

# Diversity of human African trypanosomiasis epidemiological settings requires fine-tuning control strategies to facilitate disease elimination

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**Abstract:** In 2001, the World Health Organization (WHO) established a public–private partnership to fight human African trypanosomiasis (HAT). As a result of this continuous collaboration, and in addition to the coordination with nongovernmental organizations and bilateral cooperation agencies, the number of new cases of HAT annually reported by the WHO has strikingly decreased. In 2012, HAT was included in WHO’s roadmap on neglected tropical diseases with a 2020 target date for elimination. Although the prevalence of HAT is decreasing and its elimination is targeted, control approaches must be adapted to the different epidemiological patterns in order to adopt the most adequate strategies to maintain their cost-effectiveness. These strategies must be flexible and dynamic in order to be adapted to the disease progression, as well as to the changes affecting the existing health facilities in transmission areas, including their accessibility, their capabilities, and their involvement in the elimination process. Considering the different patterns of transmission (*Trypanosoma brucei* (*T.b.*) *rhodesiense* HAT) and transmission intensity (*T.b. gambiense* HAT), different settings have been defined. In the case of *T.b. rhodesiense*, this form exists primarily where wild animals are the main parasite reservoir, and where the main parasite reservoir is cattle. In *T.b. gambiense*, this form exists in areas with high intensity transmission, areas with moderate intensity transmission, and areas with low intensity transmission. Criteria and indicators must be established to monitor and evaluate the actions implemented toward the elimination of HAT.

**Keywords:** human African trypanosomiasis, HAT, sleeping sickness, disease elimination, *T.b. rhodesiense*, *T.b. gambiense*

## Introduction

In 2001, the World Health Organization (WHO) and partners established a public–private partnership to fight human African trypanosomiasis (HAT). Since then, critical challenges have been overcome, mainly in terms of availability of drugs, technical and logistical support to disease endemic countries, harmonization of control procedures, monitoring of screening activities, and coordination with concerned nongovernmental organizations (NGOs) and bilateral cooperation agencies. That has led to both the improvement of epidemiological knowledge,<sup>1</sup> and to the 75% decline of new cases that are annually reported to the WHO.<sup>2</sup> In 2011, 6631 new cases of Gambiense and 112 new cases of Rhodesiense HAT have been reported for a total number of 6743 new cases (Table 1).

Representatives from disease-endemic countries gathered in Geneva in May 2007, having noted the general trend in the reduction of new reported cases of HAT associated

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**Table 1** The number of new cases of human African trypanosomiasis (*T.b. gambiense* and *T.b. rhodesiense*) reported to the WHO (2001–2011)

	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011
<b><i>T.b. gambiense</i></b>											
Angola	4577	3621	3115	2280	1727	1105	648	517	247	218	154
Benin	0	0	0	0	0	0	0	0	0	0	0
Burkina Faso	0	0	0	0	0	0	0	0	0	0	0
Cameroon	14	32	33	17	3	15	7	13	24	15	15
Chad	138	715	222	483	190	276	97	196	510	232	276
Central African Republic	718	572	539	738	666	460	654	1194	1054	395	132
Congo	894	1005	717	873	398	300	189	182	87	87	61
Côte d'Ivoire	92	97	68	74	42	29	13	14	8	8	10
Democratic Republic of Congo	17,322	13,853	11,481	10,369	10,269	8023	8162	7326	7183	5629	5595
Equatorial Guinea	17	32	23	22	17	13	15	11	7	8	1
Gabon	30	26	26	49	53	31	30	24	14	22	17
Gambia	–	–	–	–	–	–	–	–	–	–	–
Ghana	0	0	0	0	0	0	0	0	0	0	0
Guinea	72	132	130	95	94	48	69	90	79	68	57
Guinea Bissau	–	–	–	–	–	–	–	–	–	–	–
Liberia	–	–	–	–	–	–	–	–	–	–	–
Mali	0	0	0	0	0	0	0	0	0	0	0
Niger	–	–	–	–	–	–	–	–	–	–	–
Nigeria	14	26	31	10	21	3	0	0	0	2	3
Senegal	–	–	–	–	–	–	–	–	–	–	–
Sierra Leone	–	–	–	–	–	–	–	–	–	0	–
South Sudan <sup>a</sup>	1919	3121	3061	1742	1853	789	469	623	373	200	266
Togo	0	0	0	0	0	0	0	0	0	0	0
Uganda	310	604	517	378	311	290	120	198	99	100	44
Total reported	26,117	23,836	19,963	17,130	15,644	11,382	10,473	10,388	9685	6984	6631
<b><i>T.b. rhodesiense</i></b>											
Botswana	–	–	–	–	–	–	–	–	–	–	–
Burundi	–	–	–	–	–	–	–	–	–	–	–
Ethiopia	–	–	–	–	–	–	–	–	–	–	–
Kenya	10	11	0	0	0	1	0	0	1	0	0
Malawi	38	43	70	48	41	58	50	49	39	29	22
Mozambique	–	1	–	1	–	–	–	–	–	–	–
Namibia	–	–	–	–	–	–	–	–	–	–	–
Rwanda	–	–	–	–	–	–	–	–	–	–	–
Swaziland	–	–	–	–	–	–	–	–	–	–	–
United Republic of Tanzania	277	228	113	159	186	127	126	59	14	4	1
Uganda	426	329	338	335	473	261	119	138	129	112	82
Zambia	4	5	15	9	7	6	10	13	4	8	3
Zimbabwe	–	–	–	–	3	–	–	0	3	2	4
Total reported	755	617	536	552	710	453	305	259	190	155	112
<b>Total <i>T.b. gambiense</i> and <i>T.b. rhodesiense</i></b>											
Total reported	26,872	24,453	20,499	17,682	16,354	11,835	10,778	10,647	9875	7139	6743

**Note:** <sup>a</sup>Data for South Sudan in 2009 include a small revision (three fewer cases) of previously published figures due to a reassessment of the epidemiological data.

**Abbreviation:** *T.b.*, *Trypanosoma brucei*.

with improved surveillance system and the extended coverage of control activities; these representatives concluded that HAT could be eliminated as a public health problem.<sup>3</sup>

In 2011, the elimination of HAT was considered by the WHO Strategic and Technical Advisory Group on Neglected Tropical Diseases as a feasible goal, and this was included in WHO's roadmap on neglected tropical diseases with a 2020 target date for its elimination as a public health problem.<sup>4</sup>

In 2012, different partners including major pharmaceutical companies, the World Bank, research institutes, charitable organizations, and government development agencies of the United Kingdom and the United States of America announced the implementation of collaborative measures that were included in the London Declaration on Neglected Tropical Diseases to support the objectives and to accelerate the achievements of the WHO's roadmap.<sup>5</sup>

## Findings

The generic term, “HAT,” covers the different epidemiological situations associated with the two different forms of the disease. Control approaches need to be defined in a clear and orderly fashion in order to adopt the most adequate strategies for the control and elimination of the disease.

In the foci where *T.b. rhodesiense* prevails, two different scenarios are considered:

1. Transmission areas where wild animals, mainly kept in natural protected areas, are the main parasite reservoir. In these areas, people most at risk are those living within or around national parks and game reserves where the risk of HAT exposure is related to the movement of animals and tsetse flies arise out of reserves, especially during particular seasons of the year. Also at risk are individuals who move into the protected areas mainly for collecting firewood or honey, and for fishing or poaching. Tourists and people involved in wildlife conservation or hunting activities (eg, rangers, guides, and so on) are also considered to be at risk.<sup>6</sup> These areas where Rhodesiense HAT transmission occurs can be found around natural protected areas in southwestern Uganda, western and northern Tanzania, southern Kenya, Zambia, Malawi, and northern Zimbabwe.
2. Transmission areas where the main parasite reservoir is cattle are also at risk. Cattle often graze around villages and farming areas. Moreover, they are frequently kept in villages during the night. Occasionally, they are moved to local markets for trading. Thus, people at risk of contracting HAT in these areas are mainly cattle keepers, although all villagers are also exposed to a certain level of risk.<sup>7</sup> We found these areas of Rhodesiense HAT transmission patterns in southeastern Uganda and western Kenya. In some areas, livestock and wildlife coexist, and a mixed transmission pattern is observed (eg, in western Tanzania).

A subacute clinical form of Rhodesiense HAT has been described in Zambia and Malawi,<sup>8,9</sup> which could hint toward the role of human beings as a potential reservoir, thus humans may be involved in the transmission of the disease. In this case, active case finding could be considered in the control of this disease. However, the usual clinical presentation of Rhodesiense HAT is acute, leading the patient to seek health assistance soon after infection. As a consequence, the most effective approach in tackling Rhodesiense HAT would be the reinforcement of a HAT diagnosis, and enhancing the treatment capabilities of existing health care facilities in transmission areas. This would ensure the rapid identification and treatment of cases, which coupled with vector control,

would decrease vector density and subsequently diminish human–vector contact. The treatment of cattle with antitrypanosomal drugs combined with their elective insecticide impregnation has been shown to be effective for Rhodesiense HAT control by reducing the domestic animal reservoir and the vector density.<sup>10</sup>

Across the areas in which *T.b. gambiense* prevails, three scenarios are considered. Although some domestic and wild animals have been described as hosting parasites with an enzymatic profile comparable with *T.b. gambiense*,<sup>11,12</sup> it is uncertain whether these animals can host the parasite long enough to play an important role in the transmission of Gambiense HAT.<sup>13</sup> Therefore, humans are considered to be the main reservoir of this parasite. Consequently, differences between Gambiense HAT foci are not based on disease transmission patterns but on disease transmission intensity:<sup>14</sup>

1. Areas with high intensity transmission. These can be defined as areas where the average annual number of cases reported during the past 10 years is at least one per 1000 inhabitants. These areas are found in Chad (southern region); in the Central African Republic (northwestern and southeastern regions); Equatorial Guinea (southwestern region); Gabon (northwestern region); Congo (central and southern regions); the Democratic Republic of the Congo (in northern, northwestern, central, and eastern parts); Angola (northwest region); South Sudan (southwestern region); and Uganda (northwestern region).
2. Areas with moderate intensity transmission. These are areas where at least one case per 10,000 inhabitants has been reported yearly during the past 10 years. These areas are mainly found in the regions surrounding previously described high-intensity risk areas and in coastal Guinea, Côte d’Ivoire (central region), Nigeria (southwestern region), and Cameroon (southwestern, northwestern, and eastern regions).
3. Areas with low intensity transmission. These are areas where less than one case per 10,000 inhabitants per annum has been observed during the past 10 years. They include areas around moderate intensity transmission zones, plus a few areas in northwestern and southeastern Guinea, western and southeastern Côte d’Ivoire, southeast Sierra Leone, and central and southeastern Cameroon.

In addition to the progress on the knowledge of Gambiense HAT distribution, there are still some areas that need further investigation, such as known active foci in regions that are characterized by difficult access due to topography or security constraints. Furthermore, foci which

have not reported cases in the last decades and that have no effective surveillance system in place need to be assessed. Despite these limitations, the knowledge gap between the true and the reported number of cases has been significantly improved in recent years. An estimated multiplying factor of 12 was applied to the knowledge gap in 1995,<sup>13</sup> dropping more recently to three.<sup>15</sup>

Various estimates of the number and distribution of people at risk for both forms of the disease have been provided elsewhere (Figure 1).<sup>16</sup> Precise geographical details of disease transmission areas are displayed in a specific atlas.<sup>1</sup>

To combat Gambiense HAT, it is necessary to appropriately combine the three classical control methods (ie, active case-finding through mobile teams, passive case-finding involving available health facilities, and vector control). The selection of the most appropriate combination and “dosage” of each method has to rely on: (1) the precise understanding of the epidemiological setting, including geographical and demographical data; (2) the accessibility and capabilities of the existing health facilities; and (3) vector knowledge including the sites where vector control must be applied, and methods that have to be utilized.

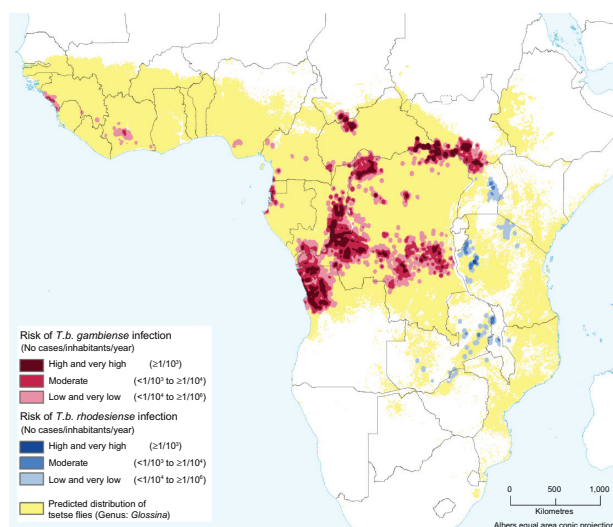
The sustainability of the control operation will depend upon the cost-effectiveness of the combination implemented. Sustainability also relies on the capacity of control programs to adapt the combination and “dosage” of the methods to the epidemiological evolution of the disease. This would avoid past situations where control and surveillance activities were

reduced instead of adapted to the prevailing situation, which led to a dramatic resurgence of cases.<sup>17</sup>

Considering the remoteness of most foci, the moderate and unspecific clinical signs and symptoms, as well as the lack of efficient health facilities in some foci, it is necessary to continue to maintain mobile teams. However, the frequency of mobile team visits and the localities to be surveyed will vary according to the intensity of the observed disease transmission in each village. In areas with high levels of transmission, mobile teams can perform visits at least once a year to the concerned localities. In areas with moderate levels of transmission, mobile teams can visit villages in the area at least every other year. Areas with low intensity transmission are not considered eligible for regular active screening unless a targeted “reactive screening” is recommended due to the reactivating of a transmission area that was detected during the surveillance process.<sup>18</sup> To implement this type of planning, a monitoring and evaluation system backed by an accurate and up-to-date mapping of disease distribution at the village level is required.

The involvement of the health system in HAT control has to be pondered across all stages. Indeed, when a mobile team visits a village, first, it does not examine the entire population, and second, it usually stays 1 day. Therefore, existing health facilities in the area are required to guarantee health care not only during the intervention periods to ensure that actively detected cases will be treated, but also between mobile team visits to ensure that the continuous diagnosis and treatment of cases that are passively detected occur. When the use of mobile teams is not considered to be the most appropriate approach, health facilities have to be able to detect HAT cases among individuals coming to seek health care services. Therefore, regardless of the intensity of transmission, the health care system must be involved in all areas where Gambiense HAT is present; what will vary is the shared responsibility between health services and mobile teams in areas of high and moderate transmission, and the sole responsibility held by health services in areas associated with low transmission rates.

The contribution of vector control to fight Gambiense HAT is basically aimed at reinforcing medical actions that are focused on the reduction of the human reservoir. Therefore, medical teams have to identify areas where diagnostic and treatment activities are not delivering the expected results, leaving human reservoirs untreated and able to maintain the transmission of the disease. This could occur due to very high transmission rates or due to low screening coverage of a given population; it could also occur due to the low



**Figure 1** Various estimates of the number and distribution of people at risk for *T.b. gambiense* and *T.b. rhodesiense* infection.<sup>16</sup>  
**Abbreviation:** *T.b.*, *Trypanosoma brucei*.

sensitivity levels of the serological and parasitological tools used. Actions based on medical approaches could also fail due to healthy human carriers that are either not detected as having the disease, or who refuse treatment.<sup>19</sup> Vector control staff members have to identify appropriate tse-tse control methods according to the vector species and the ecological setting. Effective vector control methods involving the community will have more chances of sustainability.

## Conclusion

Human African trypanosomiasis is not only a focal disease, but the foci have different patterns of transmission (Rhodesiense HAT) and transmission intensity (Gambiense HAT). When elimination is targeted and the prevalence of the disease is decreasing, control strategies must be adapted to maintain their cost-effectiveness; thus, strategies must be flexible and dynamic enough to be adapted to the disease progression, as well as to the changes affecting the existing local health services, including their level of preparedness and their involvement in the elimination process.

The classical controversy between enthusiasts of “vertical” approaches through mobile teams and those of “horizontal” approaches through primary health care services is losing its relevance since both approaches are complementary and have their specific and synergic roles in the elimination process.

Current control tools are not the most appropriate when designing fine-tuned strategies for each epidemiological setting identified. Continuous research is needed to improve the current weaknesses of the control tools currently used, mainly to engage health services in HAT elimination.

In the process of eliminating HAT, criteria and indicators must be established to monitor and evaluate the actions implemented.

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## Disclosures

The authors report no conflicts of interest in this work. The boundaries and names shown and the designations used on

the maps presented in this paper do not imply the expression of any opinion whatsoever on the part of the WHO concerning the legal status of any country, territory, city, or area, or of its authorities, or concerning the delimitation of its frontiers or boundaries. The views expressed in this paper are those of the authors and do not necessarily reflect the views of the WHO.

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