infection rate (%)	No. positive L3s (%					
	dissected	1 arous mes	1 arous rate (70)	larval stages	infection rate (70)	infective rate)					
June	261	177	67.8	3	1.15	0					
September	13	0	0.0	0	0.00	0					
October	52	19	36.5	2	3.85	0					
November	45	12	26.7	1	2.22	0					
December	30	4	13.3	0	0.00	0					
Overall	401	212	52.9	6	1.50	0					

Monthly parous and infection (microscopy) rate of S. neavei spp. in Nyagak-Bondo onchocerciasis focus during 2010



Figure 1

