OPEN LETTER

Insights from quantitative and mathematical modelling on the proposed WHO 2030 goals for Chagas disease [version 1; peer review: 2 approved]

Collaborating Group on Chagas Disease Modelling

Abstract

Chagas disease (CD) persists as one of the neglected tropical diseases (NTDs) with a particularly large impact in the Americas. The World Health Organization (WHO) recently proposed goals for CD elimination as a public health problem to be reached by 2030 by means of achieving intradomiciliary transmission interruption (IDTI), blood transfusion and transplant transmission interruption, diagnostic and treatment scaling-up and prevention and control of congenital transmission. The NTD Modelling Consortium has developed mathematical models to study Trypanosoma cruzi transmission dynamics and the potential impact of control measures. Modelling insights have shown that IDTI is feasible in areas with sustained vector control programmes and no presence of native triatomine vector populations. However, IDTI in areas with native vectors it is not feasible in a sustainable manner. Combining vector control with trypanocidal treatment can reduce the timeframes necessary to reach operational thresholds for IDTI (<2% seroprevalence in children aged <5 years), but the most informative age groups for serological monitoring are yet to be identified. Measuring progress towards the 2030 goals will require availability of vector surveillance and seroprevalence data at a fine scale, and a more active surveillance system, as well as a better understanding of the risks of vector re-colonization and disease resurgence after vector control cessation. Also, achieving scaling-up in terms of access to treatment to the expected levels (75%) will require a substantial increase in screening asymptomatic populations, which is anticipated to become very costly as CD prevalence decreases. Further modelling work includes refining and extending mathematical models (including transmission dynamics and statistical frameworks) to predict transmission at a sub-national scale, and developing quantitative tools to inform IDTI certification, post-certification and re-certification protocols. Potential perverse incentives associated with operational thresholds are discussed. These modelling insights aim to inform discussions on the goals and treatment guidelines for CD.

Keywords

Chagas disease, WHO guidelines, Elimination as a public health problem, intradomiciliary transmission interruption, trypanocidal treatment, NTD Modelling Consortium
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The Collaborating Group on Chagas Disease Modelling (CGCDM) gives a valuable review of the accomplishments and open challenges of quantitative and mathematical modelling of Chagas disease (CD). CGCDM provides a refreshingly frank critique of some goals of the World Health Organization (WHO), describing vector control in areas with sylvatic triatomine vectors as "not feasible in a sustainable manner" and the stated treatment goals as "not feasible" at all. The review provides convincing quantitative evidence for the latter assessment. It does not cite quantitative estimates to show that vector control in the face of sylvatic recolonization is unsustainable. The review would benefit from a more detailed quantitative description of what vector control efforts would be feasible in regions with multiple sources of recolonization, including sylvatic and between villages (e.g. Gürtler et al. 2007). The assessment of CGCDM is likely to be useful to scientists who want to address critical gaps in modelling Chagas disease, to officers of WHO and national governments who seek to set realistic goals and make informed policy about CD control, and to funders in search of fact-based counsel on the opportunities and risks of supporting modelling research and field action to control CD.

CGCDM strongly urges combining current vector control strategy with trypanocidal treatment of infected people. CGCDM identifies the enormous advances that will be required to achieve trypanocidal treatment of infected people. These advances include improved access to screening and diagnosis (currently <1% of individuals at risk have access in Colombia and perhaps elsewhere, according to previous sources and CGCDM), highly sensitive and highly specific tests to diagnose infection with Trypanosoma cruzi, more efficacious and less toxic drugs or other treatments than are currently available, and persuading people to complete the course of treatment. CGCDM also candidly identifies some of the risks of its own recommended strategy.
Based on experience in CD vector and disease control in the semiarid and humid Chaco of northern Argentina, currently the main hotspot of vector-borne transmission, we suggest three additional ingredients that may enhance the recipes advanced by CGCDM for interrupting vector-borne transmission of *T. cruzi* to people. We believe that these ingredients would prove valuable to CD control in regions of vector-borne transmission beyond the Argentine Chaco, but that remains to be proved. These ingredients are: (1) engaging the families and communities affected by CD in protecting themselves from CD, and modelling their engagement; (2) enriching the picture of CD dispersal using genetic and other modern techniques, and modelling that more complex picture; and (3) improving the rationality and statistical efficiency of vector control, and modelling the resulting impacts on parasite transmission and economic savings from improved vector detection and control. Our suggestions do not address other important problems CGCDM addressed, such as interrupting transmission of *T. cruzi* by blood transfusion and transplants, increasing access to expanded diagnostic and treatment programs, and preventing and controlling congenital transmission. We now sketch each proposed additional ingredient.

(1) Engaging the communities and families affected by CD in protecting themselves

CGCDM do not mention the roles of the individuals, families, and communities affected by CD in defending themselves. We think these roles could be influential, perhaps even crucial, in controlling and monitoring CD, and achieving the 2030 targets identified in Table 1. Specifically, people living in areas where vector-borne CD is endemic can influence wall and roof construction and maintenance, domestic animal management, on-demand residual insecticide spraying, and monitoring any resurgence of bugs after official interventions (e.g., Monroy et al. 2009). Community participation and training are key to achieving a highly demanding goal such as “0% colonization rate of dwellings” in the affected zones.

In the Gran Chaco region, a single village is typically composed of multiple house compounds. Each house compound consists of one (or more) domiciles (for human residents) and multiple outbuildings, such as chicken coops, goat corrals, kitchens, grain stores, and outhouses. Domiciles of different house compounds, though constructed with the same basic materials (mud bricks and poles), vary widely in the state of repair of the walls. Walls smoothly plastered provide no refuge for the triatomine vectors of CD and the human residents typically suffer lower incidence of *T. cruzi*. By contrast, houses with cracked walls shelter the bugs, have higher triatomine infestation levels (Cecere et al. 2002; Bustamante et al. 2009), and the human residents are more likely to be infected with *T. cruzi* across settings and vector species (e.g., Mott et al. 1978, Andrade et al. 1995, Bonfante-Cabarcas et al. 2011). Similarly, house compounds that have thatched roofs provide good refuges for bugs. Domiciles roofed with materials less hospitable to resting bugs usually have lower prevalence and incidence of *T. cruzi*. (Gürtler et al. 1998, Black et al. 2007). Domestic dogs have a very high prevalence of infection with *T. cruzi* and a very high probability of transmitting *T. cruzi* to uninfected bugs. When bugs thus infected are in human sleeping quarters, they have an increased likelihood of transmitting *T. cruzi* to humans. Keeping domestic dogs out of human sleeping quarters permanently could substantially reduce the incidence of human infection, according to modelling (Cohen and Gürtler 2001) and observations in Argentina (Gürtler et al. 2005, Cardinal et al. 2018). In Venezuela, the household presence of several domestic animal species (dogs, caprines, armadillos and chickens) is significantly and positively associated with human seroprevalence (Bonfante-Cabarcas et al. 2011).

Because of the long asymptomatic period of CD, residents in endemic areas often have a low awareness of the risks associated with *Triatoma infestans* infestation. Local residents could be educated about the risks of bug-mediated transmission of *T. cruzi* to their own health and the health of their children. They
could also be taught how to apply residual insecticides (individually purchased or provided by control agencies) as need arises within their domiciles and outbuildings, and to monitor the resurgence of bugs after control measures (Cecere et al. 2019, Gaspe et al. 2018). Thatch-roofed granaries are not usually monitored or sprayed, may often harbor large bug populations, and could be a potent source of bug reinestation of other structures (Cohen et al. 2017).

Modelling challenges are to synthesize what is known about the magnitudes of these effects and to compare the costs and benefits of house-to-house and community-wide education about walls, roofs, domestic dogs, spraying, and monitoring with the costs and benefits of more traditional top-down interventions. The latter have proven difficult to sustain in the long run, especially when intrusive triatomine species tend to invade domestic premises and pose a recurrent risk of recolonisation, such as with *Triatoma dimidiata* in Central America (Peterson et al. 2019) and *Rhodnius prolixus* in Colombia and Venezuela (Sanchez-Martin et al. 2006, ref. 12).

(2) Enriching the picture of CD vector dispersal

Based on modelling studies, CGCDM "suggest that a better comprehension of vectorial transmission in rural and urban settings would require understanding and quantifying of two different forms of vector dispersal, namely, dispersal between sylvatic and non-sylvatic habitats and diffusive dispersal within cities." We agree with the need to understand both forms of dispersal. We propose that additional levels of dispersal need empirical studies and modelling.

Within a house compound, bugs disperse among domiciliary and peri-domiciliary structures. Within a village, bugs disperse among house compounds. Within localities or clusters of villages, bugs disperse among villages possibly through flight and possibly through movements of people, non-human animals, and goods. At all three levels (house compound, village, cluster of villages), bugs may disperse between sylvatic and settled habitats, and all forms of bug dispersal may depend on the season (Dohna et al. 2007, 2009; Gourbière et al. 2009). Independently of the dispersal of bugs, people (and their dogs and cats) carry infections of *T. cruzi* among house compounds, villages, localities, urban and rural regions of a given country, and internationally. Four multi-level networks of dispersal interact: bugs, humans, domestic animals, and *T. cruzi* (in bugs, people, and domestic animals).

In future empirical work, genetic techniques to infer the origins and spread of infections (Biek et al. 2012; Gourbière et al. 2012; Gire et al. 2014; Grad et al. 2014) need to be tailored and applied to the origins and transmission of *T. cruzi* infections and the dispersal of the vectors. For example, Nouvellet et al. (2013) used the incidence of *T. cruzi* infections in humans and the average number of potentially infective contacts per uninfected person to estimate that the probability of *T. cruzi* transmission per potentially infective contact is 1 in 900-4000 contacts with infected bugs. If genetic techniques were used to identify markers or "barcodes" in *T. cruzi*, the appearance of these markers in people newly infected with *T. cruzi* could be used to confirm or refine these estimates.

Modelling challenges are to model the structures and infer the parameter values of these networks of interactions using available data (e.g., Dohna and Pineda-Krch 2010) and to design additional practical data collection goals for improved modelling and estimation.

(3) Improving the rationality and statistical efficiency of vector control by spraying insecticides

A decade ago, Dohna et al. (2009, p. 1) made a statement that was true then and unfortunately remains
true today: "… it is important to increase the efficiency of vector control. It is currently unknown what spatial and temporal pattern of repeated insecticide application maximizes its efficiency." Some modeling studies (reviewed in Nouvellet et al. 2015\textsuperscript{27}) produced concrete recommendations for optimizing vector control but none of these recommendations has been tested in field trials. Large-scale field trials that test well-founded model-based recommendations for cost-effective vector control are crucial, especially for areas where resources are limited and sustained vector control is required.

Mass spraying of residual insecticides in a community is traditionally triggered when the community-wide fraction of bug-infested houses (the house infestation rate) detected by a standardized search procedure exceeds a fixed threshold, typically 5% (Schofield 1994\textsuperscript{28}; Guhl 2005\textsuperscript{29}, p. 395; Aiga et al. 2012\textsuperscript{30} suggest a threshold of 8% for their region). The number of houses to be searched is rarely defined in advance, or all houses in a village are searched, which is costly and time-consuming.

A more efficient alternative could be to design a plan of sequential sampling that stops searching of houses and starts spraying when the probability that the infestation rate exceeds 5% reaches a chosen standard. For example, suppose the probability of house infestation is 5% or less. If houses are chosen randomly from a list of houses in the village, then the probability that the first three houses are all infested is at most \( (0.05)^5 = 0.000125 \), just over one chance in ten thousand. In this case, it would seem superfluous to search additional houses for bug infestation before deciding to spray the entire community. The specification in this example that "houses are chosen randomly from a list of houses in the village" is important, because if the inspectors choose geographically contiguous houses for their own convenience rather than randomly, there may be dependence in the infestation of successive neighboring houses. For example, three successive houses might be infested as a result of their proximity or shared familial indifference to plastering walls, regardless of the house infestation rate of the community. Sequential plans for detecting pest infestation rates above an economic threshold are commonly used in commercial agriculture. Similar procedures need to be developed and adapted for the efficient surveying of house infestations in the control of CD. Sampling design is of prime relevance, especially when control programs pursue certification of transmission interruption or vector elimination.

Modelling challenges include designing sequential sampling plans that take account of the consistent relationships between the mean and the variance of bug relative population sizes in different habitats within house compounds (Taylor's law; Cohen et al. 2017\textsuperscript{15}); modelling and measuring the spatial aggregation of house infestation; and estimating the economic savings resulting from more efficient vector detection and control.

The paper ably draws attention to some undesirable consequences of the certification of IDTI witnessed during the post-certification period in several countries. In general, CD vector control programs have lost substantial operational capacity. The (re)emergence and growing relevance of dengue and of other mosquito-borne viruses such as zika and chikungunya have drawn away from CD the scarce resources available for vector control. This complex scenario in part explains why ten countries have yet to hit the goal of IDTI by 2020 (Tarleton et al. 2014\textsuperscript{31}). Competition for scarce resources for vector control remains a major obstacle to hitting the 2030 targets.

The following passages contain some inaccuracies or imprecise language and should be reviewed.

In the Abstract, "Modelling insights have shown that IDTI is feasible in areas with sustained vector control programmes and no presence of native triatomine vector populations." An interested reader not informed of the main issue under discussion may not understand the underlying logic of this sentence. If there is no presence of native triatomine populations, neither IDTI nor sustained triatomine control programmes
would be pertinent, unless there are other (non-native) vectors that pose a risk of human infection with *T. cruzi*.

The Abstract begins: "Chagas disease (CD) persists as one of the neglected tropical diseases (NTDs) with a particularly large impact in the Americas" and the Background section on page 3 continues: "CD is a parasitic disease caused by the protozoan *Trypanosoma cruzi*, and transmitted mainly by domiciliated triatomine (Reduviidae) vectors (kissing bugs) in tropical areas of the Americas." In fact, the trypanosome that causes CD is transmitted mainly by domiciliated triatomine (Reduviidae) vectors (kissing bugs) in tropical, subtropical and temperate areas of the Americas. CD is not restricted to tropical areas.

The Background section, page 3, states: "...along with the urbanization process in recent decades, other transmission routes, such as blood transfusion, organ transplant and congenital have become important in both endemic and non-endemic countries." This generalization is partially inaccurate. In ref. 2, Table 2 shows no transfusion-related CD case in 21 Latin American endemic countries. This is quite plausible because screening of blood donors has achieved virtually universal coverage decades ago, with rare exceptions. A 2015 WHO publication, based on 2010 data provided by the countries, reported that "19 out of the 21 endemic countries achieved 100% screening of donated blood." The issue of transfusion-mediated transmission has been or still is pertinent to some non-endemic, developed countries.

The Background section, page 3, also states: "So far, 11 out of the 21 recognised endemic countries have been certified as having reached intradomiciliary transmission interruption (IDTI)." This statement inaccurately suggests that there is no vector-borne domestic transmission of *T. cruzi* in 11 countries. The certifications of IDTI for Central American countries are relevant for *R. prolixus*, not for *T. dimidiata*, a widespread vector involved in domestic transmission to humans (Peterson et al. 2019). Similarly, the certifications for the Southern Cone countries are relevant for *T. infestans*, but in Brazil there are other species involved in domestic transmission.

On page 4, a reference is needed to justify: "the number of triatomines/house was fitted to data from various locations prior to vector control and applied to cases where there was only one triatomine species present in the dwellings, as well as to situations with mixtures of species and developmental stages, various types of houses and bug densities per house."

Also on page 4: "... In the presence of sylvatic populations, there is a continuous introduction and colonisation of domiciliary and peri-domiciliary habitats; in these areas, traditional vector control is not feasible in a sustainable manner." A better distinction between feasibility, effectiveness and sustainability is relevant. Many decades ago, in some of these areas, such as in Venezuela, traditional vector control was feasible and very effective by current standards when it was pursued consistently (Feliciangeli et al. 2003, Bonfante-Cabarcas et al. 2011, Bartsch et al. 2017). The decentralization of healthcare services, beginning in the late 1980-1990s in Latin America, combined with other regional processes, virtually eliminated or downsized the existing vector control programs except in a few countries.

Also on page 4: "Various studies on routine vector surveillance have demonstrated that the currently used methods have low sensitivity and greatly underestimate vector density, infestation and infection rates; vector surveillance may be capturing half of infestations – and, most likely, most bugs within a house." "..."most bugs within a house" suggests that the methods may be capturing most of the bugs within the house, which disagrees with the former sentence and field observations. Perhaps the sentence intended to say that vector surveillance was detecting half of infestations, and standard vector collection methods captured mostly bugs within a house, i.e., domiciles.
On page 6: "Trypanocidal treatment with benznidazole (BNZ) or nifurtimox (NFX) has been aimed at both reducing parasitaemia and curbing disease progression. So far, there is limited evidence on the efficacy of drugs for these." There is substantial evidence on the efficacy of BNZ in suppressing or reducing parasitaemia as determined by RT-PCR (e.g., ref. 24, 28), whereas the available evidence on curbing disease progression is limited and debatable (e.g., ref. 29). The age of the patient under treatment is important: both drugs are able to achieve the serological and parasitological cure of recent chronic infections when applied to T. cruzi-seropositive children under age 15 years (ref. 28), whereas in chronic adult patients both drugs suppressed or reduced parasitaemia albeit transiently, as determined by PCR or xenodiagnosis (ref. 24, 28); the serological effects of treatment with either drug may take many years to appear.

On page 7: "With the current tools, low access to screening is the bottleneck; achieving just 10% of successful treatment at population level will require an enormous investment on improving access to screening, especially when targeting asymptomatic populations in low prevalence settings, which currently prevail in most endemic areas. This issue is debatable and likely varies geographically. In Argentina, the key obstacle has been access to treatment in rural areas: either the drugs or the treating physicians are not available to indicate treatment and manage the adverse drug-related reactions over the 60-day follow-up (Sartor et al. 2017). Health services and several NGOs (e.g. MSF) have conducted large serosurveys of T. cruzi infection in rural areas using rapid tests or dried blood spots.

Also on page 7: "Current estimates of access to diagnostics and treatment are at <1%. The pertinent reference is Ribeiro et al. (2009), who provided the <1% estimate across endemic countries. Ref. 33 reported that access to diagnostics and treatment in Colombia was 1.2%.

A few editorial errors should be corrected. In the caption of Figure 3, the identification of the coloured lines is wrong. The colour code adjacent to the upper right corner of panel B is more likely to be correct. In line 5 of paragraph 2 on page 6, BZN should be replaced by BNZ. In the captions of Fig. 2 and 3, ref. 17 should be replaced by ref. 18. Table 1 should list the 21 endemic countries, in line with recommendations issued by PAHO/WHO (2018) and other passages in the text.

References
6. De Andrade AL, Zicker F, De Oliveira RM, Da Silva IG, Silva SA, De Andrade SS, Martelli CM:


Is the rationale for the Open Letter provided in sufficient detail?
No

Does the article adequately reference differing views and opinions?
Yes

Are all factual statements correct, and are statements and arguments made adequately supported by citations?
Partly

Is the Open Letter written in accessible language?
Yes

Where applicable, are recommendations and next steps explained clearly for others to follow?
Yes

Competing Interests: No competing interests were disclosed.


We confirm that we have read this submission and believe that we have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.