# **Trends in Parasitology**

# Neglected tools for neglected diseases: mathematical models in economic evaluations --Manuscript Draft--

Manuscript Number:	TREPAR-D-14-00138R1
Article Type:	Review
Corresponding Author:	Hugo C. Turner, PhD
	London, UNITED KINGDOM
First Author:	Hugo C. Turner, PhD
Order of Authors:	Hugo C. Turner, PhD
	Martin Walker, PhD
	Michael D French, PhD
	Isobel M Blake, PhD
	Thomas S Churcher, PhD
	María-Gloria Basáñez, PhD
Abstract:	Despite many current interventions against neglected tropical diseases (NTDs) being highly cost effective, new strategies are needed to reach the World Health Organization's control and elimination goals. Here we argue for the importance of incorporating economic evaluations of new strategies in decisions regarding resource allocation. Such evaluation should ideally be conducted using dynamic transmission models that capture inherent nonlinearities in transmission and the indirect benefits ('herd effects') of interventions. A systematic review of mathematical models that have been used for economic analysis of interventions against the ten NTDs covered by the London Declaration reveals that only 16 out of 49 studies used dynamic transmission models, highlighting a fundamental - but addressable - gap in the evaluation of interventions against NTDs.

# Imperial College London

Department of Infectious Disease Epidemiology School of Public Health Faculty of Medicine Imperial College London Norfolk Place, London W2 1PG, UK Tel: +44 (0) 20 7594 3217

Hugo Turner PhD hugo.turner06@imperial.ac.uk 1<sup>st</sup> October 2014

Danielle Loughlin, Editor Trends in Parasitology

Dear Dr Loughlin,

Thank you very much for considering our manuscript entitled **"Neglected tools for neglected diseases: mathematical models in economic evaluations**".

We are very grateful to the referees and editors for their comments, and have endeavoured to address them in the revised manuscript which we now submit.

We look forward to hearing from you.

Yours sincerely,

Hugo Ture

Dr Hugo C. Turner (corresponding author on behalf of all authors)

Click here to download Manuscript: TREPAR Turner et al Neg tools for NTD Final FVR comments REVISON.docx

- Title Neglected tools for neglected diseases: mathematical models in economic evaluations
- Foot line: Neglected tropical diseases and economic evaluations:
- Corresponding Author's Institution: Imperial College London, Faculty of Medicine, School of Public Health

- List of other Authors
- Hugo C. Turner<sup>1,2</sup>, Martin Walker<sup>1,2</sup>, Michael D. French<sup>2,3</sup> Isobel M. Blake<sup>2,4</sup>, Thomas S.
- Churcher<sup>2</sup> and María-Gloria Basáñez<sup>1,2</sup>
- <sup>1</sup>London Centre for Neglected Tropical Disease Research, Department of Infectious Disease
- Epidemiology, School of Public Health, Faculty of Medicine (St. Mary's Campus), Imperial
- College London, Norfolk Place, London W2 1PG, UK
- <sup>2</sup> Department of Infectious Disease Epidemiology, School of Public Health, Faculty of
- Medicine (St. Mary's Campus), Imperial College London, Norfolk Place, London W2 1PG, UK
- <sup>3</sup> Schistosomiasis Control Initiative, Department of Infectious Disease Epidemiology, School
- of Public Health, Faculty of Medicine (St. Mary's Campus), Imperial College London,
- Norfolk Place, London W2 1PG, UK
- <sup>4</sup> MRC Centre for Outbreak Analysis & Modelling, Department of Infectious Disease
- Epidemiology, School of Public Health, Faculty of Medicine (St. Mary's Campus), Imperial
- College London, London W2 1PG, UK

1 Summ	ary
--------	-----

3 Despite many current interventions against neglected tropical diseases (NTDs) being	y highly
---	----------

- 4 cost effective, new strategies are needed to reach the World Health Organization's control
- 5 and elimination goals. Here we argue for the importance of incorporating economic
- 6 evaluations of new strategies in decisions regarding resource allocation. Such evaluation
- 7 should ideally be conducted using dynamic transmission models that capture inherent
- 8 nonlinearities in transmission and the indirect benefits ('herd effects') of interventions. A
- 9 systematic review of mathematical models that have been used for economic analysis of
- 10 interventions against the ten NTDs covered by the London Declaration reveals that only 16
- 11 out of 49 studies used dynamic transmission models, highlighting a fundamental but
- 12 addressable gap in the evaluation of interventions against NTDs.
- 13 Key words: NTDs; neglected tropical diseases, mathematical modelling; analysis theoretical
- 14 models; epidemiology; control programmes; cost effectiveness; cost benefit; economic
- 15 evaluation studies.

#### **1** Neglected Tropical Diseases: Reaching the 2020 control and/or elimination

2 goals

The neglected tropical diseases (NTDs) are a group of chronic, disabling, and disfiguring 3 4 conditions that occur especially among the rural poor and disadvantaged urban populations 5 [1]. These diseases cause a substantial health and economic burden on poor populations in Africa, Asia, and Latin America [1]. Latest estimations indicate NTDs cause approximately 6 7 534,000 deaths, and are responsible for 56.6 million disability-adjusted life-years (DALYs) 8 lost, each year [2-4], although these numbers are likely to be significantly underestimated [2]. 9 In January 2012 the United Kingdom Coalition against Neglected Tropical Diseases 10 subscribed to the London Declaration for the control and/or elimination of ten of the highest burden NTDs by the year 2020 (Table 1) [5]. These goals were inspired by the World Health 11 12 Organization's (WHO) 2020 road map for accelerating work to overcome the global burden of NTDs [6]. 13 14 The reports by the Disease Control Priorities Project (DCPP) include estimates of the cost effectiveness of interventions against diseases in the developing world (http://www.dcp-15 3.org/). These reports illustrate that many of the current interventions against NTDs are 16 highly cost effective and, in several instances, are even more so than interventions against the 17 18 so-called 'big three'; malaria, tuberculosis and human immunodeficiency virus (HIV) [7, 8]. 19 Despite this, it is also recognised that novel strategies and tools will be needed to reach the NTD control and elimination goals [5]. However, the majority of cost-effectiveness estimates 20 21 (such as those presented by the DCPP [8]) are only relevant to interventions currently in use in specific epidemiological and programmatic settings, and are often not informative to the 22 23 decision-making process of switching to new strategies in different settings.

1 In the context of the London Declaration, models will be needed to perform economic

2 evaluations to investigate what changes in policy are required to meet the WHO 2020 goals

3 in the most cost-effective way. The aim of this paper is to (a) highlight why this area of

- 4 research should be addressed with dynamic, rather than the more commonly used static
- 5 transmission models, and (b) to outline what we feel are key research needs for this area.

#### **6** Mathematical models for economic evaluation

7 The use of mathematical models to perform economic evaluations of new health care 8 interventions is an important tool for deciding how to allocate public-health resources [9]. 9 Accurately parameterised mathematical models have the advantage of enabling investigation of the cost-effectiveness of various strategies in a range of different epidemiological and 10 programmatic scenarios, and can therefore help optimise the use of resources. This is 11 particularly important for NTDs, which are most prevalent in resource-poor settings and 12 13 historically have suffered from a lack of funding (for control implementation and research), visibility, and political advocacy. Modellers must work effectively with funders and 14 politicians who will often want to demonstrate impact quickly rather than delaying decision 15 making while comprehensive evaluations are undertaken. Consequently, models must be 16 flexible and versatile enough to capture the effects and accompanying uncertainties of past 17 and ongoing intervention activities. 18

#### 19 *Static infection models*

The most widely used models for economic evaluations are so-called static (infection)
models, such as decision-tree and Markov models [10-13]. Such models are relatively
straightforward to develop, and tend to have low computational demands, facilitating rapid
simulations and accompanying sensitivity analyses [14]. In static models, individual hosts
acquire infection at a rate—the so-called force of infection—which is *uncoupled* from the

1 abundance of infection (i.e. for microparasites, the number and density of infected 2 individuals; for macroparasites, the number and density of parasites) in the population or 3 community as a whole [10, 15, 16]. This may be an appropriate framework for non-4 communicable diseases, but it is not for infectious diseases where the rate of infection is inextricably linked to: a) the abundance of infection among all individuals in a population 5 6 and, b) on how these individuals are connected and able to transmit infection to one another (e.g. by direct contact, by vectors, or by exposure to contaminated material in the 7 environment). 8

9 Consequently, although static models are suitable for the evaluation of interventions that do 10 not markedly affect the targeted parasites transmission cycle, such as those that relieve 11 disease symptoms and sequelae but do not treat infection, or those that target a very small 12 fraction of the vector, host, or parasite population (leaving a large infectious reservoir), they 13 are rarely suitable for evaluating interventions that reduce levels of infection transmission in 14 a population. In such circumstances, static models will underestimate the impact of an

15 intervention.

#### 16 Dynamic transmission models

Dynamic transmission models couple the rate of infection and the population abundance of 17 18 infection by explicitly modelling the transmission cycle of the disease in question [14, 15, 17]. They involve a substantial time investment to develop, can be computationally intensive 19 [14] and can be more difficult to parameterise (and consequently generally have greater 20 21 inherent uncertainty) than static models. However, and critically, dynamic transmission models capture the so-called 'herd effect' (or indirect effects) of interventions targeting 22 particular population groups. Such interventions include vaccination campaigns and school-23 24 based mass drug administration (MDA) programmes, whereby all individuals within a

1	population can benefit from the intervention, albeit in a differential manner, regardless of
2	whether they are within the target group or not. Figure 1 illustrates this concept by showing
3	that a school-based MDA programme treating children for Ascaris lumbricoides can have a
4	notable indirect (herd) benefit for the untreated adults whose worm burden is also reduced
5	over time, due to population-wide reductions in transmission. The impact of these indirect
6	herd effects will accumulate over time as reductions in transmission affect reductions in
7	incidence which, in turn, manifest more slowly in reduced population levels of infection.
8	Consequently, the extent that static models underestimate the impact of transmission reducing
9	interventions will increase over time. By the same token, static models may provide a
5	
10	reasonable approximation to dynamic models over a relatively short time frame. Therefore,
10	reasonable approximation to dynamic models over a relatively short time frame. Therefore,
10 11	reasonable approximation to dynamic models over a relatively short time frame. Therefore, accounting for the herd effect can be crucial to the validity, robustness and policy relevance
10 11 12	reasonable approximation to dynamic models over a relatively short time frame. Therefore, accounting for the herd effect can be crucial to the validity, robustness and policy relevance of conclusions drawn from cost-effectiveness evaluations of interventions against
10 11 12 13	reasonable approximation to dynamic models over a relatively short time frame. Therefore, accounting for the herd effect can be crucial to the validity, robustness and policy relevance of conclusions drawn from cost-effectiveness evaluations of interventions against transmissible diseases in general, and of NTDs in particular [14, 15, 18].
10 11 12 13 14	reasonable approximation to dynamic models over a relatively short time frame. Therefore, accounting for the herd effect can be crucial to the validity, robustness and policy relevance of conclusions drawn from cost-effectiveness evaluations of interventions against transmissible diseases in general, and of NTDs in particular [14, 15, 18]. Aside from dynamic models being general applicability and appropriate for cost-effectiveness

unbiased manner, the effectiveness of interventions which elicit very different effects on the

19 microfilaricide used to control River Blindness, elicits a pronounced yet transient reduction in

dynamics of infection/transmission [19]. For example, ivermectin which is used a potent

20 numbers of microfilariae (the parasite transmission stage) [20], whilst doxycycline, a

21 macrofilaricide, sterilises and kills adult worms, causing a more gradual but sustained

reduction in worm burden (and microfilariae) [21-23]. Furthermore, potential prophylactic

helminth vaccines [24], may induce notable long-term benefits while possibly having a

24 relatively small initial impact compared to MDA.

17

1 Second, dynamic models can be used to evaluate the possibility of infection elimination 2 under specific intervention strategies or intervention combinations (i.e. the reduction to zero of the incidence of infection caused by a specific agent in a defined geographic area as a 3 result of deliberate efforts; continued measures to prevent re-establishment of transmission 4 5 are required [25]). For this purpose, dynamic models should generally be stochastic, 6 capturing the reality of random chance in demographic and transmission processes. This is 7 because such random variation becomes increasingly important in determining whether a 8 disease is eliminated as infection is decreased to very low levels, a process known as 9 'stochastic fade-out'. By contrast, in (non-stochastic) deterministic models, all events occur in a pre-specified way depending on the parameter values and initial conditions of the model, 10 11 ignoring the effects of chance. Stochastic models are particularly relevant to NTDs as goals 12 shift from control to elimination because new interventions are likely to be more expensive than current tools and may only be deemed cost-effective if they lead to elimination faster. 13 avoiding future costs [26]. 14

15 Lastly, the transmission of macroparasites (such as the causative agents of guinea worm, 16 schistosomiasis, river blindness, lymphatic filariasis and soil-transmitted helminthiases) is often governed by density-dependent processes (Box 1) [27-29] which makes the effect of 17 control interventions on transmission often highly non-linear (Figure 2), i.e. decreasing the 18 mean parasite load by 50%, may not decrease transmission by 50%. This means that the 19 indirect benefits of treatment on the reduction in transmission will depend on the number and 20 distribution of parasites between hosts [17, 30]. Consequently, these macroparasites often 21 22 require dynamic transmission models tracking the intensity of the different parasite lifestages accurately to account for the impact of interventions against them [17]. 23 Density-dependent processes may also be important regulators of microparasite population 24 dynamics. Particularly important is the Allee effect (a positive or facilitating density-25

1 dependent process), which gives rise to a so-called 'breakpoint' population density below 2 which the infection cannot persist [31]. A study that fitted a dynamic transmission model to 3 data on the prevalence of ocular *Chlamydia trachomatis* (aetiological agent of trachoma) 4 before and after MDA in 24 Ethiopian villages indicated that the transmission probability was decreasing non-linearly—and more rapidly than would be expected without assistance from 5 6 an Allee effect—as the prevalence of infection in the community declined [32]. Thus, the 7 microparasite population would be less likely to recover after cessation of MDA, highlighting 8 the potential importance of accounting for these effects when evaluating elimination 9 strategies.

## 10 Current models used for economic evaluations of NTD interventions

We conducted a systematic review of mathematical modelling studies that include an 11 economic analysis of intervention strategies for the ten NTDs under the London Declaration 12 (Table 1). We performed a computer-assisted search of the bibliographic databases Medline, 13 ISI Web of Knowledge, and PubMed. We imposed no language or date restrictions. The 14 15 reference lists within retrieved publications were searched for articles that were not identified 16 in our database searches. The criteria for inclusion and exclusion of studies, and the search terms are shown in Figure 3. It is beyond the scope of this paper to conduct an in-depth 17 18 comparison of all studies for each disease, and thus we mainly focus on compiling a list of 19 studies, stratified by whether they employ static infection or dynamic transmission models. A review of the methodological quality (other than model type) of economic evaluations for 20 21 parasitic diseases is published elsewhere [33].

We found 49 model-based economic evaluations of NTD interventions, with half on Chagas
disease and schistosomiasis. Only 16 of the identified studies used dynamic transmission
models, the majority by the same small group of authors (Table 1). This is not to imply that

using a static model was always the wrong choice; this depends on both the intervention and
the parasite under investigation. However, it does highlight a methodological predilection,
which, although reasonable in certain circumstances, will increasingly become inappropriate
in the context of the elimination goals set by the London Declaration for many of the NTDs.
It is noteworthy that this bias towards static models for economic evaluations is not confined
to NTDs but a common theme in the economic analysis literature. A review by Kim and
Goldie (2008) on the cost-effectiveness of a range of different vaccination programmes also

8 found that most of the models used were static [14].

#### 9 A research gap: dynamic models for economic evaluations

Although static models are suitable for the evaluation of interventions that do not markedly 10 affect the targeted parasite's transmission cycle—such as when there is a large untargeted 11 12 reservoir of parasites or vectors, when evaluating over a short time horizon—they notably underestimate the impact of, and thus are inappropriate for evaluating, interventions that 13 reduce levels of infection transmission in a population, such as MDA or vaccination 14 15 campaigns. This is particularly important when investigating elimination strategies. The lack of dynamic models used for the economic evaluation of NTD interventions constitutes a 16 fundamental research gap and methodological shortcoming. This is of particular importance 17 in the context of NTDs which have comparatively little funding for research and policy 18 orientated analyses and are most prevalent in resource-poor settings. Thus, it is imperative 19 20 that appropriate analyses are conducted to guide the most cost-effective way of deploying 21 intervention tools and strategies [19]. Furthermore, it is important to consider that even with the development of new drugs and other intervention tools, along with the optimisation of 22 23 existing strategies, there is unlikely to be a 'silver bullet' which is the optimum solution in 24 every situation. A more likely reality is more complicated, and one that will require different combinations of existing strategies and the development of specific approaches depending on
local conditions. For example, in areas with high pre-control endemicity of soil-transmitted
helminths, it may be necessary to extend school-based deworming programmes to include
adults in the community [34-36]. Dynamic transmission models can be useful tools in
investigating the potential benefit and cost of different combinations of interventions in
different programmatic and epidemiological scenarios, guiding the most cost-effective use of
resources.

#### 8 The role of dynamic models in clinical trials and enhancing Target Product Profiles

9 It would be beneficial to proposed trials of new interventions to recognise early in the planning process the important analytical role of dynamic models so that suitable data can be 10 11 collected for parameterization and robust analysis and evaluation. In particular, data may need to be collected at a number of time points additional to those indicated for measuring 12 primary outcomes to: a) understand the dynamics of the trialled drug's action on the parasite 13 and b) permit adjustment for the potential diluting effects of reinfection post drug treatment, 14 an inevitable consequence for trials conducted in endemic settings with ongoing transmission. 15 Furthermore, dynamic models can have a role in not only evaluating new interventions, but 16 17 informing desired characteristics based on model-projected outcomes and benefits (i.e. the Target Product Profile) and helping to inform decisions on which candidates to take forward 18 19 from pre-clinical to clinical development.

# 20 Development of dynamic models: Research needs and challenges

There are research requirements for the further development of dynamic models (discussed in
[19, 37]) and for the accompanying and necessary economic evaluations. A summary of the
key research needs is given below.

#### 1 *Empirical and programmatic data*

2 A key impediment to the evaluation of NTD interventions, both in terms of economic and

3 public-health outcomes, is the lack of longitudinal data available from large-scale and long-

4 term control programmes [19] which are made available to modellers and analysts for model

5 parameterization and impact evaluation. This relative paucity of empirical studies is in itself

6 another major research gap for NTDS.

7 Robust economic evaluations can be fairly 'data hungry', ideally requiring programmatic data on pre-control endemicity from a variety of areas with different epidemiology and 8 9 programmatic attributes, including levels of therapeutic coverage and systematic noncompliance over time, and the duration and type of the intervention(s) employed. In addition 10 11 there needs to be an improved interaction and dialogue between modellers and those who 12 coordinate the data collection side of control programme. More epidemiological mapping is also necessary to understand: a) the distribution of infections (and co-infections) at different 13 14 programmatic levels (districts, regions etc.), potentially allowing a more geographically 15 targeted approach to control [38], and b) how seasonal variations in transmission may vary among areas, which may influence the benefit and optimum timing of different interventions. 16 17 Programmatic data will also be crucial to validate current and future model projections and, with individual data, to understand better the impact of individual heterogeneities in 18 responses to interventions (such as individual drug responses) on population-level benefits 19 20 and cost-effectiveness.

#### 21 Transmission breakpoints and surveillance strategies

Transmission breakpoints theoretically arise due to the operation of positive density
dependencies (Box 1) and refer to the density (e.g. number of macroparasites per host or
number of hosts infected with microparasites per unit population) below which the parasite

1 population cannot sustain itself and enters terminal decline [17]. Understanding the behaviour 2 of the host-parasite system in the vicinity of these transmission breakpoints is a priority area 3 of research [19], particularly on how different interventions may be associated with different 4 risks of infection resurgence after they are stopped or scaled down. Also, it will be important 5 to consider when interventions can be safely stopped based on the relationship between 6 transmission breakpoints and operational thresholds for intervention cessation (and starting 7 post-intervention surveillance). Economic modelling can also be helpful in informing the 8 most cost-effective operational threshold for intervention cessation and identifying the 9 methods for scaling down or stopping interventions, accounting for the sensitivity of the current diagnostic methods (and the potential economic value of new, more accurate ones). 10 Dynamic models can also be useful in investigating patterns of potential infection resurgence 11 following cessation of control and in designing cost-effective surveillance strategies to detect 12 13 this. This area will become increasing important as progress towards the 2020 NTD goals is

14 made.

15 *Health impacts* 

16 A notable gap in research is the development of dynamic models linking infection to

17 mortality and morbidity [19], a prerequisite for performing cost-unity analysis. More research

18 is needed to ascertain how disease burden relates to present, past, and cumulative experience

19 of infection (and co-infection) [19]. There is also a growing need for further

20 development/application of more comprehensive disability metrics such as the quality-

21 adjusted life year (QALY) which can more effectively capture the disease burden of NTD

22 infections [39]. Moreover, it is important that other benefits of NTD control strategies, such

as increased school attendance, cognitive development, increased productivity, and the

24 prevented economic burden [40, 41] are considered.

1 *Costs* 

2 Comprehensive costs of NTD interventions should be incorporated into the modelling 3 analyses, in particular the relative cost of alternative/complementary strategies, which may 4 change with scale and can be influenced by other related intervention activities. For example, 5 the relative increase in the programme cost of increasing the treatment frequency of MDA, or 6 using complementary focal vector control, will likely change depending on the scale that the 7 strategy is adopted within a given geographical area (e.g. across a whole district or just in 8 high risk areas). Additionally, further investigation into what drives pronounced variation in 9 the costs of interventions among countries [42, 43], and how this may influence the relative 10 costs of alternative strategies, is urgently needed. The potential increase in costs as 11 programmes are expanded to cover hard-to-reach groups also needs consideration [44]. 12 Moreover, multiple perspectives should be investigated, not just from the health care providers (i.e. the control programmes and non-governmental organization (NGO) donors), 13 but also that from the pharmaceutical companies that donate or discount drugs, as different 14 15 intervention strategies will influence the costs associated with these parties in different ways.

16 *Economic evaluation framework* 

17 Many alternative intervention strategies (such as increasing treatment frequency of MDA) 18 will be aimed at accelerating progress towards elimination, rather than delivering additional 19 health gains. Therefore, cost-effectiveness ratios alone, which compare interventions using cost per DALY averted (or an alternative measure of health gain, such as in terms of their 20 value for money in obtaining health gains) will not necessarily be the most informative metric 21 22 by which to judge the 'best' or most appropriate intervention strategy. This highlights a need for a new paradigm in economic evaluation frameworks which better appraise the long-term 23 24 benefits of elimination. For example, in economic evaluations it is standard practice to discount both the costs and benefits occurring in the future. However, it has been suggested 25

1 that applying a constant discount rate may undervalue the future, an important consideration 2 when evaluating elimination strategies [45]. These economic evaluation frameworks should 3 potentially consider the implications of using non-constant discount rates (and /or using 4 different rates for costs and effects) [45, 46]. It is also important that future economic 5 evaluations incorporate appropriate multivariate sensitivity analysis to investigate the 6 robustness of their results to changes in key assumptions, as well as exploring the 7 generalisability of their results outside of the particular programmatic and epidemiological 8 context. Univariate sensitivity analysis (i.e. only changing one parameter value at a time) may 9 not capture important correlations in the effects induced by different parameters [33].

#### 10 The implications of integrated control

NTD interventions are increasingly becoming integrated to target more than one disease or 11 groups of diseases at once, and there is a growing need for modelling studies evaluating 12 13 alternative interventions to account for this. For instance, onchocerciasis and lymphatic filariasis intervention activities (as well as those for schistosomiasis and soil-transmitted 14 15 helminthiases) are often carried out simultaneously, and therefore a change in strategy for one 16 (such as increasing the treatment frequency) will likely have an additional impact on the other 17 diseases in co-endemic areas. There is also an increasing drive to integrate chemotherapybased NTD interventions with complementary socioeconomic and nutritional improvements 18 such as WASH (water, sanitation, and hygiene) and school-feeding. However, the 19 implications of this integration on the optimum control and monitoring and evaluation 20 strategy (for the combination of targeted diseases) [47] and programme costs have not been 21 fully investigated. 22

## 23 Model validation and comparisons

24 More funding and resources need to be made available for model validation; before

25 undertaking detailed evaluations of competing intervention strategies, one must be able to

1	demonstrate that a model adequately describes epidemiological patterns both in endemic
2	parasite (pathogen) populations and, crucially, in populations perturbed by control activities.
3	This process is often held back by a lack of appropriate longitudinal data from sentinel
4	populations. Indeed, the gathering of such data is often not prioritized highly enough despite
5	being essential to robust policy-relevant model-based evaluations.
6	To inform policy in a reliable and robust manner, it will be essential that model comparison
7	exercises are performed [19], exploring the reasons for any disparity between the results and
8	conclusions of different models on the economic value of different control strategies.
9	By explicitly considering transmission, dynamic models have greater inherent uncertainty
10	than their static counterparts, in both parameterization and structure. This makes their
11	validation even more important. However, adequately reflecting uncertainty in our
12	understanding of infections and their transmission dynamics has an important role in model-
13	based decision making; policy makers should be presented with both the expectation and the
14	accompanying uncertainty associated with epidemiological and economic projections alike.
14 15	accompanying uncertainty associated with epidemiological and economic projections alike.
	accompanying uncertainty associated with epidemiological and economic projections alike.
15	
15 16	Concluding Remarks
15 16 17	<b>Concluding Remarks</b> The lack of economic evaluations using dynamic transmission models demonstrates an
15 16 17 18	<b>Concluding Remarks</b> The lack of economic evaluations using dynamic transmission models demonstrates an important research gap and methodological shortcoming in the evaluation of NTD control
15 16 17 18 19	<b>Concluding Remarks</b> The lack of economic evaluations using dynamic transmission models demonstrates an important research gap and methodological shortcoming in the evaluation of NTD control and elimination interventions. This is particularly important in the context of the London
15 16 17 18 19 20	<b>Concluding Remarks</b> The lack of economic evaluations using dynamic transmission models demonstrates an important research gap and methodological shortcoming in the evaluation of NTD control and elimination interventions. This is particularly important in the context of the London Declaration, where dynamic models will be nesseccary to investigate what changes to policy
15 16 17 18 19 20 21	<b>Concluding Remarks</b> The lack of economic evaluations using dynamic transmission models demonstrates an important research gap and methodological shortcoming in the evaluation of NTD control and elimination interventions. This is particularly important in the context of the London Declaration, where dynamic models will be nesseccary to investigate what changes to policy are required to meet the 2020 NTD control and elimination goals. Despite being easier to

1	We believe that reaching the 2020 NTD goals will not only depend on the continuing and
2	burgeoning implementation of existing intervention strategies, and the development of new
3	more effective intervention tools, but also on the concomitant development of judiciously
4	formulated and well parameterized dynamic transmission models with which to undertake
5	economic evaluations. Such evaluations will inform policy by determining the optimum way
6	in which intervention tools should be deployed in fundamentally resource-poor settings.
7	In order for this to happen effectively, a wide range of research needs should be addressed
8	including, but not limited to: i) more empirical studies and greater availability of
9	programmatic data to further parameterise/develop these dynamic transmission models, and
10	to improve estimated health effects, ii) the advancement of economic evaluation frameworks
11	and collection of detailed cost data that account for integrated NTD control programmes, and
12	iii) rigorous model validation and comparison.
13 14 15	Acknowledgements:
14	Acknowledgements: We thank Zulma Cucunubá Pérez and Pierre Nouvellet for their comments and feedback on the Chagas disease models.
14 15 16	We thank Zulma Cucunubá Pérez and Pierre Nouvellet for their comments and feedback on
14 15 16 17	We thank Zulma Cucunubá Pérez and Pierre Nouvellet for their comments and feedback on the Chagas disease models.
14 15 16 17 18 19 20 21	We thank Zulma Cucunubá Pérez and Pierre Nouvellet for their comments and feedback on the Chagas disease models. Abbreviations: Disability-adjusted life-years (DALYs); Disease Control Priorities Project (DCPP); Human immunodeficiency virus (HIV); Mass drug administration (MDA); Neglected tropical diseases (NTDs); Non-governmental organization (NGO); Quality-adjusted life year
14 15 16 17 18 19 20 21 22	We thank Zulma Cucunubá Pérez and Pierre Nouvellet for their comments and feedback on the Chagas disease models. Abbreviations: Disability-adjusted life-years (DALYs); Disease Control Priorities Project (DCPP); Human immunodeficiency virus (HIV); Mass drug administration (MDA); Neglected tropical diseases (NTDs); Non-governmental organization (NGO); Quality-adjusted life year
14 15 16 17 18 19 20 21 22 23	We thank Zulma Cucunubá Pérez and Pierre Nouvellet for their comments and feedback on the Chagas disease models. Abbreviations: Disability-adjusted life-years (DALYs); Disease Control Priorities Project (DCPP); Human immunodeficiency virus (HIV); Mass drug administration (MDA); Neglected tropical diseases (NTDs); Non-governmental organization (NGO); Quality-adjusted life year
14 15 16 17 18 19 20 21 22 23 23 24	We thank Zulma Cucunubá Pérez and Pierre Nouvellet for their comments and feedback on the Chagas disease models. Abbreviations: Disability-adjusted life-years (DALYs); Disease Control Priorities Project (DCPP); Human immunodeficiency virus (HIV); Mass drug administration (MDA); Neglected tropical diseases (NTDs); Non-governmental organization (NGO); Quality-adjusted life year
14 15 16 17 18 19 20 21 22 23 24 25	We thank Zulma Cucunubá Pérez and Pierre Nouvellet for their comments and feedback on the Chagas disease models. Abbreviations: Disability-adjusted life-years (DALYs); Disease Control Priorities Project (DCPP); Human immunodeficiency virus (HIV); Mass drug administration (MDA); Neglected tropical diseases (NTDs); Non-governmental organization (NGO); Quality-adjusted life year

29

# **References**

2 3	1.	Hotez, P.J., et al., <i>Control of neglected tropical diseases</i> . N Engl J Med, 2007. <b>357</b> (10), 1018-27.
	2	
4 5	2.	Hotez, P., et al., <i>The Lancet's chronic diseases series</i> . Lancet, 2006. <b>367</b> (9510), 563-4; author reply 564-5.
6 7	3.	Hotez, P., et al., <i>The Global Burden of Disease Study 2010: implications for the neglected tropical diseases.</i> PLoS Negl Trop Dis, 2014. 8(7), e2865.
	4	
8 9	4.	Hotez, P.J., et al., <i>Incorporating a rapid-impact package for neglected tropical diseases with programs for HIV/AIDS, tuberculosis, and malaria.</i> PLoS Med, 2006. <b>3</b> (5), e102.
10	5.	London Declaration on Neglected Tropical Diseases. <i>Ending the neglect and reaching 2020</i>
11		goals. 2013; Available from:
12		http://unitingtocombatntds.org/downloads/press/ntd_event_london_declaration_on_ntds.pdf
13	6.	World Health Organization. Accelerating work to overcome the global impact of neglected
14	0.	tropical diseases– A roadmap for implementation. 2012; Available from:
15		http://www.who.int/neglected_diseases/NTD_RoadMap_2012_Fullversion.pdf
16	7.	Laxminarayan, R., et al., Advancement of global health: key messages from the Disease
17	7.	Control Priorities Project. Lancet, 2006. <b>367</b> (9517), 1193-208.
18	8.	Jamison, D.T., et al., <i>Disease control priorities in developing countries</i> . 2nd edn. Oxford
	0.	
19 20		University Press, 2006; Available from: <u>http://elibrary.worldbank.org/doi/abs/10.1596/978-0-</u> 8213-6179-5
21	9.	Weinstein, M.C., et al., Principles of good practice for decision analytic modeling in health-
22		care evaluation: Report of the ISPOR task force on good research practices—modeling
23		<i>studies</i> . Value Health, 2003. <b>6</b> (1), 9-17.
24	10.	Lugner, A.K., et al., Dynamic versus static models in cost-effectiveness analyses of anti-viral
25	101	<i>drug therapy to mitigate an influenza pandemic</i> . Health Econ, 2010. <b>19</b> (5), 518-31.
26	11.	Sonnenberg, F.A. and Beck, J.R., <i>Markov models in medical decision making: a practical</i>
27	11.	guide. Med Decis Making, 1993. <b>13</b> (4), 322-38.
28	12.	Briggs, A. and Sculpher, M., An introduction to Markov modelling for economic evaluation.
29	12.	Pharmacoeconomics, 1998. <b>13</b> (4), 397-409.
30	13.	Bala, M.V. and Mauskopf, J.A., <i>Optimal assignment of treatments to health states using a</i>
	15.	
31		<i>Markov decision model: an introduction to basic concepts.</i> Pharmacoeconomics, 2006. <b>24</b> (4),
32	14	345-54. Kim S.V. and Caldia S.L. Cast effectiveness making a function for a first second s
33	14.	Kim, S.Y. and Goldie, S.J., Cost-effectiveness analyses of vaccination programmes : a
34	1.7	focused review of modelling approaches. Pharmacoeconomics, 2008. <b>26</b> (3), 191-215.
35	15.	Edmunds, W.J., et al., <i>Evaluating the cost-effectiveness of vaccination programmes: a</i>
36		dynamic perspective. Stat Med, 1999. 18(23), 3263-82.
37	16.	Koopman, J., Modeling infection transmission. Annu Rev Public Health, 2004. 25, 303-26.
38	17.	Anderson, R.M. and May, R.M. Infectious Diseases of Humans: Dynamics and Control 1992,
39		Oxford Science Publications.
40	18.	Brennan, A., et al., A taxonomy of model structures for economic evaluation of health
41		technologies. Health Econ, 2006. 15(12), 1295-310.
42	19.	Basáñez, M.G., et al., A research agenda for helminth diseases of humans: modelling for
43		control and elimination. PLoS Negl Trop Dis, 2012. 6(4), e1548.
44	20.	Basáñez, M.G., et al., Effect of single-dose ivermectin on Onchocerca volvulus: a systematic
45		review and meta-analysis. Lancet Infect Dis, 2008. 8(5), 310-322.
46	21.	Hoerauf, A., et al., Depletion of Wolbachia endobacteria in Onchocerca volvulus by
47		doxycycline and microfilaridermia after ivermectin treatment. Lancet, 2001. 357(9266),
48		1415-1416.
49	22.	Hoerauf, A., et al., Doxycycline in the treatment of human onchocerciasis: Kinetics of
50		Wolbachia endobacteria reduction and of inhibition of embryogenesis in female Onchocerca
51		<i>worms</i> . Microbes Infect, 2003. <b>5</b> (4), 261-73.

1 23. Hoerauf, A., et al., Wolbachia endobacteria depletion by doxycycline as antifilarial therapy 2 has macrofilaricidal activity in onchocerciasis: a randomized placebo-controlled study. Med 3 Microbiol Immunol, 2008. 197(3), 295-311. 4 24. Lustigman, S., et al., Ch 22 Vaccines linked to chemotherapy: A new approach to control 5 helminth infections, in Parasitic Helminths: Targets, Screens, Drugs and Vaccines (editor: 6 Caffrey, C. R.). 2012, Wiley-VCH Verlag. 7 25. Dowdle, W.R., Hopkins, D.R. eds, The eradication of infectious diseases: report of the 8 Dahlem Workshop on the Eradication of Infections Diseases. 1998, John Wiley & Sons, 9 Chichester. 10 26. Geoffard, P.Y. and Philipson, T., Disease eradication: private versus public vaccination. Am 11 Econ Rev, 1997. 87(1), 222-230. 12 27. Churcher, T.S., et al., Density dependence and the control of helminth parasites. J Anim Ecol 2006. 75(6), 1313-1320. 13 14 28. Basáñez, M.G., et al., Onchocerca-Simulium interactions and the population and 15 evolutionary biology of Onchocerca volvulus. Adv Parasitol, 2009. 68, 263-313. 16 29. Dietz, K., Density-dependence in parasite transmission dynamics. Parasitol Today, 1988. 17 **4**(4), 91-7. Churcher, T.S., et al., Density dependence and overdispersion in the transmission of helminth 18 30. 19 parasites. Parasitology, 2005. 131(1), 121-132. 20 31. Chidambaram, J.D., et al., Mass antibiotics for trachoma and the Allee effect. Lancet Infect 21 Dis. 2005. 5(4), 194-6. 22 32. Lietman, T.M., et al., The epidemiological dynamics of infectious trachoma may facilitate 23 elimination. Epidemics, 2011. 3(2), 119-24. Walker, D. and Fox-Rushby, J., Economic evaluation of parasitic diseases: a critique of the 24 33. 25 internal and external validity of published studies. Trop Med Int Health, 2000. 5(4), 237-49. 26 34. Anderson, R.M., et al., How effective is school-based deworming for the community-wide 27 control of soil-transmitted helminths? PLoS Negl Trop Dis, 2013. 7(2), e2027. 28 Truscott, J., et al., Can chemotherapy alone eliminate the transmission of soil transmitted 35. 29 helminths? Parasit Vectors, 2014. 7(1), 266. 30 36. Anderson, R.M., et al., The coverage and frequency of mass drug administration required to 31 eliminate persistent transmission of soil-transmitted helminths. Philos Trans R Soc Lond B Biol Sci, 2014. 369(1645), 20130435. 32 33 37. Kealey, A. and Smith, R., Neglected tropical diseases: infection, modeling, and control. J 34 Health Care Poor Underserved, 2010. 21(1), 53-69. 35 38. Sturrock, H.J., et al., Optimal survey designs for targeting chemotherapy against soiltransmitted helminths: effect of spatial heterogeneity and cost-efficiency of sampling. Am J 36 37 Trop Med Hyg, 2010. 82(6), 1079-87. 38 39. King, C.H., Health metrics for helminth infections. Acta Trop, 2013. 39 40. Lee, B.Y., et al., Global economic burden of Chagas disease: a computational simulation 40 model. Lancet Infect Dis, 2013. 13(4), 342-8. 41 41. Conteh, L., et al., Socioeconomic aspects of neglected tropical diseases. Lancet, 2010. 42 **375**(9710), 239-47. 43 42. Goldman, A.S., et al., National mass drug administration costs for lymphatic filariasis 44 elimination. PLoS Negl Trop Dis, 2007. 1(1), e67. 45 43. Keating, J., et al., Lymphatic filariasis and onchocerciasis prevention, treatment, and control 46 costs across diverse settings: A systematic review. Acta Tropica, 2014. 135, 86-95. 47 44. Johns, B. and Torres, T.T., Costs of scaling up health interventions: a systematic review. 48 Health Policy Plan, 2005. 20(1), 1-13. 49 45. Walker, D.G. and Lupp, J., Guide for preparing an eradication investment case. 2011; 50 Available from: http://eic-guidelines.org/ 51 46. Tediosi, F., et al., Developing eradication investment cases for onchocerciasis, lymphatic 52 filariasis, and human African trypanosomiasis: rationale and main challenges. PLoS Negl Trop Dis, 2013. 7(11), e2446. 53

<ul> <li>for urogenital schistosomiasis and active trachoma in Burkina Faso before chemotherapy using sentinel sites. BMC Infect Dis, 2011. 11, 191.</li> <li>48. Murray, C.J.L. and Lopez, A.D., Quantifying disability: data, methods and World Health Organ, 1994. 72(3), 481-94.</li> <li>49. May, R.M., Togetherness among schistosomes: its effects on the dynamics of Math Biosci, 1977. 35(3–4), 301-343.</li> <li>50. Lechat, M.F., et al., Selection of MDT strategies through epidemiometric mod Other Mycobact Dis, 1990. 58(2), 296-301.</li> <li>51. Lechat, M.F., et al., Application of an economic model to the study of lepros Int J Lepr Other Mycobact Dis, 1978. 46(1), 14-24.</li> <li>52. Chu, B.K., et al., The economic benefits resulting from the first 8 years of the Programme to Eliminate Lymphatic Filariasis (2000-2007). PLoS Negl Trop</li> </ul>	results. Bull of the infection. odeling. Int J Lepr
<ul> <li><i>chemotherapy using sentinel sites.</i> BMC Infect Dis, 2011. 11, 191.</li> <li>Murray, C.J.L. and Lopez, A.D., <i>Quantifying disability: data, methods and</i> World Health Organ, 1994. 72(3), 481-94.</li> <li>May, R.M., <i>Togetherness among schistosomes: its effects on the dynamics of</i> Math Biosci, 1977. 35(3–4), 301-343.</li> <li>Lechat, M.F., et al., <i>Selection of MDT strategies through epidemiometric me</i> Other Mycobact Dis, 1990. 58(2), 296-301.</li> <li>Lechat, M.F., et al., <i>Application of an economic model to the study of lepros</i> Int J Lepr Other Mycobact Dis, 1978. 46(1), 14-24.</li> <li>Chu, B.K., et al., <i>The economic benefits resulting from the first 8 years of th Programme to Eliminate Lymphatic Filariasis (2000-2007).</i> PLoS Negl Trophotometric Matheratics and Matheratic Science S</li></ul>	results. Bull of the infection. odeling. Int J Lepr
<ol> <li>4 48. Murray, C.J.L. and Lopez, A.D., Quantifying disability: data, methods and World Health Organ, 1994. 72(3), 481-94.</li> <li>6 49. May, R.M., Togetherness among schistosomes: its effects on the dynamics of Math Biosci, 1977. 35(3–4), 301-343.</li> <li>8 50. Lechat, M.F., et al., Selection of MDT strategies through epidemiometric me Other Mycobact Dis, 1990. 58(2), 296-301.</li> <li>10 51. Lechat, M.F., et al., Application of an economic model to the study of lepros Int J Lepr Other Mycobact Dis, 1978. 46(1), 14-24.</li> <li>12 52. Chu, B.K., et al., The economic benefits resulting from the first 8 years of the Programme to Eliminate Lymphatic Filariasis (2000-2007). PLoS Negl Trop</li> </ol>	of the infection. odeling. Int J Lepr
<ul> <li>World Health Organ, 1994. 72(3), 481-94.</li> <li>May, R.M., <i>Togetherness among schistosomes: its effects on the dynamics of</i> Math Biosci, 1977. 35(3–4), 301-343.</li> <li>Lechat, M.F., et al., <i>Selection of MDT strategies through epidemiometric ma</i> Other Mycobact Dis, 1990. 58(2), 296-301.</li> <li>Lechat, M.F., et al., <i>Application of an economic model to the study of lepros</i> Int J Lepr Other Mycobact Dis, 1978. 46(1), 14-24.</li> <li>Chu, B.K., et al., <i>The economic benefits resulting from the first 8 years of th</i> <i>Programme to Eliminate Lymphatic Filariasis (2000-2007)</i>. PLoS Negl Tropped 2007.</li> </ul>	of the infection. odeling. Int J Lepr
<ol> <li>May, R.M., Togetherness among schistosomes: its effects on the dynamics of Math Biosci, 1977. 35(3–4), 301-343.</li> <li>Lechat, M.F., et al., Selection of MDT strategies through epidemiometric ma Other Mycobact Dis, 1990. 58(2), 296-301.</li> <li>Lechat, M.F., et al., Application of an economic model to the study of lepros Int J Lepr Other Mycobact Dis, 1978. 46(1), 14-24.</li> <li>Chu, B.K., et al., The economic benefits resulting from the first 8 years of th Programme to Eliminate Lymphatic Filariasis (2000-2007). PLoS Negl Trop</li> </ol>	odeling. Int J Lepr
<ul> <li>Math Biosci, 1977. 35(3-4), 301-343.</li> <li>50. Lechat, M.F., et al., Selection of MDT strategies through epidemiometric may Other Mycobact Dis, 1990. 58(2), 296-301.</li> <li>51. Lechat, M.F., et al., Application of an economic model to the study of leprose Int J Lepr Other Mycobact Dis, 1978. 46(1), 14-24.</li> <li>52. Chu, B.K., et al., The economic benefits resulting from the first 8 years of the Programme to Eliminate Lymphatic Filariasis (2000-2007). PLoS Negl Tropology 10, 1978.</li> </ul>	odeling. Int J Lepr
<ul> <li>9 Other Mycobact Dis, 1990. 58(2), 296-301.</li> <li>10 51. Lechat, M.F., et al., <i>Application of an economic model to the study of lepros</i>.</li> <li>11 Int J Lepr Other Mycobact Dis, 1978. 46(1), 14-24.</li> <li>12 52. Chu, B.K., et al., <i>The economic benefits resulting from the first 8 years of the</i>.</li> <li>13 <i>Programme to Eliminate Lymphatic Filariasis (2000-2007)</i>. PLoS Negl Tropped 10.</li> </ul>	
<ol> <li>Lechat, M.F., et al., Application of an economic model to the study of leprose</li> <li>Int J Lepr Other Mycobact Dis, 1978. 46(1), 14-24.</li> <li>Chu, B.K., et al., The economic benefits resulting from the first 8 years of the Programme to Eliminate Lymphatic Filariasis (2000-2007). PLoS Negl Trop</li> </ol>	sy control costs.
<ol> <li>Int J Lepr Other Mycobact Dis, 1978. 46(1), 14-24.</li> <li>Chu, B.K., et al., <i>The economic benefits resulting from the first 8 years of th</i> <i>Programme to Eliminate Lymphatic Filariasis (2000-2007)</i>. PLoS Negl Tropped 100.</li> </ol>	sy control costs.
<ol> <li>S2. Chu, B.K., et al., <i>The economic benefits resulting from the first 8 years of the</i></li> <li><i>Programme to Eliminate Lymphatic Filariasis (2000-2007)</i>. PLoS Negl Tropped 1</li> </ol>	
13 Programme to Eliminate Lymphatic Filariasis (2000-2007). PLoS Negl Tro	
	e Global
	p Dis, 2010. <b>4</b> (6),
14 e708.	•
15 53. Michael, E., et al., Global eradication of lymphatic filariasis: The value of c	chronic disease
16 <i>control in parasite elimination programmes.</i> PLoS One, 2008. <b>3</b> (8), e2936.	
17 54. Stolk, W.A., et al., Modeling the impact and costs of semiannual mass drug	administration
18 for accelerated elimination of Lymphatic Filariasis. PLoS Negl Trop Dis, 2	
19 55. Baltussen, R. and Smith, A., Cost effectiveness of strategies to combat visio	n and hearing loss
20 in sub-Saharan Africa and South East Asia: mathematical modelling study.	-
21 p. e615.	· · · · · · · · · · · · · · · · · · ·
22 56. Baltussen, R., et al., Cost-effectiveness of trachoma control in seven world r	regions.
23 Ophthalmic Epidemiol 2005. <b>12</b> (2), 91-101.	0
24 57. Frick, K.D., et al., Modeling the economic net benefit of a potential vaccina	tion program
25 against ocular infection with Chlamydia trachomatis. Vaccine, 2004. 22(5-0	
26 58. Blake, I.M., et al., <i>Targeting antibiotics to households for trachoma control</i>	
27 Dis, 2010. <b>4</b> (11), e862.	<i>b b b b b b b b b b</i>
28 59. Fèvre, E.M., et al., <i>Estimating the burden of rhodesiense sleeping sickness a</i>	luring an outbreak
<i>in Serere, eastern Uganda</i> . BMC Public Health, 2008. <b>8</b> .	
30 60. WHO, Control and surveillance of African trypanosomiasis. Report of a WI	HO Expert
31 <i>Committee</i> . World Health Organ Tech Rep Ser, 1998. <b>881</b> , I-VI, 1-114.	
32 61. Lutumba, P., et al., The efficiency of different detection strategies of human	African
<ul> <li>trypanosomiasis by <i>T. b. gambiense</i>. Trop Med Int Health, 2005. <b>10</b>(4), 347</li> </ul>	
34 62. Politi, C., et al., <i>Cost-effectiveness analysis of alternative treatments of Afric</i>	
<ul> <li><i>trypanosomiasis in Uganda</i>. Health Econ, 1995. 4(4), 273-87.</li> </ul>	Suit Suite terrise
	sease control
36 63. Shaw, A.P., Comparative analysis of the costs and benefits of alternative di	
<ul> <li>36 63. Shaw, A.P., Comparative analysis of the costs and benefits of alternative diastrategies: vector control versus human case finding and treatment. Ann So</li> </ul>	
<ul> <li>Shaw, A.P., Comparative analysis of the costs and benefits of alternative di</li> <li>strategies: vector control versus human case finding and treatment. Ann So</li> <li>1989. 69(Suppl 1), 237-53.</li> </ul>	c Belg Med Trop,
<ul> <li>36 63. Shaw, A.P., Comparative analysis of the costs and benefits of alternative dia strategies: vector control versus human case finding and treatment. Ann So 1989. 69(Suppl 1), 237-53.</li> <li>39 64. Korte, R., et al., Cost and effectiveness of different approaches to schistoson</li> </ul>	c Belg Med Trop,
<ul> <li>Shaw, A.P., Comparative analysis of the costs and benefits of alternative dia strategies: vector control versus human case finding and treatment. Ann So 1989. 69(Suppl 1), 237-53.</li> <li>Korte, R., et al., Cost and effectiveness of different approaches to schistoson Africa. Trop Med Parasitol, 1986. 37(2), 149-52.</li> </ul>	c Belg Med Trop, niasis control in
<ul> <li>36 63. Shaw, A.P., Comparative analysis of the costs and benefits of alternative dia strategies: vector control versus human case finding and treatment. Ann So 1989. 69(Suppl 1), 237-53.</li> <li>39 64. Korte, R., et al., Cost and effectiveness of different approaches to schistoson Africa. Trop Med Parasitol, 1986. 37(2), 149-52.</li> <li>41 65. Kirigia, J.M., et al., A cost-benefit analysis of preventive schistosomiasis int</li> </ul>	c Belg Med Trop, niasis control in
<ul> <li>Shaw, A.P., Comparative analysis of the costs and benefits of alternative dia strategies: vector control versus human case finding and treatment. Ann So 1989. 69(Suppl 1), 237-53.</li> <li>Korte, R., et al., Cost and effectiveness of different approaches to schistoson Africa. Trop Med Parasitol, 1986. 37(2), 149-52.</li> <li>Kirigia, J.M., et al., A cost-benefit analysis of preventive schistosomiasis int Kenya. Afr J Health Sci, 2000. 7(3-4), 5-11.</li> </ul>	c Belg Med Trop, niasis control in rerventions in
<ol> <li>Shaw, A.P., Comparative analysis of the costs and benefits of alternative dia strategies: vector control versus human case finding and treatment. Ann So 1989. 69(Suppl 1), 237-53.</li> <li>Korte, R., et al., Cost and effectiveness of different approaches to schistoson Africa. Trop Med Parasitol, 1986. 37(2), 149-52.</li> <li>Kirigia, J.M., et al., A cost-benefit analysis of preventive schistosomiasis int Kenya. Afr J Health Sci, 2000. 7(3-4), 5-11.</li> <li>Guyatt, H.L. and Evans, D., Desirable characteristics of a schistosomiasis version.</li> </ol>	nc Belg Med Trop, niasis control in rerventions in vaccine: some
<ol> <li>Shaw, A.P., Comparative analysis of the costs and benefits of alternative di strategies: vector control versus human case finding and treatment. Ann So 1989. 69(Suppl 1), 237-53.</li> <li>Korte, R., et al., Cost and effectiveness of different approaches to schistoson Africa. Trop Med Parasitol, 1986. 37(2), 149-52.</li> <li>Kirigia, J.M., et al., A cost-benefit analysis of preventive schistosomiasis int Kenya. Afr J Health Sci, 2000. 7(3-4), 5-11.</li> <li>Guyatt, H.L. and Evans, D., Desirable characteristics of a schistosomiasis v implications of a cost-effectiveness analysis. Acta Trop, 1995. 59(3), 197-20.</li> </ol>	c Belg Med Trop, niasis control in erventions in vaccine: some 09.
<ol> <li>Shaw, A.P., Comparative analysis of the costs and benefits of alternative dia strategies: vector control versus human case finding and treatment. Ann So 1989. 69(Suppl 1), 237-53.</li> <li>Korte, R., et al., Cost and effectiveness of different approaches to schistoson Africa. Trop Med Parasitol, 1986. 37(2), 149-52.</li> <li>Kirigia, J.M., et al., A cost-benefit analysis of preventive schistosomiasis int Kenya. Afr J Health Sci, 2000. 7(3-4), 5-11.</li> <li>Guyatt, H.L. and Evans, D., Desirable characteristics of a schistosomiasis v implications of a cost-effectiveness analysis. Acta Trop, 1995. 59(3), 197-20.</li> <li>Prescott, N.M., The economics of schistosomiasis chemotherapy. Parasitol T</li> </ol>	c Belg Med Trop, niasis control in erventions in vaccine: some 09.
<ol> <li>Shaw, A.P., Comparative analysis of the costs and benefits of alternative dia strategies: vector control versus human case finding and treatment. Ann So 1989. 69(Suppl 1), 237-53.</li> <li>Korte, R., et al., Cost and effectiveness of different approaches to schistoson Africa. Trop Med Parasitol, 1986. 37(2), 149-52.</li> <li>Kirigia, J.M., et al., A cost-benefit analysis of preventive schistosomiasis int Kenya. Afr J Health Sci, 2000. 7(3-4), 5-11.</li> <li>Guyatt, H.L. and Evans, D., Desirable characteristics of a schistosomiasis v implications of a cost-effectiveness analysis. Acta Trop, 1995. 59(3), 197-20.</li> <li>Prescott, N.M., The economics of schistosomiasis chemotherapy. Parasitol 7 21-4.</li> </ol>	nc Belg Med Trop, niasis control in rerventions in vaccine: some 09. Foday, 1987. <b>3</b> (1),
<ol> <li>Shaw, A.P., Comparative analysis of the costs and benefits of alternative dia strategies: vector control versus human case finding and treatment. Ann So 1989. 69(Suppl 1), 237-53.</li> <li>Korte, R., et al., Cost and effectiveness of different approaches to schistoson Africa. Trop Med Parasitol, 1986. 37(2), 149-52.</li> <li>Kirigia, J.M., et al., A cost-benefit analysis of preventive schistosomiasis int Kenya. Afr J Health Sci, 2000. 7(3-4), 5-11.</li> <li>Guyatt, H.L. and Evans, D., Desirable characteristics of a schistosomiasis v implications of a cost-effectiveness analysis. Acta Trop, 1995. 59(3), 197-20.</li> <li>Prescott, N.M., The economics of schistosomiasis chemotherapy. Parasitol 7 21-4.</li> <li>Evans, D.B. and Guyatt, H.L.,Human behaviour, cost-effectiveness analysis</li> </ol>	nc Belg Med Trop, niasis control in rerventions in vaccine: some 09. Foday, 1987. <b>3</b> (1), and research and
<ol> <li>Shaw, A.P., Comparative analysis of the costs and benefits of alternative dia strategies: vector control versus human case finding and treatment. Ann So 1989. 69(Suppl 1), 237-53.</li> <li>Korte, R., et al., Cost and effectiveness of different approaches to schistoson Africa. Trop Med Parasitol, 1986. 37(2), 149-52.</li> <li>Kirigia, J.M., et al., A cost-benefit analysis of preventive schistosomiasis int Kenya. Afr J Health Sci, 2000. 7(3-4), 5-11.</li> <li>Guyatt, H.L. and Evans, D., Desirable characteristics of a schistosomiasis v implications of a cost-effectiveness analysis. Acta Trop, 1995. 59(3), 197-20.</li> <li>Prescott, N.M., The economics of schistosomiasis chemotherapy. Parasitol T 21-4.</li> <li>Evans, D.B. and Guyatt, H.L.,Human behaviour, cost-effectiveness analysis development priorities: the case of a schistosomiasis vaccine. Trop Med Int</li> </ol>	nc Belg Med Trop, niasis control in rerventions in vaccine: some 09. Foday, 1987. <b>3</b> (1), and research and
<ol> <li>Shaw, A.P., Comparative analysis of the costs and benefits of alternative dia strategies: vector control versus human case finding and treatment. Ann So 1989. <b>69</b>(Suppl 1), 237-53.</li> <li>Korte, R., et al., Cost and effectiveness of different approaches to schistoson Africa. Trop Med Parasitol, 1986. <b>37</b>(2), 149-52.</li> <li>Kirigia, J.M., et al., A cost-benefit analysis of preventive schistosomiasis int Kenya. Afr J Health Sci, 2000. <b>7</b>(3-4), 5-11.</li> <li>Guyatt, H.L. and Evans, D., Desirable characteristics of a schistosomiasis v implications of a cost-effectiveness analysis. Acta Trop, 1995. <b>59</b>(3), 197-20.</li> <li>Prescott, N.M., The economics of schistosomiasis chemotherapy. Parasitol T 21-4.</li> <li>Evans, D.B. and Guyatt, H.L.,Human behaviour, cost-effectiveness analysis development priorities: the case of a schistosomiasis vaccine. Trop Med Int <b>2</b>(11), A47-A54.</li> </ol>	niasis control in niasis control in erventions in vaccine: some 09. Foday, 1987. <b>3</b> (1), and research and Health, 1997.
<ol> <li>Shaw, A.P., Comparative analysis of the costs and benefits of alternative dia strategies: vector control versus human case finding and treatment. Ann So 1989. 69(Suppl 1), 237-53.</li> <li>Korte, R., et al., Cost and effectiveness of different approaches to schistoson Africa. Trop Med Parasitol, 1986. 37(2), 149-52.</li> <li>Kirigia, J.M., et al., A cost-benefit analysis of preventive schistosomiasis int Kenya. Afr J Health Sci, 2000. 7(3-4), 5-11.</li> <li>Guyatt, H.L. and Evans, D., Desirable characteristics of a schistosomiasis v implications of a cost-effectiveness analysis. Acta Trop, 1995. 59(3), 197-20.</li> <li>Prescott, N.M., The economics of schistosomiasis chemotherapy. Parasitol T 21-4.</li> <li>Evans, D.B. and Guyatt, H.L.,Human behaviour, cost-effectiveness analysis development priorities: the case of a schistosomiasis vaccine. Trop Med Int 2(11), A47-A54.</li> <li>King, C.H., et al., Utility of Repeated Praziquantel Dosing in the Treatment</li> </ol>	nc Belg Med Trop, niasis control in rerventions in vaccine: some 09. Today, 1987. <b>3</b> (1), and research and Health, 1997.
<ol> <li>Shaw, A.P., Comparative analysis of the costs and benefits of alternative dia strategies: vector control versus human case finding and treatment. Ann So 1989. 69(Suppl 1), 237-53.</li> <li>Korte, R., et al., Cost and effectiveness of different approaches to schistoson Africa. Trop Med Parasitol, 1986. 37(2), 149-52.</li> <li>Kirigia, J.M., et al., A cost-benefit analysis of preventive schistosomiasis int Kenya. Afr J Health Sci, 2000. 7(3-4), 5-11.</li> <li>Guyatt, H.L. and Evans, D., Desirable characteristics of a schistosomiasis v implications of a cost-effectiveness analysis. Acta Trop, 1995. 59(3), 197-20.</li> <li>Prescott, N.M., The economics of schistosomiasis chemotherapy. Parasitol 7 21-4.</li> <li>Evans, D.B. and Guyatt, H.L.,Human behaviour, cost-effectiveness analysis development priorities: the case of a schistosomiasis vaccine. Trop Med Int 2(11), A47-A54.</li> <li>King, C.H., et al., Utility of Repeated Praziquantel Dosing in the Treatment Schistosomiasis in High-Risk Communities in Africa: A Systematic Review.</li> </ol>	nc Belg Med Trop, niasis control in rerventions in vaccine: some 09. Today, 1987. <b>3</b> (1), and research and Health, 1997.
<ol> <li>Shaw, A.P., Comparative analysis of the costs and benefits of alternative di strategies: vector control versus human case finding and treatment. Ann So 1989. <b>69</b>(Suppl 1), 237-53.</li> <li>Korte, R., et al., Cost and effectiveness of different approaches to schistoson Africa. Trop Med Parasitol, 1986. <b>37</b>(2), 149-52.</li> <li>Kirigia, J.M., et al., A cost-benefit analysis of preventive schistosomiasis int Kenya. Afr J Health Sci, 2000. <b>7</b>(3-4), 5-11.</li> <li>Guyatt, H.L. and Evans, D., Desirable characteristics of a schistosomiasis v implications of a cost-effectiveness analysis. Acta Trop, 1995. <b>59</b>(3), 197-20.</li> <li>Prescott, N.M., The economics of schistosomiasis chemotherapy. Parasitol T 21-4.</li> <li>Evans, D.B. and Guyatt, H.L.,Human behaviour, cost-effectiveness analysis development priorities: the case of a schistosomiasis vaccine. Trop Med Int <b>2</b>(11), A47-A54.</li> <li>King, C.H., et al., Utility of Repeated Praziquantel Dosing in the Treatment Schistosomiasis in High-Risk Communities in Africa: A Systematic Review. Dis, 2011. <b>5</b>(9), 1321</li> </ol>	nc Belg Med Trop, niasis control in rerventions in vaccine: some 09. Foday, 1987. <b>3</b> (1), and research and Health, 1997.
<ol> <li>Shaw, A.P., Comparative analysis of the costs and benefits of alternative dia strategies: vector control versus human case finding and treatment. Ann So 1989. 69(Suppl 1), 237-53.</li> <li>Korte, R., et al., Cost and effectiveness of different approaches to schistoson Africa. Trop Med Parasitol, 1986. 37(2), 149-52.</li> <li>Kirigia, J.M., et al., A cost-benefit analysis of preventive schistosomiasis int Kenya. Afr J Health Sci, 2000. 7(3-4), 5-11.</li> <li>Guyatt, H.L. and Evans, D., Desirable characteristics of a schistosomiasis v implications of a cost-effectiveness analysis. Acta Trop, 1995. 59(3), 197-20.</li> <li>Prescott, N.M., The economics of schistosomiasis chemotherapy. Parasitol 7 21-4.</li> <li>Evans, D.B. and Guyatt, H.L.,Human behaviour, cost-effectiveness analysis development priorities: the case of a schistosomiasis vaccine. Trop Med Int 2(11), A47-A54.</li> <li>King, C.H., et al., Utility of Repeated Praziquantel Dosing in the Treatment Schistosomiasis in High-Risk Communities in Africa: A Systematic Review.</li> </ol>	nc Belg Med Trop, niasis control in erventions in vaccine: some 09. Today, 1987. <b>3</b> (1), e and research and Health, 1997. of Plos Negl Trop ntion for reducing

<ol> <li>model. AmJ 1109 Med Phyg. 1977. 20(3): 502-16.</li> <li>Gyatt, H.L. and Chan, M.S., An investigation into the interaction between drug efficacy and drug price of praciquantel in determining the cost-effectiveness of school-targeted treatment for Schoistosma mamsoni using a population dynamic model. Trop Med Int Health, 1998. 30(6), 425-35.</li> <li>Carabin, H., et al., A population dynamic approach to evaluating the impact of school attendance on the unit cost and effectiveness of school-based schistosomiasis chemotherapy programmes. Parasitology, 2000. 121 (P 12), 171-83.</li> <li>Ndeffo Mbah, M.L., et al., <i>Projecting the Long-Term Impact of School- or Community-Based Mass-Treatment Interventions for Control of Schistosom Infection.</i> Plos Negl Trop Dis, 2013. 7(8).</li> <li>Wang, X.X., et al., <i>Projecting the Long-Term Impact of School- or Community-Based Mass-Treatment Interventions for Control of Schistosoma Infection.</i> Plos Negl Trop Dis, 2012. 6(11).</li> <li>Guyatt, H.L., <i>Different approaches to modelling the cost-effectiveness of schistosomiasis control.</i> Men Inst Oswaldo Cruz, 1998. 93 (Suppl 1), 75-84.</li> <li>Turner, H.C., et al., <i>Reaching the London Declaration on Neglected Tropical Diseases goals for onchocerciasis: an economic evaluation of increasing the frequency of ivermectin treatment in Africa.</i> Clin Infect Dis, 2014. 59(7): 223-32.</li> <li>Coffeng, L.E., et al., <i>African Trogramme for Onchocerciasis Control 1995–2015: Model-estimatel health impact and cosn.</i> FLoS Negl Trop Dis, 2013. 7(1), e2032.</li> <li>Lee, B.Y., et al., <i>The potential economic value of a hookworm vaccine.</i> Vaccine, 2011. 29(6), 1201-10.</li> <li>Guyatt, H.L., et al., <i>Control of Ascaris infection by chemotherapy: which is the most cost-effective option?</i> Trans R Soc Trop Med Hyg, 1995. 89(1), 16-20.</li> <li>Guyatt, H.L., et al., <i>Control of Ascaris infection by chemotherapy: which is the most cost-effective option?</i> Trans R Soc Trop Med Hyg, 1995. 89(1), 16-20.<th>1</th><th>71.</th><th>Rosenfield, P.L., et al., <i>Development and verification of a schistosomiasis transmission</i></th></li></ol>	1	71.	Rosenfield, P.L., et al., <i>Development and verification of a schistosomiasis transmission</i>
<ul> <li>drug price of praziguantel in determining the cost-effectiveness of school-targeted treatment for Schistosoma mansoni using a population dynamic approach to evaluating the impact of school attendance on the unit cost and effectiveness of school-based schistosomiasis chemotherapy programmes. Parasitology, 2000. 121 (Pt 2), 171-83.</li> <li>Ndeffo Mbah, ML, et al., <i>Potential cost-effectiveness of school-assed schistosomiasis treatment for</i> reducing HIV transmission in Africa - The case of Zimbabwean women. Plos Negl Trop Dis, 2013, 78).</li> <li>Wang, XX., et al., <i>Projecting the Long-Term Impact of School- or Community-Based Mass-</i> <i>Treatment Interventions for Control of Schistosoma Infection</i>. Plos Negl Trop Dis, 2012. 6(11).</li> <li>Guyatt, H.L., Different approaches to modelling the cost-effectiveness of schistosomiasis control. Mem Inst Oswaldo Cruz, 1998. 93 (Suppl 1), 75-84.</li> <li>Turner, H.C., et al., <i>Reaching the London Declaration on Neglected Tropical Diseases goals for onchocerciasis: an economic evaluation of increasing the frequency of ivernectin treatment in Africa. Clin Infect Dis, 2014. 59(1): 923-32.</i></li> <li>Coffeng, L.E., et al., <i>African Programme for Onchocerciasis Control 1995–2015: Model-</i> estimated health impact and cost. PLoS Negl Trop Dis, 2013. 7(1), e2032.</li> <li>Lee, B.Y., et al., <i>The potential economic value of a hookworm</i> vaccine. Vaccine, 2011. 29(6), 1201-10.</li> <li>Guyatt, H.L., et al., <i>Control of Ascaris infection by chemotherapy: which is the most cost- effective option</i>? Trans R Soc Trop Med Hyg, 1995. 89(1), 16-20.</li> <li>Guyatt, H.L., et al., <i>Control of Ascaris infection by chemotherapy: which is the most cost- effective option</i>? Trans R Soc Trop Med Hyg, 1995. 89(1), 16-20.</li> <li>Custer, B., et al., <i>Modeling the economic value of a Chagas' disease therapeutic vaccine</i>. Hum Vaccine &amp; Immunother, 2012. 8(9), 123-1301.</li> <li>Sciurti, E., et al., <i>Cost-effectiveness of pathogen reduction methods for ELI</i></li></ul>	2		<i>model</i> . Am J Trop Med Hyg, 1977. <b>26</b> (3), 505-16.
<ul> <li>for Schistosoma mansoni using a population dynamic model. Trop Med Int Health, 1998. 3(6), 425-35.</li> <li>Carabin, H., et al., A population dynamic approach to evaluating the impact of school attendance on the unit cost and effectiveness of school-based schistosomiasis chemotherapy programmes. Parasislogy, 2000. 121 (P. 1), 171-83.</li> <li>Ndeffo Mbah, M.L., et al., Potential cost-effectiveness of schistosomiasis treatment for reducing HIV transmission in Africa - The case of Zimbabwean women. Plos Negl Trop Dis, 2013. 7(8).</li> <li>Wang, X.X., et al., Projecting the Long-Term Impact of School- or Community-Based Mass- Treatment Interventions for Control of Schistosoma Infection. Plos Negl Trop Dis, 2013. 7(8).</li> <li>Guyatt, H.L., Different approaches to modelling the cost-effectiveness of schistosomiasis control. Mem Inst Oswaldo Cruz, 1998. 93 (Suppl 1), 75-84.</li> <li>Turner, H.C., et al., Reaching the London Declaration on Neglected Tropical Dissess goals for onchocertais: an economic evaluation of increasing the frequency of ivermectin treatment in Africa. Clin Infect Dis, 2014. 59(7): 923-32.</li> <li>Coffeng, I.E., et al., African Programme for Onchocerciasis Control 1995-2015: Model- estimated health impact and cost. PLoS Negl Trop Dis, 2013. 7(1), e2032.</li> <li>Lee, B.Y., et al., The potential economic value of a hookworm vaccine. Vaccine, 2011. 29(6), 1201-10.</li> <li>Guyatt, H.L., et al., Control of Ascaris infection by chemotherapy: which is the most cost- effective option? Trans R Soc Trop Med Hyg, 1995. 89(1), 16-20.</li> <li>Guyatt, H.L., et al., Control of Ascaris infection by chemotherapy: which is the most cost- effective option? Trans R Soc Trop Med Hyg, 1993. 1301.</li> <li>Sicuri, E., et al., Economic evaluation of Chagas disease screening of pregnant Latin American women and of their infants in a non endemic area. Acta Trop, 2011. 118(2), 110-7.</li> <li>Lee, B.Y., et al., Mcodeling the economic value of a Chagas' disease</li></ul>		72.	
<ul> <li>3(6), 425-35.</li> <li>773. Carabin, H., et al., A population dynamic approach to evaluating the impact of school arendance on the unit cost and effectiveness of school-based schistosomiasis chemotherapy programmes. Parasitology, 2000. 121 (Pt 2), 171-83.</li> <li>74. Ndeffo Mbah, M.L., et al., Projecting the Long-Term Impact of School- or Community-Based Mass-Treatment Interventions for Control of Schistosoma Infection. Plos Negl Trop Dis, 2013. 7(8).</li> <li>75. Wang, X.X., et al., Projecting the Long-Term Impact of School- or Community-Based Mass-Treatment Interventions for Control of Schistosoma Infection. Plos Negl Trop Dis, 2012. 6(11).</li> <li>76. Guyatt, H.L., Different approaches to modelling the cost-effectiveness of schistosomiasis control. Mem Inst Oswaldo Cruz, 1998. 93 (Suppt 1), 75-84.</li> <li>77. Turner, H.C., et al., Reaching the London Declaration on Neglected Tropical Diseases goals for onchocerciasis: an economic evaluation of increasing the frequency of ivermectin treatment in Africa. Clin Infect Dis, 2014. 59(7): 923-32.</li> <li>78. Coffeng, L.E., et al., African Programme for Onchocerciasis Control 1995-2015: Model-estimated health impact and cost. PLoS Negl Trop Dis, 2013. 7(1), e2032.</li> <li>79. Lee, B.Y., et al., The potential economic value of a hookworm vaccine. Vaccine, 2011. 29(6), 1201-10.</li> <li>80. Guyatt, H.L., et al., A population dynamic approach to the cost-effectiveness analysis of mass antheliminic treatment: effectics of treatment for equency on Ascaris infection. Trans R Soc Trop Med Hyg, 1993. 87(5), 570-5.</li> <li>82. Lee, B.Y., et al., The potential economic value of a Chagas' disease therapeutic vaccine. Hum Vaccin &amp; Immunother, 2012. 8(9), 1293-1301.</li> <li>83. Sicuri, E., et al., The potential economic value of a Chagas' disease therapeutic vaccine. Hum Vaccin &amp; Immunother, 2012. 8(9), 1293-1301.</li> <li>84. Lee, B.Y., et al., The potential economic value of a Trypanosoma cruzi (Chagas disease) vaccine in Latin America. PUoS Negl Trop Dis, 2010</li></ul>			
<ol> <li>Carabin, H., et al., A population dynamic approach to evaluating the impact of school attendance on the unit cost and effectiveness of school-based schistosomiasis treatment for reducing HV transmission in Africa - The case of Zimbabwean women. Plos Negl Trop Dis, 2013, 7(8).</li> <li>Ndeffo Mbah, M.L., et al., Potential cost-effectiveness of schiotosomiasis treatment for reducing HV transmission in Africa - The case of Zimbabwean women. Plos Negl Trop Dis, 2013, 7(8).</li> <li>Wang, X.X., et al., Projecting the Long-Term Impact of School- or Community-Based Mass- Treatment Interventions for Control of Schistosoma Infection. Plos Negl Trop Dis, 2012. 6(11).</li> <li>Guyatt, H.L., Different approaches to modelling the cost-effectiveness of schistosomiasis control. Mem Inst Oswaldo Cruz, 1998, 93 (Suppl 1), 75-84.</li> <li>Turmer, H.C., et al., Reaching the London Declaration on Neglected Tropical Diseases goals for onchocerciasis: an economic evaluation of increasing the frequency of ivermectin treatment in Africa. Clin Infect Dis, 2014, 59(7): 923-32.</li> <li>Coffeng, L.E., et al., African Programme for Onchocerciasis: Control 1995-2015: Model- estimated health impact and cost. PLoS Negl Trop Dis, 2013. 7(1), e2032.</li> <li>Lee, B.Y., et al., The potential economic value of a hookworm vaccine. Vaccine, 2011. 29(6), 1201-10.</li> <li>Guyatt, H.L., et al., A control of Ascaris infection by chemotherapy: which is the most cost- effective opino? Trans R Soc Trop Med Hyg. 1995. 80(1), 16-20.</li> <li>Guyatt, H.L., et al., A population dynamic approach to the cost-effectiveness analysis of mass anthelminit: treatment: effect Play 2013. 80(1), 2031.</li> <li>Sicuri, E., et al., Economic evaluation of Chagas disease therapeutic vaccine. Hum Vaccin &amp; Immunother, 2012. 8(9), 1293-1301.</li> <li>Sicuri, E., et al., Economic evaluation of Chagas disease screening of pregnant Latin American woment effective enpiso of pathogen reduction technology as assessed using a mul</li></ol>			
<ul> <li>attendance on the unit cost and effectiveness of school-based schistosomiasis chemotherapy programmes. Parasitology. 2000. 121 (Pt 2), 171-83.</li> <li>74. Ndeffö Mbah, ML., et al., Potential cost-effectiveness of schistosomiasis treatment for reducing HIV transmission in Africa - The case of Zimbabwean women. Plos Negl Trop Dis, 2013, 7(8).</li> <li>75. Wang, X.X., et al., Projecting the Long-Term Impact of School- or Community-Based Mass- Treatment Interventions for Control of Schistosoma Infection. Plos Negl Trop Dis, 2012.</li> <li>6(11).</li> <li>76. Guyatt, H.L., Different approaches to modelling the cost-effectiveness of schistosomiasis control. Mem Inst Oswaldo Cruz, 1998. 93 (Suppl 1), 75-84.</li> <li>77. Turner, H.C., et al., Reaching the London Declaration on Neglected Tropical Diseases goals for onchocerciasis: an economic evaluation of increasing the frequency of ivermectin treatment in Africa. Clin Infect Dis, 2014. 59(7): 923-32.</li> <li>78. Coffeng, L.E., et al., African Programme for Onchocerciasis Control 1995-2015: Model- estimated health impact and cost. PLoS Negl Trop Dis, 2013. 7(1), e2023.</li> <li>79. Lee, B.Y., et al., The potential economic value of a hookworm vaccine. Vaccine, 2011. 29(6), 1201-10.</li> <li>80. Guyatt, H.L., et al., A population dynamic approach to the cost-effectiveness analysis of mass anthelimitir treatment: effects of treatment frequency on Ascaris infection. Trans R Soc Trop Med Hyg. 1993. 87(5), 570-5.</li> <li>82. Lee, B.Y., et al., Modeling the economic value of a Chagas' disease therapeut cvaccine. Hum Vaccin &amp; Immunother, 2012. 8(9), 1293-1301.</li> <li>83. Sicuri, E., et al., Economic evaluation of Chagas disease screening of pregnant Latin American women and of their infants in a non endemic area. Acta Trop. 2011. 118(2), 110-7.</li> <li>84. Lee, B.Y., et al., The cost-effectiveness of guitogen reduction technology as assessed using a multiple risk reduction model. Transfusion, 2010. 50(11), 2461-24</li></ul>			
<ul> <li>programmes. Parasitology, 2000. 121 (Pt 2), 171-83.</li> <li>Ndefio Mbah, M.L., et al., Potential cost-effectiveness of schistosomiasis treatment for</li> <li>reducing HIV transmission in Africa - The case of Zimbabwean vomen. Plos Negl Trop Dis, 2013. 7(8).</li> <li>Wang, X.X., et al., Projecting the Long-Term Impact of School- or Community-Based Mass- Treatment Interventions for Control of Schistosoma Infection. Plos Negl Trop Dis, 2012.</li> <li>6(11).</li> <li>Guyatt, H.L., Different approaches to modelling the cost-effectiveness of schistosomiasis control. Mem Inst Oswaldo Cruz, 1998. 93 (Suppl 1), 75-84.</li> <li>Turner, H.C., et al., Reaching the London Declaration on Neglected Tropical Diseases goals for onchocericasis: an economic evaluation of increasing the frequency of ivermectin treatment in Africa. Clin Infect Dis, 2014. 59(7): 923-32.</li> <li>Coffeng, L.E., et al., African Programme for Onchocerciasis Control 1995-2015: Model- estimated health impact and cost. PLoS Negl Trop Dis, 2013. 7(1), e2032.</li> <li>Lee, B.Y., et al., The potential economic value of a hookworm vaccine. Vaccine, 2011. 29(6), 1201-10.</li> <li>Guyatt, H.L., et al., Control of Ascaris infection by chemotherapy: which is the most cost- effective option? Trans R Soc Trop Med Hyg, 1995. 89(1), 16-20.</li> <li>Guyatt, H.L., et al., Control of Ascaris infection by chemotherapy: which is the most cost- effective option? Trans R Soc Trop Med Hyg, 1995. 89(1), 16-20.</li> <li>Lee, B.Y., et al., Modeling the economic value of a Chagas' disease therapeutic vaccine. Hum Vaccin &amp; Immunother, 2012. 8(9), 1293-1301.</li> <li>Sticuri, E., et al., Economic value of a Chagas' disease therapeutic vaccine. Hum Vaccine &amp; Inmunother, 2012. 8(9), 1293-1301.</li> <li>Sticuri, E., et al., Economic value of a Trypanosoma cruzi (Chagas disease) vaccine in Lain America. PLoS Negl Trop Dis, 2000. 50(11), 2461-2473.</li> <li>Kagapova, M., et al., Cost-effectiveness of chagas disease v</li></ul>		73.	
<ol> <li>Ndeffo Mbah, M.L., et al., Potential cost-effectiveness of schistosomiasis treatment for reducing HIV transmission in Africa - The case of Zinbabwean women. Plos Negl Trop Dis, 2013, 7(8).</li> <li>Wang, X.X., et al., Projecting the Long-Term Impact of School- or Community-Based Mass- Treatment Interventions for Control of Schistosoma Infection. Plos Negl Trop Dis, 2012. 6(11).</li> <li>Guyatt, H.L., Different approaches to modelling the cost-effectiveness of schistosomiasis control. Mem Inst Oswaldo Cruz, 1998. 93 (Suppl 1), 75-84.</li> <li>Turner, H.C., et al., Reaching the London Declaration on Neglected Tropical Diseases goals for onchocerciasis: an economic evaluation of increasing the frequency of ivermectin treatment in Africa. Clin Infect Dis, 2014. 59(7): 923-32.</li> <li>Coffeng, L.E., et al., African Programme for Onchocerciasis Control 1995–2015: Model- estimated health impact and cost. PLoS Negl Trop Dis, 2013. 7(1), e2032.</li> <li>Lee, B.Y., et al., The potential economic value of a hookworm vaccine. Vaccine, 2011. 29(6), 1201-10.</li> <li>Guyatt, H.L., et al., Control of Ascaris infection by chemotherapy: which is the most cost- effective option? Trans R Soc Trop Med Hyg. 1995. 89(1), 16-20.</li> <li>Guyatt, H.L., et al., A population dynamic approach to the cost-effectiveness analysis of mass anthelmintic treatment: effects of treatment frequency on Ascaris infection. Trans R Soc Trop Med Hyg. 1993. 87(5), 570-5.</li> <li>Lee, B.Y., et al., Modeling the economic value of a Chagas' disease therapeutic vaccine. Hum Vaccin &amp; Immunother, 2012. 8(9), 1293-1301.</li> <li>Sicuri, E., et al., Economic evaluation of Chagas disease screening of pregnant Latin American women and of their infants in a non endemic area. Acta Trop, 2011. 118(2), 110-7.</li> <li>Lee, B.Y., et al., The potential economics, 2010, 20(1), 2461-2473.</li> <li>Agapova, M., et al., Cost-effectiveness of publogen reduction technology as assessed using a multiple risk reduction</li></ol>			
<ul> <li>reducing HIV transmission in Africa - The case of Zimbabwean women. Plos Negl Trop Dis, 2013. 7(8).</li> <li>Wang, X.X., et al., Projecting the Long-Term Impact of School- or Community-Based Mass-Treatment Interventions for Control of Schistosoma Infection. Plos Negl Trop Dis, 2012. 6(11).</li> <li>Guyatt, H.L., Different approaches to modelling the cost-effectiveness of schistosomiasis control. Mem Inst Oswaldo Cruz, 1998. 93 (Suppl 1), 75-84.</li> <li>Turner, H.C., et al., Reaching the London Declaration on Neglected Tropical Diseases goals for onchocerciasis: an economic evaluation of increasing the frequency of ivermectin treatment in Africa. Clin Infect Dis, 2014. 59(7): 923-32.</li> <li>Coffteng, L.E., et al., African Programme for Onchocerciasis Control 1995–2015: Model-estimated health impact and cost. PLoS Negl Trop Dis, 2013. 7(1), e2032.</li> <li>Lee, B.Y., et al., The potential economic value of a hookworm vaccine. Vaccine, 2011. 29(6), 1201-10.</li> <li>Guyatt, H.L., et al., A population dynamic approach to the cost-effectiveness analysis of mass anthelminitic treatment: effects of treatment frequency on Ascaris infection. Trans R Soc Trop Med Hyg, 1995. 89(1), 16-20.</li> <li>Guyatt, H.L., et al., A population dynamic approach to the cost-effectiveness analysis of mass anthelminitic treatment: effects of treatment frequency on Ascaris infection. Trans R Soc Trop Med Hyg, 1993. 87(5), 570-5.</li> <li>Lee, B.Y., et al., Modeling the economic value of a Chagas' disease therapeutic vaccine. Hum Vaccin &amp; Immunother, 2012. 8(9), 1293-1301.</li> <li>Sicuri, E., et al., Economic value of Chagas disease screening of pregnant Latin America value on of Chagas disease screening of pregnant Latin America value on model. Transfusion, 2010. 50(10), 2220-32.</li> <li>Lee, B.Y., et al., Cost-effectiveness of screening the US blood supply for Trypanosoma cruzi. Transfusion, 2010. 50(10), 2220-32.</li> <li>Xuacuez-Prokopec, G.M., et al., Cost-effectiveness of</li></ul>			
<ol> <li>2013. 7(8).</li> <li>Wang, X.X., et al., Projecting the Long-Term Impact of School- or Community-Based Mass-Treatment Interventions for Control of Schistosoma Infection. Plos Negl Trop Dis, 2012.</li> <li>6(11).</li> <li>Guyatt, H.L., Different approaches to modelling the cost-effectiveness of schistosomiasis control. Mem Inst Oswaldo Cruz, 1998. <b>93</b> (Suppl 1), 75-84.</li> <li>Turner, H.C., et al., Reaching the London Declaration on Neglected Tropical Diseases goals for onchocerciasis: an economic evaluation of increasing the frequency of ivermectin treatment in Africa. Clin Infect Dis, 2014. <b>59</b>(7): 923-32.</li> <li>Coffeng, L.E., et al., African Programme for Onchocerciasis Control 1995–2015: Modelestimated health impact and cost. PLoS Negl Trop Dis, 2013. <b>7</b>(1), e2032.</li> <li>Lee, B.Y., et al., The potential economic value of a hookworm vaccine. Vaccine, 2011. <b>29</b>(6), 1201-10.</li> <li>Guyatt, H.L., et al., Control of Ascaris infection by chemotherapy: which is the most cost-effective option? Trans R Soc Trop Med Hyg, 1995. <b>89</b>(1), 16-20.</li> <li>Guyatt, H.L., et al., A population dynamic approach to the cost-effectiveness analysis of mass anthelimitic treatment: effects of treatment frequency on Ascaris infection. Trans R Soc Trop Med Hyg, 1993. <b>87</b>(5), 570-5.</li> <li>Lee, B.Y., et al., Modeling the economic value of a Chagas' disease therapeutic vaccine. Hum Vaccin &amp; Immunother, 2012. <b>8</b>(9), 1293-1301.</li> <li>Sicuri, E., et al., Economic evaluation of Chagas disease screening of pregnant Latin American women and of their infants in a non endemic area. Acta Trop, 2011. <b>118</b>(2), 110-7.</li> <li>Lee, B.Y., et al., The cost-effectiveness of pathogen reduction technology as assessed using a multiple risk reduction model. Transfusion, 2010. <b>5</b>(11), 2461-2473.</li> <li>Agapova, M., et al., Cost-effectiveness of pathogen fueducion technology as assessed using a multiple risk reduction model. Schoffectiveness of chagas disease vector contro</li></ol>		74.	
<ol> <li>Wang, X.X., et al., Projecting the Long-Term Impact of School- or Community-Based Mass- Treatment Interventions for Control of Schistosoma Infection. Plos Negl Trop Dis, 2012. 6(11).</li> <li>Guyatt, H.L., Different approaches to modelling the cost-effectiveness of schistosomiasis control. Mem Inst Oswaldo Cruz, 1998. 93 (Suppl 1), 75-84.</li> <li>Turner, H.C., et al., Reaching the London Declaration on Neglected Tropical Diseases goals for onchocerciasis: an economic evaluation of increasing the frequency of ivermectin treatment in Africa. Clin Infect Dis, 2014. 59(7): 923-32.</li> <li>Coffteng, L.E., et al., African Programme for Onchocerciasis Control 1995–2015: Model- estimated health impact and cost. PLoS Negl Trop Dis, 2013. 7(1), e2032.</li> <li>Lee, B.Y., et al., The potential economic value of a hookworm vaccine. Vaccine, 2011. 29(6), 1201-10.</li> <li>Guyatt, H.L., et al., A population dynamic approach to the cost-effectiveness analysis of mass anthelminitic treatment: effects of treatment frequency on Ascaris infection. Trans R Soc Trop Med Hyg, 1993. 87(5), 570-5.</li> <li>Lee, B.Y., et al., Modeling the economic value of a Chagas' disease therapeutic vaccine. Hum Vaccin &amp; Immunother, 2012. 8(9), 1293-1301.</li> <li>Sicuri, E., et al., <i>Atopotential economic value of a Chagas' disease therapeutic vaccine</i>. Hum Vaccin &amp; Immunother, 2012. 8(9), 1293-1301.</li> <li>Sicuri, E., et al., The potential economic value of a Trypanosoma cruzi (Chagas disease) vaaccine in Latin America. PLoS Negl Trop Dis, 2010. 4(12), e916.</li> <li>Custer, B., et al., The cost-effectiveness of pathogen reduction technology as assessed using a multiple risk reduction model. Transfusion, 2010. 50(11), 2461-2473.</li> <li>Agapova, M., et al., Cost-effectiveness of fochagas disease vector control strategies in Northwestern Argentina. PLoS Negl Trop Dis, 2009. 3(1), e363.</li> <li>Vasquez-Prokopec, G.M., et al., Cost-effectiveness of chagas disease vector control strategies</li></ol>			
<ul> <li><i>Treatment Interventions for Control of Schistosoma Infection.</i> Plos Negl Trop Dis, 2012. 6(11).</li> <li>Guyatt, H.L., Different approaches to modelling the cost-effectiveness of schistosomiasis control. Mem Inst Oswaldo Cruz, 1998. 93 (Suppl 1), 75-84.</li> <li>Turner, H.C., et al., Reaching the London Declaration on Neglected Tropical Diseases goals for onchocerciasis: an economic evaluation of increasing the frequency of ivermectin treatment in Africa. Clin Infect Dis, 2014. 59(7): 923-32.</li> <li>Coffeng, L.E., et al., African Programme for Onchocerciasis Control 1995-2015: Model- estimated health impact and cost. PLoS Negl Trop Dis, 2013. 7(1), e2032.</li> <li>Lee, B.Y., et al., The potential economic value of a hookworm vaccine. Vaccine, 2011. 29(6), 1201-10.</li> <li>Guyatt, H.L., et al., Control of Ascaris infection by chemotherapy: which is the most cost- effective option? Trans R Soc Trop Med Hyg, 1995. 89(1), 16-20.</li> <li>Guyatt, H.L., et al., A population dynamic approach to the cost-effectiveness analysis of mass anthelminic treatment: effects of treatment frequency on Ascaris Infection. Trans R Soc Trop Med Hyg, 1993. 87(5), 570-5.</li> <li>Lee, B.Y., et al., Modeling the economic value of a Chagas' disease therapeutic vaccine. Hum Vaccin &amp; Immunother, 2012. 8(9), 1293-1301.</li> <li>Sicuri, E., et al., <i>Economic evaluation of Chagas disease screening of pregnant Latin American women and of their infants in a non endenic area.</i> Acta Trop, 2011. 118(2), 110-7.</li> <li>Lee, B.Y., et al., <i>The potential economic value of a Trypanosoma cruzi (Chagas disease)</i> vaccine in Latin America. PLoS Negl Trop Dis, 2010. 4(12), e916.</li> <li>Custer, B., et al., <i>The cost-effectiveness of pathogen reduction technology as assessed using a multiple risk reduction model</i>. Transfusion, 2010. 50(11), 2461-2473.</li> <li>Agapova, M., et al., <i>Cost-effectiveness of inflormation methods for ELISA serology testing of Trypanosoma cruzi in California blood bank</i></li></ul>			
<ol> <li>6(11),</li> <li>6(11),</li> <li>Guyatt, H.L., Different approaches to modelling the cost-effectiveness of schistosomiasis control. Mem Inst Oswaldo Cruz, 1998. 93 (Suppl 1), 75-84.</li> <li>77. Turner, H.C., et al., Reaching the London Declaration on Neglected Tropical Diseases goals for onchocerciasis: an economic evaluation of increasing the frequency of ivermectin treatment in Africa. Clin Infect Dis, 2014. 59(7), 923-32.</li> <li>78. Coffeng, L.E., et al., African Programme for Onchocerciasis Control 1995–2015: Model- estimated health impact and cost. PLoS Negl Trop Dis, 2013. 7(1), e2032.</li> <li>79. Lee, B.Y., et al., The potential economic value of a hookworm vaccine. Vaccine, 2011. 29(6), 1201-10.</li> <li>80. Guyatt, H.L., et al., Control of Ascaris infection by chemotherapy: which is the most cost- effective option? Trans R Soc Trop Med Hyg, 1995. 89(1), 16-20.</li> <li>81. Guyatt, H.L., et al., A population dynamic approach to the cost-effectiveness analysis of mass anthelminitic treatment: effects of treatment frequency on Ascaris infection. Trans R Soc Trop Med Hyg, 1993. 87(5), 570-5.</li> <li>82. Lee, B.Y., et al., Modeling the economic value of a Chagas' disease therapeutic vaccine. Hum Vaccin &amp; Immunother, 2012. 8(9), 1293-1301.</li> <li>83. Sicuti, E., et al., <i>Enotomic evaluation of Chagas disease screening of pregnant Latin American women and of their infants in a non endemic area.</i> Acta Trop, 2011. 118(2), 110-7.</li> <li>84. Lee, B.Y., et al., The potential economic value of a Trypanosoma cruci (Chagas disease) vaccine &amp; I.Atin American. PLoS Negl Trop Dis, 2010. 4(12), e916.</li> <li>85. Custer, B., et al., The cost-effectiveness of pathogen reduction technology as assessed using a multiple risk reduction model. Transfusion, 2010. 50(1), 2461-2473.</li> <li>86. Agapova, M., et al., Cost-effectiveness of chagas disease sector control strategies in Northwestern Argentina. PLoS Negl Trop Dis, 2009. 3(1), e363.</li> <li>89. Wil</li></ol>		75.	• • • • • •
<ol> <li>Guyatt, H.L., Different approaches to modelling the cost-effectiveness of schistosomiasis control. Mem Inst Oswaldo Cruz, 1998. 93 (Suppl 1), 75-84.</li> <li>Turner, H.C., et al., Reaching the London Declaration on Neglected Tropical Diseases goals for onchocerciasis: an economic evaluation of increasing the frequency of ivermectin treatment in Africa. Clin Infect Dis, 2014. 59(7): 923-32.</li> <li>Coffeng, L.E., et al., African Programme for Onchocerciasis Control 1995–2015: Model- estimated health impact and cost. PLoS Negl Trop Dis, 2013. 7(1), e2032.</li> <li>Lee, B.Y., et al., The potential economic value of a hookworm vaccine. Vaccine, 2011. 29(6), 1201-10.</li> <li>Guyatt, H.L., et al., Control of Ascaris infection by chemotherapy: which is the most cost- effective option? Trans R Soc Trop Med Hyg, 1995. 89(1), 16-20.</li> <li>Guyatt, H.L., et al., A population dynamic approach to the cost-effectiveness analysis of mass anthelminic treatment: effects of treatment frequency on Ascaris infection. Trans R Soc Trop Med Hyg, 1993. 87(5), 570-5.</li> <li>Lee, B.Y., et al., Modeling the economic value of a Chagas' disease therapeutic vaccine. Hum Vaccin &amp; Immunother, 2012. 8(9), 1293-1301.</li> <li>Sicuri, E., et al., Economic evaluation of Chagas disease screening of pregnant Latin American women and of their infants in a non endemic area. Acta Trop, 2011. 118(2), 110-7.</li> <li>Lee, B.Y., et al., The cost-effectiveness of pathogen reduction technology as assessed using a multiple risk reduction model. Transfusion, 2010. 4(12), e916.</li> <li>Custer, B., et al., The cost-effectiveness of pathogen reduction technology as assessed using a multiple risk reduction model. Transfusion, 2010. 50(11), 2461-2473.</li> <li>Agapova, M., et al., Cost-effectiveness of pathogen reduction technology as assessed using a multiple risk reduction model. Transfusion, 2010. 4(12), e916.</li> <li>Vazquez-Prokopec, G.M., et al., Cost-effectiveness of chagas disease vector con</li></ol>			
<ol> <li>control. Mem Inst Oswaldo Cruz, 1998. 93 (Suppl 1), 75-84.</li> <li>Turner, H.C., et al., Reaching the London Declaration on Neglected Tropical Diseases goals for onchocerciasis: an economic evaluation of increasing the frequency of ivermectin treatment in Africa. Clin Infect Dis, 2014. 59(7): 923-32.</li> <li>Coffeng, L.E., et al., African Programme for Onchocerciasis Control 1995–2015: Model- estimated health impact and cost. PLoS Negl Trop Dis, 2013. 7(1), e2032.</li> <li>Lee, B.Y., et al., The potential economic value of a hookworm vaccine. Vaccine, 2011. 29(6), 1201-10.</li> <li>Guyatt, H.L., et al., Control of Ascaris infection by chemotherapy: which is the most cost- effective option? Trans R Soc Trop Med Hyg, 1995. 89(1), 16-20.</li> <li>Guyatt, H.L., et al., A population dynamic approach to the cost-effectiveness analysis of mass anthelminitic treatment: effects of treatment frequency on Ascaris infection. Trans R Soc Trop Med Hyg, 1993. 87(5), 570-5.</li> <li>Lee, B.Y., et al., Modeling the economic value of a Chagas' disease therapeutic vaccine. Hum Vaccin &amp; Immunother, 2012. 8(9), 1293-1301.</li> <li>Sicuri, E., et al., Economic evaluation of Chagas disease screening of pregnant Latin American women and of their infants in a non endemic area. Acta Trop. 2011. 118(2), 110-7.</li> <li>Lee, B.Y., et al., The potential economic value of a Trypanosoma cruzi (Chagas disease) vaccine in Latin America. PLoS Negl Trop Dis, 2010. 4(12), e916.</li> <li>Custer, B., et al., The cost-effectiveness of pathogen reduction technology as assessed using a multiple risk reduction model. Transfusion, 2010. 50(11), 2461-2473.</li> <li>Agapova, M., et al., Cost-effectiveness of reneming the US blood supply for Trypanosoma cruzi. Transfusion, 2010. 50(10), 2220-32.</li> <li>Vazquez-Prokopec, G.M., et al., Cost-effectiveness of chagas disease vector control strategies in Northwestern Argenitma. PLoS Negl Trop Dis, 2009. 3(1), e363.</li> <li>Wilson, L.S., et al., Co</li></ol>		-	
<ol> <li>Turner, H.C., et al., Reaching the London Declaration on Neglected Tropical Diseases goals for onchocerciasis: an economic evaluation of increasing the frequency of ivermectin treatment in Africa. Clin Infect Dis, 2014. <b>39</b>(7): 923-32.</li> <li>Coffeng, L.E., et al., African Programme for Onchocerciasis Control 1995–2015: Model- estimated health impact and cost. PLoS Negl Trop Dis, 2013. <b>7</b>(1), e2032.</li> <li>Lee, B.Y., et al., The potential economic value of a hookworm vaccine. Vaccine, 2011. <b>29</b>(6), 1201-10.</li> <li>Guyatt, H.L., et al., Control of Ascaris infection by chemotherapy: which is the most cost- effective option? Trans R Soc Trop Med Hyg, 1995. <b>89</b>(1), 16-20.</li> <li>Guyatt, H.L., et al., A population dynamic approach to the cost-effectiveness analysis of mass anthelmintic treatment: effects of treatment frequency on Ascaris infection. Trans R Soc Trop Med Hyg, 1993. <b>87</b>(5), 570-5.</li> <li>Lee, B.Y., et al., <i>Modeling the economic value of a Chagas' disease therapeutic vaccine</i>. Hum Vaccin &amp; Immunother, 2012. <b>8</b>(9), 1293-1301.</li> <li>Sicuri, E., et al., <i>Economic evaluation of Chagas disease screening of pregnant Latin American women and of their infants in a non endemic area.</i> Acta Trop, 2011. <b>118</b>(2), 110-7.</li> <li>Lee, B.Y., et al., <i>The potential economic value of a Trypanosoma cruzi (Chagas disease) vaccine in Latin America.</i> PLoS Negl Trop Dis, 2010. <b>4</b>(12), e916.</li> <li>Custer, B., et al., <i>The cost-effectiveness of pathogen reduction technology as assessed using a multiple risk reduction model.</i> Transfusion, 2010. <b>50</b>(11), 2461-2473.</li> <li>Agapova, M., et al., <i>Cost-effectiveness of chagas disease vector control strategies in Northwestern Argentina.</i> PLoS Negl Trop Dis, 2009. <b>3</b>(1), e363.</li> <li>Wilson, L.S., et al., <i>Cost-effectiveness of Chagas disease vector control strategies in Northwestern Argentina.</i> PLoS Negl Trop Dis, 2005. <b>73</b>(5), 901-10.</li> <li>Schoffeld, C.J. and Dias, J.C., A cost-benefit analy</li></ol>		76.	• • • • • • •
<ul> <li>for onchocerciasis: an economic evaluation of increasing the frequency of ivermectin treatment in Africa. Clin Infect Dis, 2014. 59(7): 923-32.</li> <li>78. Coffeng, L.E., et al., African Programme for Onchocerciasis Control 1995–2015: Model- estimated health impact and cost. PLOS Negl Trop Dis, 2013. 7(1), e2032.</li> <li>79. Lee, B.Y., et al., The potential economic value of a hookworm vaccine. Vaccine, 2011. 29(6), 1201-10.</li> <li>80. Guyatt, H.L., et al., Control of Ascaris infection by chemotherapy: which is the most cost- effective option? Trans R Soc Trop Med Hyg, 1995. 89(1), 16-20.</li> <li>81. Guyatt, H.L., et al., A population dynamic approach to the cost-effectiveness analysis of mass anthelmintic treatment: effects of treatment frequency on Ascaris infection. Trans R Soc Trop Med Hyg, 1993. 87(5), 570-5.</li> <li>82. Lee, B.Y., et al., Modeling the economic value of a Chagas' disease therapeutic vaccine. Hum Vaccin &amp; Immunother, 2012. 8(9), 1293-1301.</li> <li>83. Sicuri, E., et al., Economic evaluation of Chagas disease screening of pregnant Latin American women and of their infants in a non endemic area. Acta Trop, 2011. 118(2), 110-7.</li> <li>84. Lee, B.Y., et al., The potential economic value of a Trypanosoma cruzi (Chagas disease) vaccine in Latin America. PLoS Negl Trop Dis, 2010. 4(12), e916.</li> <li>85. Custer, B., et al., The cost-effectiveness of pathogen reduction technology as assessed using a multiple risk reduction model. Transfusion, 2010. 50(11), 2461-2473.</li> <li>86. Agapova, M., et al., Cost-effectiveness of screening the US blood supply for Trypanosoma cruzi, Transfusion, 2010. 50(10), 2220-32.</li> <li>87. Vazquez-Prokopec, G.M., et al., Cost-effectiveness of chagas disease vector control strategies in Northwestern Argentina. PLoS Negl Trop Dis, 2009. 3(1), e363.</li> <li>88. Wilson, L.S., et al., Cost-effectiveness of Chagas disease interventions in latin america and the Caribbear: Markov models. Am J Trop Med Hyg,</li></ul>			
<ul> <li>treatment in Africa. Clin Infect Dis, 2014. 59(7): 923-32.</li> <li>Treatment in Africa. Clin Infect Dis, 2013. 50(7): 923-32.</li> <li>Coffeng, L.E., et al., African Programme for Onchocerciais Control 1995–2015: Model- estimated health impact and cost. PLoS Negl Trop Dis, 2013. 7(1), e2032.</li> <li>Lee, B.Y., et al., The potential economic value of a hookworm vaccine. Vaccine, 2011. 29(6), 1201-10.</li> <li>Guyatt, H.L., et al., Control of Ascaris infection by chemotherapy: which is the most cost- effective option? Trans R Soc Trop Med Hyg, 1995. 89(1), 16-20.</li> <li>Guyatt, H.L., et al., A population dynamic approach to the cost-effectiveness analysis of mass anthelmintic treatment: effects of treatment frequency on Ascaris infection. Trans R Soc Trop Med Hyg, 1993. 87(5), 570-5.</li> <li>Lee, B.Y., et al., Modeling the economic value of a Chagas' disease therapeutic vaccine. Hum Vaccin &amp; Immunother, 2012. 8(9), 1293-1301.</li> <li>Sicuri, E., et al., <i>Economic evaluation of Chagas disease screening of pregnant Latin American women and of their infants in a non endemic area</i>. Acta Trop, 2011. 118(2), 110-7.</li> <li>Lee, B.Y., et al., The potential economic value of a Trypanosoma cruzi (Chagas disease) vaccine in Latin America. PLoS Negl Trop Dis, 2010. 4(12), e916.</li> <li>Custer, B., et al., The cost-effectiveness of screening the US blood supply for Trypanosoma cruzi. Transfusion, 2010. 50(10), 2220-32.</li> <li>Vazquez-Prokopec, G.M., et al., Cost-effectiveness of chagas disease vector control strategies in Northwestern Argentina. PLoS Negl Trop Dis, 2009. 3(1), e363.</li> <li>Wilson, L.S., et al., Cost-effectiveness of Chagas disease vector control strategies in Northwestern Argentina. PLoS Negl Trop Dis, 2009. 3(1), e363.</li> <li>Wilson, L.S., et al., Cost-effectiveness of Chagas disease interventions in latin america and the Caribbean: Markov models. Am J Trop Med Hyg, 2005. 73(5), 901-10.</li> <li>Schofield, C.J. and Dias, J.C., A</li></ul>		77.	
<ol> <li>R. Coffeng, L.E., et al., African Programme for Onchocerciasis Control 1995–2015: Model- estimated health impact and cost. PLoS Negl Trop Dis, 2013. 7(1), e2032.</li> <li>I.ee, B.Y., et al., The potential economic value of a hookworm vaccine. Vaccine, 2011. 29(6), 1201-10.</li> <li>Guyatt, H.L., et al., Control of Ascaris infection by chemotherapy: which is the most cost- effective option? Trans R Soc Trop Med Hyg, 1995. 89(1), 16–20.</li> <li>Guyatt, H.L., et al., A population dynamic approach to the cost-effectiveness analysis of mass antheliminic treatment: effects of treatment frequency on Ascaris infection. Trans R Soc Trop Med Hyg, 1993. 87(5), 570-5.</li> <li>E. Lee, B.Y., et al., Modeling the economic value of a Chagas' disease therapeutic vaccine. Hum Vaccin &amp; Immunother, 2012. 8(9), 1293-1301.</li> <li>Sicuri, E., et al., Economic evaluation of Chagas disease screening of pregnant Latin American women and of their infants in a non endemic area. Acta Trop, 2011. 118(2), 110-7.</li> <li>Lee, B.Y., et al., The ootential economic value of a Trypanosoma cruzi (Chagas disease) vaccine in Latin America. PLoS Negl Trop Dis, 2010. 4(12), e916.</li> <li>Custer, B., et al., The cost-effectiveness of pathogen reduction technology as assessed using a multiple risk reduction model. Transfusion, 2010. 50(11), 2461-2473.</li> <li>Agapova, M., et al., Cost-effectiveness of screening the US blood supply for Trypanosoma cruzi. Transfusion, 2010. 50(10), 2220-32.</li> <li>Vazquez-Prokopec, G.M., et al., Cost-effectiveness of chagas disease vector control strategies in Northwestern Argentina. PLoS Negl Trop Dis, 2009. 3(1), e363.</li> <li>Wilson, L.S., et al., Cost-effectiveness of Chagas disease interventions in latin america and the Caribbean: Markov models. Am J Trop Med Hyg, 2005. 73(5), 901-10.</li> <li>Schofield, C.J. and Dias, J.C., A cost-benefit analysis of Chagas disease control. Mem Inst Oswaldo Cruz, 1991. 86(3), 285-95.</li> <li>Barbu, C., et al., E</li></ol>			
<ul> <li>estimated health impact and cost. PLoS Negl Trop Dis, 2013. 7(1), e2032.</li> <li>estimated health impact and cost. PLoS Negl Trop Dis, 2013. 7(1), e2032.</li> <li>Lee, B.Y., et al., <i>The potential economic value of a hookworm vaccine</i>. Vaccine, 2011. 29(6), 1201-10.</li> <li>Guyatt, H.L., et al., <i>Control of Ascaris infection by chemotherapy: which is the most cost-effective option</i>? Trans R Soc Trop Med Hyg, 1995. 89(1), 16-20.</li> <li>Guyatt, H.L., et al., <i>A population dynamic approach to the cost-effectiveness analysis of mass anthelminic treatment: effects of treatment frequency on Ascaris infection</i>. Trans R Soc Trop Med Hyg, 1993. 87(5), 570-5.</li> <li>Lee, B.Y., et al., <i>Modeling the economic value of a Chagas' disease therapeutic vaccine</i>. Hum Vaccin &amp; Immunother, 2012. 8(9), 1293-1301.</li> <li>Sicuri, E., et al., <i>Economic evaluation of Chagas disease screening of pregnant Latin America women and of their infants in a non endemic area</i>. Acta Trop, 2011. 118(2), 110-7.</li> <li>Lee, B.Y., et al., <i>The potential economic value of a Trypanosoma cruzi (Chagas disease) vaccine in Latin America</i>. PLoS Negl Trop Dis, 2010. 4(12), e916.</li> <li>Custer, B., et al., <i>The cost-effectiveness of screening the US blood supply for Trypanosoma cruzi</i>. Transfusion, 2010. 50(10), 220-32.</li> <li>Vazquez-Prokopec, G.M., et al., <i>Cost-effectiveness of chagas disease vector control strategies in Northwestern Argentina</i>. PLoS Negl Trop Dis, 2009. 3(1), e363.</li> <li>Wilson, L.S., et al., <i>Cost-effectiveness of Chagas disease interventions in latin america and the Caribbean: Markov models</i>. Am J Trop Med Hyg, 2005. 73(5), 901-10.</li> <li>Schofield, C.J. and Dias, J.C., A cost-benefit analysis of Chagas disease control. Mem Inst Oswaldo Cruz, 1991. 86(3), 285-95.</li> <li>Barbu, C., et al., <i>Evaluation of spatially targeted strategies to control non-domiciliated Triatom adminidia vector of Chagas diseases</i>. PLoS Negl Trop Dis, 2011. 5(5), e1045.</li> <li>Castillo-Riquelme, M., et al., <i>Modelling geographic v</i></li></ul>		-	
<ol> <li>P. Lee, B.Y., et al., <i>The potential economic value of a hookworm vaccine</i>. Vaccine, 2011. 29(6), 1201-10.</li> <li>Guyatt, H.L., et al., <i>Control of Ascaris infection by chemotherapy: which is the most cost-effective option</i>? Trans R Soc Trop Med Hyg, 1995. 89(1), 16-20.</li> <li>Guyatt, H.L., et al., <i>A population dynamic approach to the cost-effectiveness analysis of mass anthelmintic treatment: effects of treatment frequency on Ascaris infection</i>. Trans R Soc Trop Med Hyg, 1993. 87(5), 570-5.</li> <li>Lee, B.Y., et al., <i>Modeling the economic value of a Chagas' disease therapeutic vaccine</i>. Hum Vaccin &amp; Immunother, 2012. 8(9), 1293-1301.</li> <li>Sicuri, E., et al., <i>Economic evaluation of Chagas disease screening of pregnant Latin American women and of their infants in a non endemic area.</i> Acta Trop, 2011. 118(2), 110-7.</li> <li>Lee, B.Y., et al., <i>The potential economic value of a Trypanosoma cruzi (Chagas disease) vaccine in Latin America.</i> PLoS Negl Trop Dis, 2010. 4(12), e916.</li> <li>Custer, B., et al., <i>Cost-effectiveness of pathogen reduction technology as assessed using a multiple risk reduction model.</i> Transfusion, 2010. 50(11), 2461-2473.</li> <li>Agapova, M., et al., <i>Cost-effectiveness of screening the US blood supply for Trypanosoma cruzi.</i> Transfusion, 2010. 50(10), 2220-32.</li> <li>Vazquez-Prokopec, G.M., et al., <i>Cost-effectiveness of chagas disease vector control strategies in Northwestern Argentina.</i> PLoS Negl Trop Dis, 2009. 3(1), e363.</li> <li>Wilson, L.S., et al., <i>Cost-effectiveness of Chagas disease therventions in latin america and the Caribbean: Markw models.</i> Am J Trop Med and Hyg, 2008. 79(1), 53-68.</li> <li>Wilson, L.S., et al., <i>Cost-effectiveness of Chagas disease control.</i> Mem Inst Oswaldo Cruz, 1991. 86(3), 285-95.</li> <li>Barbu, C., et al., <i>Evaluation of spatially targeted strategies to control non-domiciliated Triatom admidiata vector of Chagas disease: 1he control non-domiciliated Triatom admidiata vector of </i></li></ol>		78.	
<ol> <li>1201-10.</li> <li>Guyatt, H.L., et al., <i>Control of Ascaris infection by chemotherapy: which is the most cost-effective option?</i> Trans R Soc Trop Med Hyg, 1995. <b>89</b>(1), 16-20.</li> <li>Guyatt, H.L., et al., <i>A population dynamic approach to the cost-effectiveness analysis of mass anthelmintic treatment: effects of treatment frequency on Ascaris infection</i>. Trans R Soc Trop Med Hyg, 1993. <b>87</b>(5), 570-5.</li> <li>Lee, B.Y., et al., <i>Modeling the economic value of a Chagas' disease therapeutic vaccine</i>. Hum Vaccin &amp; Immunother, 2012. <b>8</b>(9), 1293-1301.</li> <li>Sicuri, E., et al., <i>Economic evaluation of Chagas disease screening of pregnant Latin American women and of their infants in a non endemic area</i>. Acta Trop, 2011. <b>118</b>(2), 110-7.</li> <li>Lee, B.Y., et al., <i>The potential economic value of a Trypanosoma cruzi (Chagas disease) vaccine in Latin America</i>. PLoS Negl Trop Dis, 2010. <b>4</b>(12), e916.</li> <li>Custer, B., et al., <i>The cost-effectiveness of pathogen reduction technology as assessed using a multiple risk reduction model</i>. Transfusion, 2010. <b>50</b>(11), 2461-2473.</li> <li>Agapova, M., et al., <i>Cost-effectiveness of screening the US blood supply for Trypanosoma cruzi</i>. Transfusion, 2010. <b>50</b>(10), 2220-32.</li> <li>Vazquez-Prokopee, G.M., et al., <i>Cost-effectiveness of chagas disease vector control strategies in Northwestern Argentina</i>. PLoS Negl Trop Dis, 2009. <b>3</b>(1), e363.</li> <li>Wilson, L.S., et al., <i>Cost-effectiveness of Chagas disease interventions in latin america and the Caribbean: Markov models</i>. Am J Trop Med Hyg, 2005. <b>73</b>(5), 901-10.</li> <li>Schoffeld, C.J. and Dias, J.C., A cost-benefit analysis of Chagas disease control. Mem Inst Oswaldo Cruz, 1991. <b>86</b>(3), 285-95.</li> <li>Barbu, C., et al., <i>Evaluation of spatially targeted strategies to control non-domiciliated Triatoma dimidiata vector of Chagas disease: the example of Chagas disease.</i> J Health Econ, 2008. <b>27</b>(2), 405-26.</li> <li>Basombrio, M.A., et al., <i>A cost-be</i></li></ol>		70	
<ol> <li>80. Guyatt, H.L., et al., Control of Ascaris infection by chemotherapy: which is the most cost-effective option? Trans R Soc Trop Med Hyg, 1995. 89(1), 16-20.</li> <li>81. Guyatt, H.L., et al., A population dynamic approach to the cost-effectiveness analysis of mass anthelimitic treatment: effects of treatment frequency on Ascaris infection. Trans R Soc Trop Med Hyg, 1993. 87(5), 570-5.</li> <li>82. Lee, B.Y., et al., Modeling the economic value of a Chagas' disease therapeutic vaccine. Hum Vaccin &amp; Immunother, 2012. 8(9), 1293-1301.</li> <li>83. Sicuri, E., et al., Economic evaluation of Chagas disease screening of pregnant Latin American women and of their infants in a non endemic area. Acta Trop, 2011. 118(2), 110-7.</li> <li>84. Lee, B.Y., et al., The potential economic value of a Trypanosoma cruzi (Chagas disease) vaccine in Latin America. PLoS Negl Trop Dis, 2010. 4(1), e916.</li> <li>85. Custer, B., et al., The cost-effectiveness of pathogen reduction technology as assessed using a multiple risk reduction model. Transfusion, 2010. 50(11), 2461-2473.</li> <li>86. Agapova, M., et al., Cost-effectiveness of screening the US blood supply for Trypanosoma cruzi. Transfusion, 2010. 50(11), 2461-2473.</li> <li>87. Vazquez-Prokopec, G.M., et al., Cost-effectiveness of chagas disease vector control strategies in Northwestern Argentina. PLoS Negl Trop Dis, 2009. 3(1), e363.</li> <li>88. Wilson, L.S., et al., Cost-effectiveness of Chagas disease interventions in latin america and the Caribbean: Markov models. Am J Trop Med Hyg, 2005. 73(5), 901-10.</li> <li>89. Wilson, L.S., et al., Cost-effectiveness of Chagas disease control. Mem Inst Oswaldo Cruz, 1991. 86(3), 285-95.</li> <li>91. Barbu, C., et al., Evaluation of spatially targeted strategies to control non-domiciliated Triatoma dimidiata vector of Chagas disease. PLoS Negl Trop Dis, 2011. 5(5), e1045.</li> <li>92. Castillo-Riquelme, M., et al., Modelling geographic variation in the cost-effectiveness of co</li></ol>		79.	
<ul> <li>effective option? Trans R Soc Trop Med Hyg, 1995. 89(1), 16-20.</li> <li>Guyatt, H.L., et al., A population dynamic approach to the cost-effectiveness analysis of mass anthelminit treatment: effects of treatment frequency on Ascaris infection. Trans R Soc Trop Med Hyg, 1993. 87(5), 570-5.</li> <li>Lee, B.Y., et al., Modeling the economic value of a Chagas' disease therapeutic vaccine. Hum Vaccin &amp; Immunother, 2012. 8(9), 1293-1301.</li> <li>Sicuri, E., et al., Economic evaluation of Chagas disease screening of pregnant Latin American women and of their infants in a non endemic area. Acta Trop, 2011. 118(2), 110-7.</li> <li>Lee, B.Y., et al., The potential economic value of a Trypanosoma cruzi (Chagas disease) vaccine in Latin America. PLoS Negl Trop Dis, 2010. 4(12), e916.</li> <li>Custer, B., et al., The cost-effectiveness of pathogen reduction technology as assessed using a multiple risk reduction model. Transfusion, 2010. 50(11), 2461-2473.</li> <li>Agapova, M., et al., Cost-effectiveness of screening the US blood supply for Trypanosoma cruzi. Transfusion, 2010. 50(10), 2220-32.</li> <li>Vazquez-Prokopec, G.M., et al., Cost-effectiveness of chagas disease vector control strategies in Northwestern Argentina. PLoS Negl Trop Dis, 2009. 3(1), e363.</li> <li>Wilson, L.S., et al., Cost-effectiveness of Chagas disease interventions in latin america and the Caribbean: Markov models. Am J Trop Med Hyg, 2005. 73(5), 901-10.</li> <li>Schofield, C.J. and Dias, J.C., A cost-benefit analysis of Chagas disease control. Mem Inst Oswaldo Cruz, 1991. 86(3), 285-95.</li> <li>Barbu, C., et al., Evaluation of spatially targeted strategies to control non-domiciliated Triatoma dimidiata vector of Chagas disease. PLoS Negl Trop Dis, 2011. 5(5), e1045.</li> <li>Castillo-Riquelme, M., et al., Modelling geographic variation in the cost-effectiveness of control policies for infectious vector diseases: the example of Chagas disease. J Health Econ, 2008. 27(2), 405-26.</li> <li>Basombrio, M.A., et al., A cost-benefit analysis of</li></ul>		00	
<ol> <li>81. Guyatt, H.L., et al., A population dynamic approach to the cost-effectiveness analysis of mass anthelmintic treatment: effects of treatment frequency on Ascaris infection. Trans R Soc Trop Mcd Hyg, 1993. 87(5), 570-5.</li> <li>82. Lee, B.Y., et al., Modeling the economic value of a Chagas' disease therapeutic vaccine. Hum Vaccin &amp; Immunother, 2012. 8(9), 1293-1301.</li> <li>83. Sicuri, E., et al., Economic evaluation of Chagas disease screening of pregnant Latin American women and of their infants in a non endemic area. Acta Trop, 2011. 118(2), 110-7.</li> <li>84. Lee, B.Y., et al., The potential economic value of a Trypanosoma cruzi (Chagas disease) vaccine in Latin America. PLoS Negl Trop Dis, 2010. 4(12), e916.</li> <li>85. Custer, B., et al., The cost-effectiveness of pathogen reduction technology as assessed using a multiple risk reduction model. Transfusion, 2010. 50(11), 2461-2473.</li> <li>86. Agapova, M., et al., Cost-effectiveness of screening the US blood supply for Trypanosoma cruzi. Transfusion, 2010. 50(10), 2220-32.</li> <li>87. Vazquez-Prokopec, G.M., et al., Cost-effectiveness of chagas disease vector control strategies in Northwestern Argentina. PLoS Negl Trop Dis, 2009. 3(1), e363.</li> <li>88. Wilson, L.S., et al., Cost-effectiveness of Internetiation methods for ELISA serology testing of Trypanosoma cruzi in California blood banks. Am J of Trop Med and Hyg, 2008. 79(1), 53-68.</li> <li>89. Wilson, L.S., et al., Cost-effectiveness of Chagas disease interventions in latin america and the Caribbean: Markov models. Am J Trop Med Hyg, 2005. 73(5), 901-10.</li> <li>90. Schofield, C.J. and Dias, J.C., A cost-benefit analysis of Chagas disease control. Mem Inst Oswaldo Cruz, 1991. 86(3), 285-95.</li> <li>91. Barbu, C., et al., Evaluation of spatially targeted strategies to control non-domiciliated Triatoma dimidiata vector of Chagas disease: PLoS Negl Trop Dis, 2011. 5(5), e1045.</li> <li>92. Castillo-Riquelme, M., et al., Modelling geogr</li></ol>		80.	• • • • •
<ul> <li>anthelmintic treatment: effects of treatment frequency on Ascaris infection. Trans R Soc Trop Med Hyg, 1993. 87(5), 570-5.</li> <li>Lee, B. Y., et al., Modeling the economic value of a Chagas' disease therapeutic vaccine. Hum Vaccin &amp; Immunother, 2012. 8(9), 1293-1301.</li> <li>Sicuri, E., et al., Economic evaluation of Chagas disease screening of pregnant Latin American women and of their infants in a non endemic area. Acta Trop, 2011. 118(2), 110-7.</li> <li>Lee, B. Y., et al., The potential economic value of a Trypanosoma cruzi (Chagas disease) vaccine in Latin America. PLoS Negl Trop Dis, 2010. 4(12), e916.</li> <li>Custer, B., et al., The cost-effectiveness of pathogen reduction technology as assessed using a multiple risk reduction model. Transfusion, 2010. 50(11), 2461-2473.</li> <li>Agapova, M., et al., Cost-effectiveness of screening the US blood supply for Trypanosoma cruzi. Transfusion, 2010. 50(10), 2220-32.</li> <li>Vazquez-Prokopec, G.M., et al., Cost-effectiveness of chagas disease vector control strategies in Northwestern Argentina. PLoS Negl Trop Dis, 2009. 3(1), e363.</li> <li>Wilson, L.S., et al., Cost-effectiveness of implementation methods for ELISA serology testing of Trypanosoma cruzi in California blood banks. Am J of Trop Med and Hyg, 2008. 79(1), 53-68.</li> <li>Wilson, L.S., et al., Cost-effectiveness of Chagas disease interventions in latin america and the Caribbean: Markov models. Am J Trop Med Hyg, 2005. 73(5), 901-10.</li> <li>Schofield, C.J. and Dias, J.C., A cost-benefit analysis of Chagas disease control. Mem Inst Oswaldo Cruz, 1991. 86(3), 285-95.</li> <li>Barbu, C., et al., Evaluation of spatially targeted strategies to control non-domiciliated Triatoma dimidiata vector of Chagas disease: PLoS Negl Trop Dis, 2011. 5(5), e1045.</li> <li>Castillo-Riquelme, M., et al., Modelling geographic variation in the cost-effectiveness of control policies for infectious vector diseases: the example of Chagas disease. J Health Econ, 2008. 27(2), 405-26.</li> <li>Basombrio, M.A., et</li></ul>		0.1	
<ul> <li>Med Hyg, 1993. 87(5), 570-5.</li> <li>Lee, B.Y., et al., Modeling the economic value of a Chagas' disease therapeutic vaccine. Hum Vaccin &amp; Immunother, 2012. 8(9), 1293-1301.</li> <li>Sicuri, E., et al., Economic evaluation of Chagas disease screening of pregnant Latin American women and of their infants in a non endemic area. Acta Trop, 2011. 118(2), 110-7.</li> <li>Lee, B.Y., et al., The potential economic value of a Trypanosoma cruzi (Chagas disease) vaccine in Latin America. PLoS Negl Trop Dis, 2010. 4(12), e916.</li> <li>Custer, B., et al., The cost-effectiveness of pathogen reduction technology as assessed using a multiple risk reduction model. Transfusion, 2010. 50(11), 2461-2473.</li> <li>Agapova, M., et al., Cost-effectiveness of screening the US blood supply for Trypanosoma cruzi. Transfusion, 2010. 50(10), 2220-32.</li> <li>Vazquez-Prokopec, G.M., et al., Cost-effectiveness of chagas disease vector control strategies in Northwestern Argentina. PLoS Negl Trop Dis, 2009. 3(1), e363.</li> <li>Wilson, L.S., et al., Cost-effectiveness of Chagas disease vector control strategies of Trypanosoma cruzi in California blood banks. Am J of Trop Med and Hyg, 2008. 79(1), 53-68.</li> <li>Wilson, L.S., et al., Cost-effectiveness of Chagas disease interventions in latin america and the Caribbean: Markov models. Am J Trop Med Hyg, 2005. 73(5), 901-10.</li> <li>Schofield, C.J. and Dias, J.C., A cost-benefit analysis of Chagas disease control. Mem Inst Oswaldo Cruz, 1991. 86(3), 285-95.</li> <li>Barbu, C., et al., Evaluation of spatially targeted strategies to control non-domiciliated Triatoma dimidiata vector of Chagas disease: PLoS Negl Trop Dis, 2011. 5(5), e1045.</li> <li>Castillo-Riquelme, M., et al., Modelling geographic variation in the cost-effectiveness of control policies for infectious vector diseases: the example of Chagas disease. J Health Econ, 2008. 27(2), 405-26.</li> <li>Basombrio, M.A., et al., A cost-benefit analysis of Chagas disease control in north-western</li> </ul>		81.	
<ol> <li>Lee, B.Y., et al., Modeling the economic value of a Chagas' disease therapeutic vaccine. Hum Vaccin &amp; Immunother, 2012. 8(9), 1293-1301.</li> <li>Sicuri, E., et al., Economic evaluation of Chagas disease screening of pregnant Latin American women and of their infants in a non endemic area. Acta Trop, 2011. 118(2), 110-7.</li> <li>Lee, B.Y., et al., The potential economic value of a Trypanosoma cruzi (Chagas disease) vaccine in Latin America. PLoS Negl Trop Dis, 2010. 4(12), e916.</li> <li>Custer, B., et al., The cost-effectiveness of pathogen reduction technology as assessed using a multiple risk reduction model. Transfusion, 2010. 50(11), 2461-2473.</li> <li>Agapova, M., et al., Cost-effectiveness of screening the US blood supply for Trypanosoma cruzi. Transfusion, 2010. 50(10), 2220-32.</li> <li>Vazquez-Prokopec, G.M., et al., Cost-effectiveness of chagas disease vector control strategies in Northwestern Argentina. PLoS Negl Trop Dis, 2009. 3(1), e363.</li> <li>Wilson, L.S., et al., Cost-effectiveness of Chagas disease vector control strategies of Trypanosoma cruzi in California blood banks. Am J of Trop Med and Hyg, 2008. 79(1), 53-68.</li> <li>Wilson, L.S., et al., Cost-effectiveness of Chagas disease interventions in latin america and the Caribbean: Markov models. Am J Trop Med Hyg, 2005. 73(5), 901-10.</li> <li>Schofteld, C.J. and Dias, J.C., A cost-benefit analysis of Chagas disease control. Mem Inst Oswaldo Cruz, 1991. 86(3), 285-95.</li> <li>Barbu, C., et al., Evaluation of spatially targeted strategies to control non-domiciliated Triatoma dimidiata vector of Chagas disease. PLoS Negl Trop Dis, 2011. 5(5), e1045.</li> <li>Castillo-Riquelme, M., et al., Modelling geographic variation in the cost-effectiveness of control policies for infectious vector diseases: the example of Chagas disease. J Health Econ, 2008. 27(2), 405-26.</li> <li>Basombrio, M.A., et al., A cost-benefit analysis of Chagas disease control in north-western</li> </ol>			
<ul> <li>Vaccin &amp; Immunother, 2012. 8(9), 1293-1301.</li> <li>Sicuri, E., et al., Economic evaluation of Chagas disease screening of pregnant Latin American women and of their infants in a non endemic area. Acta Trop, 2011. 118(2), 110-7.</li> <li>Lee, B.Y., et al., The potential economic value of a Trypanosoma cruzi (Chagas disease) vaccine in Latin America. PLoS Negl Trop Dis, 2010. 4(12), e916.</li> <li>Custer, B., et al., The cost-effectiveness of pathogen reduction technology as assessed using a multiple risk reduction model. Transfusion, 2010. 50(11), 2461-2473.</li> <li>Agapova, M., et al., Cost-effectiveness of screening the US blood supply for Trypanosoma cruzi. Transfusion, 2010. 50(10), 2220-32.</li> <li>Vazquez-Prokopec, G.M., et al., Cost-effectiveness of chagas disease vector control strategies in Northwestern Argentina. PLoS Negl Trop Dis, 2009. 3(1), e363.</li> <li>Wilson, L.S., et al., Cost-effectiveness of implementation methods for ELISA serology testing of Trypanosoma cruzi in California blood banks. Am J of Trop Med and Hyg, 2008. 79(1), 53-68.</li> <li>Wilson, L.S., et al., Cost-effectiveness of Chagas disease interventions in latin america and the Caribbean: Markov models. Am J Trop Med Hyg, 2005. 73(5), 901-10.</li> <li>Schofield, C.J. and Dias, J.C., A cost-benefit analysis of Chagas disease control. Mem Inst Oswaldo Cruz, 1991. 86(3), 285-95.</li> <li>Barbu, C., et al., Evaluation of spatially targeted strategies to control non-domiciliated Triatoma dimidiata vector of Chagas disease: PLoS Negl Trop Dis, 2011. 5(5), e1045.</li> <li>Castillo-Riquelme, M., et al., Modelling geographic variation in the cost-effectiveness of control policies for infectious vector diseases: the example of Chagas disease. J Health Econ, 2008. 27(2), 405-26.</li> <li>Basombrio, M.A., et al., A cost-benefit analysis of Chagas disease control in north-western</li> </ul>		00	
<ol> <li>83. Sicuri, E., et al., <i>Economic evaluation of Chagas disease screening of pregnant Latin</i> <i>American women and of their infants in a non endemic area.</i> Acta Trop, 2011. 118(2), 110-7.</li> <li>84. Lee, B.Y., et al., <i>The potential economic value of a Trypanosoma cruzi (Chagas disease)</i> <i>vaccine in Latin America.</i> PLoS Negl Trop Dis, 2010. 4(12), e916.</li> <li>85. Custer, B., et al., <i>The cost-effectiveness of pathogen reduction technology as assessed using a</i> <i>multiple risk reduction model.</i> Transfusion, 2010. 50(11), 2461-2473.</li> <li>86. Agapova, M., et al., <i>Cost-effectiveness of screening the US blood supply for Trypanosoma</i> <i>cruzi.</i> Transfusion, 2010. 50(10), 2220-32.</li> <li>87. Vazquez-Prokopec, G.M., et al., <i>Cost-effectiveness of chagas disease vector control strategies</i> <i>in Northwestern Argentina.</i> PLoS Negl Trop Dis, 2009. 3(1), e363.</li> <li>88. Wilson, L.S., et al., <i>Cost-effectiveness of implementation methods for ELISA serology testing</i> <i>of Trypanosoma cruzi in California blood banks.</i> Am J of Trop Med and Hyg, 2008. 79(1), 53-68.</li> <li>89. Wilson, L.S., et al., <i>Cost-effectiveness of Chagas disease interventions in latin america and</i> <i>the Caribbean: Markov models.</i> Am J Trop Med Hyg, 2005. 73(5), 901-10.</li> <li>90. Schofield, C.J. and Dias, J.C., <i>A cost-benefit analysis of Chagas disease control.</i> Mem Inst Oswaldo Cruz, 1991. 86(3), 285-95.</li> <li>91. Barbu, C., et al., <i>Evaluation of spatially targeted strategies to control non-domiciliated</i> <i>Triatoma dimidiata vector of Chagas disease.</i> PLoS Negl Trop Dis, 2011. 5(5), e1045.</li> <li>92. Castillo-Riquelme, M., et al., <i>Modelling geographic variation in the cost-effectiveness of</i> <i>control policies for infectious vector diseases: the example of Chagas disease.</i> J Health Econ, 2008. 27(2), 405-26.</li> <li>93. Basombrio, M.A., et al., <i>A cost-benefit analysis of Chagas disease control in north-western</i></li> </ol>		82.	
<ul> <li>American women and of their infants in a non endemic area. Acta Trop, 2011. 118(2), 110-7.</li> <li>Ke, B.Y., et al., <i>The potential economic value of a Trypanosoma cruzi (Chagas disease)</i></li> <li>vaccine in Latin America. PLoS Negl Trop Dis, 2010. 4(12), e916.</li> <li>Custer, B., et al., <i>The cost-effectiveness of pathogen reduction technology as assessed using a</i></li> <li><i>multiple risk reduction model</i>. Transfusion, 2010. 50(11), 2461-2473.</li> <li>Agapova, M., et al., <i>Cost-effectiveness of screening the US blood supply for Trypanosoma</i></li> <li><i>cruzi</i>. Transfusion, 2010. 50(10), 2220-32.</li> <li>Vazquez-Prokopec, G.M., et al., <i>Cost-effectiveness of chagas disease vector control strategies</i></li> <li><i>in Northwestern Argentina</i>. PLoS Negl Trop Dis, 2009. 3(1), e363.</li> <li>Wilson, L.S., et al., <i>Cost-effectiveness of Chagas disease interventions in latin america and</i></li> <li><i>the Caribbean: Markov models</i>. Am J Trop Med Hyg, 2005. 73(5), 901-10.</li> <li>Schofield, C.J. and Dias, J.C., <i>A cost-benefit analysis of Chagas disease control</i>. Mem Inst</li> <li>Oswaldo Cruz, 1991. 86(3), 285-95.</li> <li>Barbu, C., et al., <i>Evaluation of spatially targeted strategies to control non-domiciliated</i></li> <li><i>Triatoma dimidiata vector of Chagas disease</i>. PLoS Negl Trop Dis, 2011. 5(5), e1045.</li> <li>Castillo-Riquelme, M., et al., <i>Modelling geographic variation in the cost-effectiveness of</i></li> <li>Sasombrio, M.A., et al., <i>A cost-benefit analysis of Chagas disease</i>. J Health Econ, 2008. 27(2), 405-26.</li> </ul>		02	
<ol> <li>84. Lee, B.Y., et al., <i>The potential economic value of a Trypanosoma cruzi (Chagas disease)</i></li> <li>vaccine in Latin America. PLoS Negl Trop Dis, 2010. 4(12), e916.</li> <li>85. Custer, B., et al., <i>The cost-effectiveness of pathogen reduction technology as assessed using a</i></li> <li><i>multiple risk reduction model</i>. Transfusion, 2010. 50(11), 2461-2473.</li> <li>86. Agapova, M., et al., <i>Cost-effectiveness of screening the US blood supply for Trypanosoma</i></li> <li><i>cruzi</i>. Transfusion, 2010. 50(10), 2220-32.</li> <li>87. Vazquez-Prokopec, G.M., et al., <i>Cost-effectiveness of chagas disease vector control strategies</i></li> <li><i>in Northwestern Argentina</i>. PLoS Negl Trop Dis, 2009. 3(1), e363.</li> <li>88. Wilson, L.S., et al., <i>Cost-effectiveness of implementation methods for ELISA serology testing</i></li> <li><i>of Trypanosoma cruzi in California blood banks</i>. Am J of Trop Med and Hyg, 2008. 79(1),</li> <li>53-68.</li> <li>89. Wilson, L.S., et al., <i>Cost-effectiveness of Chagas disease interventions in latin america and</i></li> <li><i>the Caribbean: Markov models</i>. Am J Trop Med Hyg, 2005. 73(5), 901-10.</li> <li>90. Schofield, C.J. and Dias, J.C., <i>A cost-benefit analysis of Chagas disease control</i>. Mem Inst</li> <li>Oswaldo Cruz, 1991. 86(3), 285-95.</li> <li>91. Barbu, C., et al., <i>Evaluation of spatially targeted strategies to control non-domiciliated</i></li> <li><i>Triatoma dimidiata vector of Chagas disease</i>. PLoS Negl Trop Dis, 2011. 5(5), e1045.</li> <li>92. Castillo-Riquelme, M., et al., <i>Modelling geographic variation in the cost-effectiveness of</i></li> <li>27(2), 405-26.</li> <li>93. Basombrio, M.A., et al., <i>A cost-benefit analysis of Chagas disease control in north-western</i></li> </ol>		83.	
<ul> <li>vaccine in Latin America. PLoS Negl Trop Dis, 2010. 4(12), e916.</li> <li>Custer, B., et al., The cost-effectiveness of pathogen reduction technology as assessed using a multiple risk reduction model. Transfusion, 2010. 50(11), 2461-2473.</li> <li>Agapova, M., et al., Cost-effectiveness of screening the US blood supply for Trypanosoma cruzi. Transfusion, 2010. 50(10), 2220-32.</li> <li>Vazquez-Prokopec, G.M., et al., Cost-effectiveness of chagas disease vector control strategies in Northwestern Argentina. PLoS Negl Trop Dis, 2009. 3(1), e363.</li> <li>Wilson, L.S., et al., Cost-effectiveness of implementation methods for ELISA serology testing of Trypanosoma cruzi in California blood banks. Am J of Trop Med and Hyg, 2008. 79(1), 53-68.</li> <li>Wilson, L.S., et al., Cost-effectiveness of Chagas disease interventions in latin america and the Caribbean: Markov models. Am J Trop Med Hyg, 2005. 73(5), 901-10.</li> <li>Schofield, C.J. and Dias, J.C., A cost-benefit analysis of Chagas disease control. Mem Inst Oswaldo Cruz, 1991. 86(3), 285-95.</li> <li>Barbu, C., et al., Evaluation of spatially targeted strategies to control non-domiciliated Triatoma dimidiata vector of Chagas disease. PLoS Negl Trop Dis, 2011. 5(5), e1045.</li> <li>Castillo-Riquelme, M., et al., Modelling geographic variation in the cost-effectiveness of control policies for infectious vector diseases: the example of Chagas disease. J Health Econ, 2008. 27(2), 405-26.</li> <li>Basombrio, M.A., et al., A cost-benefit analysis of Chagas disease control in north-western</li> </ul>		04	
<ol> <li>85. Custer, B., et al., <i>The cost-effectiveness of pathogen reduction technology as assessed using a</i> <i>multiple risk reduction model.</i> Transfusion, 2010. <b>50</b>(11), 2461-2473.</li> <li>86. Agapova, M., et al., <i>Cost-effectiveness of screening the US blood supply for Trypanosoma</i> <i>cruzi.</i> Transfusion, 2010. <b>50</b>(10), 2220-32.</li> <li>87. Vazquez-Prokopec, G.M., et al., <i>Cost-effectiveness of chagas disease vector control strategies</i> <i>in Northwestern Argentina.</i> PLoS Negl Trop Dis, 2009. <b>3</b>(1), e363.</li> <li>88. Wilson, L.S., et al., <i>Cost-effectiveness of implementation methods for ELISA serology testing</i> <i>of Trypanosoma cruzi in California blood banks.</i> Am J of Trop Med and Hyg, 2008. <b>79</b>(1), 53-68.</li> <li>89. Wilson, L.S., et al., <i>Cost-effectiveness of Chagas disease interventions in latin america and</i> <i>the Caribbean: Markov models.</i> Am J Trop Med Hyg, 2005. <b>73</b>(5), 901-10.</li> <li>90. Schofield, C.J. and Dias, J.C., <i>A cost-benefit analysis of Chagas disease control.</i> Mem Inst Oswaldo Cruz, 1991. <b>86</b>(3), 285-95.</li> <li>91. Barbu, C., et al., <i>Evaluation of spatially targeted strategies to control non-domiciliated</i> <i>Triatoma dimidiata vector of Chagas disease.</i> PLoS Negl Trop Dis, 2011. <b>5</b>(5), e1045.</li> <li>92. Castillo-Riquelme, M., et al., <i>Modelling geographic variation in the cost-effectiveness of</i> <i>control policies for infectious vector diseases: the example of Chagas disease.</i> J Health Econ, 2008. <b>27</b>(2), 405-26.</li> <li>93. Basombrio, M.A., et al., <i>A cost-benefit analysis of Chagas disease control in north-western</i></li> </ol>		84.	
<ul> <li>multiple risk reduction model. Transfusion, 2010. 50(11), 2461-2473.</li> <li>Agapova, M., et al., <i>Cost-effectiveness of screening the US blood supply for Trypanosoma cruzi</i>. Transfusion, 2010. 50(10), 2220-32.</li> <li>Vazquez-Prokopec, G.M., et al., <i>Cost-effectiveness of chagas disease vector control strategies in Northwestern Argentina</i>. PLoS Negl Trop Dis, 2009. 3(1), e363.</li> <li>Wilson, L.S., et al., <i>Cost-effectiveness of implementation methods for ELISA serology testing of Trypanosoma cruzi in California blood banks</i>. Am J of Trop Med and Hyg, 2008. 79(1), 53-68.</li> <li>Wilson, L.S., et al., <i>Cost-effectiveness of Chagas disease interventions in latin america and the Caribbean: Markov models</i>. Am J Trop Med Hyg, 2005. 73(5), 901-10.</li> <li>Schofield, C.J. and Dias, J.C., <i>A cost-benefit analysis of Chagas disease control</i>. Mem Inst Oswaldo Cruz, 1991. 86(3), 285-95.</li> <li>Barbu, C., et al., <i>Evaluation of spatially targeted strategies to control non-domiciliated Triatoma dimidiata vector of Chagas disease</i>. PLoS Negl Trop Dis, 2011. 5(5), e1045.</li> <li>Castillo-Riquelme, M., et al., <i>Modelling geographic variation in the cost-effectiveness of control policies for infectious vector diseases: the example of Chagas disease</i>. J Health Econ, 2008. 27(2), 405-26.</li> <li>Basombrio, M.A., et al., <i>A cost-benefit analysis of Chagas disease control in north-western</i></li> </ul>		05	
<ol> <li>86. Agapova, M., et al., Cost-effectiveness of screening the US blood supply for Trypanosoma cruzi. Transfusion, 2010. 50(10), 2220-32.</li> <li>87. Vazquez-Prokopec, G.M., et al., Cost-effectiveness of chagas disease vector control strategies in Northwestern Argentina. PLoS Negl Trop Dis, 2009. 3(1), e363.</li> <li>88. Wilson, L.S., et al., Cost-effectiveness of implementation methods for ELISA serology testing of Trypanosoma cruzi in California blood banks. Am J of Trop Med and Hyg, 2008. 79(1), 53-68.</li> <li>89. Wilson, L.S., et al., Cost-effectiveness of Chagas disease interventions in latin america and the Caribbean: Markov models. Am J Trop Med Hyg, 2005. 73(5), 901-10.</li> <li>90. Schofield, C.J. and Dias, J.C., A cost-benefit analysis of Chagas disease control. Mem Inst Oswaldo Cruz, 1991. 86(3), 285-95.</li> <li>91. Barbu, C., et al., Evaluation of spatially targeted strategies to control non-domiciliated Triatoma dimidiata vector of Chagas disease. PLoS Negl Trop Dis, 2011. 5(5), e1045.</li> <li>92. Castillo-Riquelme, M., et al., Modelling geographic variation in the cost-effectiveness of control policies for infectious vector diseases: the example of Chagas disease. J Health Econ, 2008. 27(2), 405-26.</li> <li>93. Basombrio, M.A., et al., A cost-benefit analysis of Chagas disease control in north-western</li> </ol>		85.	
<ul> <li><i>cruzi.</i> Transfusion, 2010. 50(10), 2220-32.</li> <li>87. Vazquez-Prokopec, G.M., et al., <i>Cost-effectiveness of chagas disease vector control strategies</i> <i>in Northwestern Argentina.</i> PLoS Negl Trop Dis, 2009. 3(1), e363.</li> <li>88. Wilson, L.S., et al., <i>Cost-effectiveness of implementation methods for ELISA serology testing</i> <i>of Trypanosoma cruzi in California blood banks.</i> Am J of Trop Med and Hyg, 2008. 79(1), 53-68.</li> <li>89. Wilson, L.S., et al., <i>Cost-effectiveness of Chagas disease interventions in latin america and</i> <i>the Caribbean: Markov models.</i> Am J Trop Med Hyg, 2005. 73(5), 901-10.</li> <li>90. Schofield, C.J. and Dias, J.C., <i>A cost-benefit analysis of Chagas disease control.</i> Mem Inst Oswaldo Cruz, 1991. 86(3), 285-95.</li> <li>91. Barbu, C., et al., <i>Evaluation of spatially targeted strategies to control non-domiciliated</i> <i>Triatoma dimidiata vector of Chagas disease.</i> PLoS Negl Trop Dis, 2011. 5(5), e1045.</li> <li>92. Castillo-Riquelme, M., et al., <i>Modelling geographic variation in the cost-effectiveness of</i> <i>control policies for infectious vector diseases: the example of Chagas disease.</i> J Health Econ, 2008. 27(2), 405-26.</li> <li>93. Basombrio, M.A., et al., <i>A cost-benefit analysis of Chagas disease control in north-western</i></li> </ul>		96	•
<ol> <li>87. Vazquez-Prokopec, G.M., et al., Cost-effectiveness of chagas disease vector control strategies in Northwestern Argentina. PLoS Negl Trop Dis, 2009. 3(1), e363.</li> <li>88. Wilson, L.S., et al., Cost-effectiveness of implementation methods for ELISA serology testing of Trypanosoma cruzi in California blood banks. Am J of Trop Med and Hyg, 2008. 79(1), 53-68.</li> <li>89. Wilson, L.S., et al., Cost-effectiveness of Chagas disease interventions in latin america and the Caribbean: Markov models. Am J Trop Med Hyg, 2005. 73(5), 901-10.</li> <li>90. Schofield, C.J. and Dias, J.C., A cost-benefit analysis of Chagas disease control. Mem Inst Oswaldo Cruz, 1991. 86(3), 285-95.</li> <li>91. Barbu, C., et al., Evaluation of spatially targeted strategies to control non-domiciliated Triatoma dimidiata vector of Chagas disease. PLoS Negl Trop Dis, 2011. 5(5), e1045.</li> <li>92. Castillo-Riquelme, M., et al., Modelling geographic variation in the cost-effectiveness of control policies for infectious vector diseases: the example of Chagas disease. J Health Econ, 2008. 27(2), 405-26.</li> <li>93. Basombrio, M.A., et al., A cost-benefit analysis of Chagas disease control in north-western</li> </ol>		80.	
<ul> <li><i>in Northwestern Argentina.</i> PLoS Negl Trop Dis, 2009. 3(1), e363.</li> <li>Wilson, L.S., et al., <i>Cost-effectiveness of implementation methods for ELISA serology testing</i> <i>of Trypanosoma cruzi in California blood banks.</i> Am J of Trop Med and Hyg, 2008. 79(1), 53-68.</li> <li>Wilson, L.S., et al., <i>Cost-effectiveness of Chagas disease interventions in latin america and</i> <i>the Caribbean: Markov models.</i> Am J Trop Med Hyg, 2005. 73(5), 901-10.</li> <li>Schofield, C.J. and Dias, J.C., <i>A cost-benefit analysis of Chagas disease control.</i> Mem Inst Oswaldo Cruz, 1991. 86(3), 285-95.</li> <li>Barbu, C., et al., <i>Evaluation of spatially targeted strategies to control non-domiciliated</i> <i>Triatoma dimidiata vector of Chagas disease.</i> PLoS Negl Trop Dis, 2011. 5(5), e1045.</li> <li>Castillo-Riquelme, M., et al., <i>Modelling geographic variation in the cost-effectiveness of</i> <i>control policies for infectious vector diseases: the example of Chagas disease.</i> J Health Econ, 2008. 27(2), 405-26.</li> <li>Basombrio, M.A., et al., <i>A cost-benefit analysis of Chagas disease control in north-western</i></li> </ul>		07	
<ul> <li>Wilson, L.S., et al., Cost-effectiveness of implementation methods for ELISA serology testing of Trypanosoma cruzi in California blood banks. Am J of Trop Med and Hyg, 2008. 79(1), 53-68.</li> <li>Wilson, L.S., et al., Cost-effectiveness of Chagas disease interventions in latin america and the Caribbean: Markov models. Am J Trop Med Hyg, 2005. 73(5), 901-10.</li> <li>Schofield, C.J. and Dias, J.C., A cost-benefit analysis of Chagas disease control. Mem Inst Oswaldo Cruz, 1991. 86(3), 285-95.</li> <li>Barbu, C., et al., Evaluation of spatially targeted strategies to control non-domiciliated Triatoma dimidiata vector of Chagas disease. PLoS Negl Trop Dis, 2011. 5(5), e1045.</li> <li>Castillo-Riquelme, M., et al., Modelling geographic variation in the cost-effectiveness of control policies for infectious vector diseases: the example of Chagas disease. J Health Econ, 2008. 27(2), 405-26.</li> <li>Basombrio, M.A., et al., A cost-benefit analysis of Chagas disease control in north-western</li> </ul>		07.	
<ul> <li><i>of Trypanosoma cruzi in California blood banks.</i> Am J of Trop Med and Hyg, 2008. <b>79</b>(1),</li> <li>53-68.</li> <li>Wilson, L.S., et al., <i>Cost-effectiveness of Chagas disease interventions in latin america and</i></li> <li><i>the Caribbean: Markov models.</i> Am J Trop Med Hyg, 2005. <b>73</b>(5), 901-10.</li> <li>Schofield, C.J. and Dias, J.C., <i>A cost-benefit analysis of Chagas disease control.</i> Mem Inst</li> <li>Oswaldo Cruz, 1991. <b>86</b>(3), 285-95.</li> <li>Barbu, C., et al., <i>Evaluation of spatially targeted strategies to control non-domiciliated</i></li> <li><i>Triatoma dimidiata vector of Chagas disease.</i> PLoS Negl Trop Dis, 2011. <b>5</b>(5), e1045.</li> <li>Castillo-Riquelme, M., et al., <i>Modelling geographic variation in the cost-effectiveness of</i></li> <li><i>control policies for infectious vector diseases: the example of Chagas disease.</i> J Health Econ, 2008. <b>27</b>(2), 405-26.</li> <li>Basombrio, M.A., et al., <i>A cost-benefit analysis of Chagas disease control in north-western</i></li> </ul>		99	
<ul> <li>53-68.</li> <li>Wilson, L.S., et al., <i>Cost-effectiveness of Chagas disease interventions in latin america and</i></li> <li><i>the Caribbean: Markov models.</i> Am J Trop Med Hyg, 2005. <b>73</b>(5), 901-10.</li> <li>Schofield, C.J. and Dias, J.C., <i>A cost-benefit analysis of Chagas disease control.</i> Mem Inst</li> <li>Oswaldo Cruz, 1991. <b>86</b>(3), 285-95.</li> <li>Barbu, C., et al., <i>Evaluation of spatially targeted strategies to control non-domiciliated</i></li> <li><i>Triatoma dimidiata vector of Chagas disease.</i> PLoS Negl Trop Dis, 2011. <b>5</b>(5), e1045.</li> <li>Castillo-Riquelme, M., et al., <i>Modelling geographic variation in the cost-effectiveness of</i></li> <li><i>control policies for infectious vector diseases: the example of Chagas disease.</i> J Health Econ, 2008. <b>27</b>(2), 405-26.</li> <li>Basombrio, M.A., et al., <i>A cost-benefit analysis of Chagas disease control in north-western</i></li> </ul>		00.	
<ul> <li>Wilson, L.S., et al., <i>Cost-effectiveness of Chagas disease interventions in latin america and</i> <i>the Caribbean: Markov models.</i> Am J Trop Med Hyg, 2005. <b>73</b>(5), 901-10.</li> <li>Schofield, C.J. and Dias, J.C., <i>A cost-benefit analysis of Chagas disease control.</i> Mem Inst Oswaldo Cruz, 1991. <b>86</b>(3), 285-95.</li> <li>Barbu, C., et al., <i>Evaluation of spatially targeted strategies to control non-domiciliated</i> <i>Triatoma dimidiata vector of Chagas disease.</i> PLoS Negl Trop Dis, 2011. <b>5</b>(5), e1045.</li> <li>Castillo-Riquelme, M., et al., <i>Modelling geographic variation in the cost-effectiveness of</i> <i>control policies for infectious vector diseases: the example of Chagas disease.</i> J Health Econ, 2008. <b>27</b>(2), 405-26.</li> <li>Basombrio, M.A., et al., <i>A cost-benefit analysis of Chagas disease control in north-western</i></li> </ul>			
<ul> <li>the Caribbean: Markov models. Am J Trop Med Hyg, 2005. 73(5), 901-10.</li> <li>Schofield, C.J. and Dias, J.C., A cost-benefit analysis of Chagas disease control. Mem Inst Oswaldo Cruz, 1991. 86(3), 285-95.</li> <li>Barbu, C., et al., Evaluation of spatially targeted strategies to control non-domiciliated Triatoma dimidiata vector of Chagas disease. PLoS Negl Trop Dis, 2011. 5(5), e1045.</li> <li>Castillo-Riquelme, M., et al., Modelling geographic variation in the cost-effectiveness of control policies for infectious vector diseases: the example of Chagas disease. J Health Econ, 2008. 27(2), 405-26.</li> <li>Basombrio, M.A., et al., A cost-benefit analysis of Chagas disease control in north-western</li> </ul>		80	
<ul> <li>Schofield, C.J. and Dias, J.C., A cost-benefit analysis of Chagas disease control. Mem Inst Oswaldo Cruz, 1991. 86(3), 285-95.</li> <li>Barbu, C., et al., Evaluation of spatially targeted strategies to control non-domiciliated Triatoma dimidiata vector of Chagas disease. PLoS Negl Trop Dis, 2011. 5(5), e1045.</li> <li>Castillo-Riquelme, M., et al., Modelling geographic variation in the cost-effectiveness of control policies for infectious vector diseases: the example of Chagas disease. J Health Econ, 2008. 27(2), 405-26.</li> <li>Basombrio, M.A., et al., A cost-benefit analysis of Chagas disease control in north-western</li> </ul>		07.	
<ul> <li>48 Oswaldo Cruz, 1991. 86(3), 285-95.</li> <li>49 91. Barbu, C., et al., Evaluation of spatially targeted strategies to control non-domiciliated 50 Triatoma dimidiata vector of Chagas disease. PLoS Negl Trop Dis, 2011. 5(5), e1045.</li> <li>51 92. Castillo-Riquelme, M., et al., Modelling geographic variation in the cost-effectiveness of 52 control policies for infectious vector diseases: the example of Chagas disease. J Health Econ, 53 2008. 27(2), 405-26.</li> <li>54 93. Basombrio, M.A., et al., A cost-benefit analysis of Chagas disease control in north-western</li> </ul>		00	
<ul> <li>Barbu, C., et al., Evaluation of spatially targeted strategies to control non-domiciliated Triatoma dimidiata vector of Chagas disease. PLoS Negl Trop Dis, 2011. 5(5), e1045.</li> <li>Castillo-Riquelme, M., et al., Modelling geographic variation in the cost-effectiveness of control policies for infectious vector diseases: the example of Chagas disease. J Health Econ, 2008. 27(2), 405-26.</li> <li>Basombrio, M.A., et al., A cost-benefit analysis of Chagas disease control in north-western</li> </ul>		<i>J</i> 0.	
<ul> <li>Triatoma dimidiata vector of Chagas disease. PLoS Negl Trop Dis, 2011. 5(5), e1045.</li> <li>Castillo-Riquelme, M., et al., Modelling geographic variation in the cost-effectiveness of control policies for infectious vector diseases: the example of Chagas disease. J Health Econ, 2008. 27(2), 405-26.</li> <li>Basombrio, M.A., et al., A cost-benefit analysis of Chagas disease control in north-western</li> </ul>		01	
<ul> <li>51 92. Castillo-Riquelme, M., et al., Modelling geographic variation in the cost-effectiveness of</li> <li>52 control policies for infectious vector diseases: the example of Chagas disease. J Health Econ,</li> <li>53 2008. 27(2), 405-26.</li> <li>54 93. Basombrio, M.A., et al., A cost-benefit analysis of Chagas disease control in north-western</li> </ul>		<i>9</i> 1.	
<ul> <li>52 control policies for infectious vector diseases: the example of Chagas disease. J Health Econ,</li> <li>53 2008. 27(2), 405-26.</li> <li>54 93. Basombrio, M.A., et al., A cost-benefit analysis of Chagas disease control in north-western</li> </ul>		92	
<ul> <li>2008. 27(2), 405-26.</li> <li>93. Basombrio, M.A., et al., A cost-benefit analysis of Chagas disease control in north-western</li> </ul>		14.	
54 93. Basombrio, M.A., et al., A cost-benefit analysis of Chagas disease control in north-western			
		93	

1 2	94.	Meheus, F., et al., Cost-effectiveness analysis of combination therapies for visceral leishmaniasis in the Indian subcontinent. PLoS Negl Trop Dis, 2010. 4(9).
3	95.	Vanlerberghe, V., et al., Drug policy for visceral leishmaniasis: a cost-effectiveness analysis.
4 5	96.	Trop Med Int Health, 2007. <b>12</b> (2), 274-83. Boelaert, M., et al., <i>How better drugs could change kala-azar control. Lessons from a cost-</i>
6 7	97.	effectiveness analysis. Trop Med Int Health, 2002. 7(11), 955-9. Boelaert, M., et al., Cost-effectiveness of competing diagnostic-therapeutic strategies for
8	97.	visceral leishmaniasis. Bull World Health Organ, 1999. <b>77</b> (8), 667-74.
9 10	98.	Lee, B.Y., et al., <i>The economic value of a visceral leishmaniasis vaccine in Bihar state, India.</i> Am J Trop Med Hyg, 2012. <b>86</b> (3), 417-25.
		Am J Trop Med Hyg, 2012. <b>60</b> ( <i>3)</i> , 417-23.
11		
12 13		
14		
15		
16		
17		
18		
19		
20		
21		
22		
23		
24		
25		
26		
27		
28		
29		
30		
31		
32		
33		

Glossary	/
----------	---

Cost-effectiveness analysis: a method for assessing the relative gains in health
generated by a health intervention compared to the costs.

- Disability-Adjusted Life-Years (DALYs): a time-based measure of disease burden accounting for years of life lost due to premature mortality and healthy years of life lost due to disability[48].
  - **Density dependent process:** a demographic or transmission process whose rate is regulated by the density of parasites in the host or population.

**Dynamic transmission model**: a model that links the rate at which individual hosts acquire new infections with the abundance of infection among all hosts in a population.

Elimination of infection: reduction to zero of the incidence of infection caused by a specific agent in a defined geographic area as a result of deliberate efforts; continued measures to prevent reestablishment of transmission are require.

Neglected Tropical Diseases (NTDs): a group of chronic, disabling, and disfiguring conditions that occur especially among the rural poor and disadvantaged urban populations.

**Macroparasites:** parasites that are large enough to be seen with the naked eye (e.g. parasitic worms) and which do not multiply directly within the definitive host.

- Static transmission model: a model that assumes the rate at which individual hosts acquire new infections is *independent* of the abundance of infection among all hosts in a population.
- Herd effect: the indirect benefit afforded to individuals not directly targeted by an intervention that arises from the reduction in transmission that ensues from an effective intervention.

Transmission breakpoint: the (non-zero) parasite density below which a parasite
 population cannot maintain itself and is driven into terminal decline and eventual elimination.

#### **Box 1 Density-dependent processes**

A process is density-dependent when the rate at which it occurs is determined by the number (density) of parasites within the host or in the parasite population as a whole. These processes can either restrict or enhance transmission and can act at multiple points in a parasite's life cycle [29].

#### Negative density dependence

Negative (down-regulatory, constraining or restricting) density-dependence occurs when the rate at which a process occurs decreases with increasing parasite density. For example, the per capita rate of larval development within the vectors of filarial nematodes often decreases as parasite density increases [28]. Interventions that lead to a reduction in parasite abundance can cause a relaxation of these negative density-dependent restrictions, increasing per capita rates of transmission, making parasite populations resilient to perturbation, and leading to a lower than hoped impact of an intervention.

#### Positive density dependence

Positive (facilitating or enhancing) density-dependent processes occur when rates increase with increasing parasite intensity. The Allee effect is an example of a positive density dependency. A more specific example is the probability that a female worm is mated within the definitive host [49] increases with parasite density as there is a greater chance of worms of different sexes finding each other and mating. Positive density-dependence can make elimination of a parasite population more likely as they can cause the reproductive success of a parasite to decrease with decreasing parasite densities giving rise to breakpoints below which the population cannot sustain itself and enters terminal decline.

2

3

4

5

6

7

8

9

10

Total 0	Static Models	<b>Dynamic Models</b> <sub>5</sub>	
		0	6
2	0	2 <sup>[50, 51]</sup>	7
			8
3	1 [52]	2 <sup>[53, 54]</sup>	Q
4	3 [55-57]	1 [58]	10
	150 (2)		11
5	5 <sup>[59-63]</sup>	0	12
13	6 [64-69]	7 [70-76]	13
2	0	2 <sup>[77, 78]</sup>	14
3	1 <sup>[79]</sup>	2 [80, 81]	15
			16
12	12 <sup>[82-93]</sup>	0	17
5	5 <sup>[94-98]</sup>	0	18
			19
	0 2 3 4 5 13 2 3 12	$\begin{array}{cccc} 0 & 0 \\ 2 & 0 \\ 3 & 1^{[52]} \\ 4 & 3^{[55-57]} \\ 5 & 5^{[59-63]} \\ 13 & 6^{[64-69]} \\ 2 & 0 \\ 3 & 1^{[79]} \\ 12 & 12^{[82-93]} \end{array}$	$\begin{array}{c ccccc} 0 & 0 & 0 \\ 2 & 0 & 2^{[50, 51]} \\ 3 & 1^{[52]} & 2^{[53, 54]} \\ 4 & 3^{[55-57]} & 1^{[58]} \\ 5 & 5^{[59-63]} & 0 \\ 13 & 6^{[64-69]} & 7^{[70-76]} \\ 2 & 0 & 2^{[77, 78]} \\ 3 & 1^{[79]} & 2^{[80, 81]} \\ 12 & 12^{[82-93]} & 0 \end{array}$

# Table 1: A summary of identified modelling studies presenting economic evaluations across the diseases considered by the London Declaration on NTDs.

- 1 Figure 1: The model projected indirect benefit of treating children (2-14 year olds) for
- 2 Ascaris lumbricoide for untreated adults. The results were obtained using a fully aged
- 3 structured deterministic dynamic transmission model, described in more detail in [35, 36].
- 4 Results assume a low transmission setting (a basic reproduction number, R0, of 2) and a high
- 5 coverage (80%).

# 6 Figure 2: Schematic representation of the nonlinear relationship between parasite

- 7 **density and transmission.** *The blue line indicates the situation for a linear relationship*
- 8 between parasite density and transmission; the red line illustrates the potential impact of
- 9 *negative density dependence processes on the relationship.*
- 10 Figure 3: Selection criteria for inclusion and exclusion of modelling studies presenting
- 11 economic evaluations of neglected tropical disease interventions. *The search terms*
- 12 *included all of the variants of the following terms: mathematical model; dynamic model static*
- 13 model; decision trees; cost benefit analysis; cost effectiveness analysis; economics, and
- 14 *economic evaluation.*
- 15
- 16
- 17
- 18
- 19

Dear Editor,

Thank you very much for considering our manuscript. We greatly appreciate the reviewer's and editor's suggestions for improvement and have endeavoured to address the issue raised (see below).

#### **Editor's Comments**

<u>Editor's comment 1</u>. Abstract. I have below a suggested modification to the abstract that bears in mind the recommendation of the piece being most suitable as an Opinion article that brings forth the authors' perspective on a topic:

Despite many current interventions against neglected tropical diseases (NTDs) being highly cost effective, new strategies are needed to reach the World Health Organization's control and elimination goals. Here we argue for the importance of incorporating economic evaluations of new strategies in decisions regarding resource allocation. Such evaluation should ideally be conducted using dynamic transmission models that capture inherent nonlinearities in transmission and the indirect benefits ('herd effect') of interventions. A systematic review of mathematical models that have been used for economic analysis of interventions against the ten NTDs covered by the London Declaration reveals that only 16 out of 49 studies used dynamic transmission models, highlighting a fundamental – but addressable – gap in the evaluation of interventions against NTDs.

#### Authors' response

Thank you, for the suggested edits. We have incorporated them into the manuscript.

<u>Editor's comment 2</u>. Aims of the article in the introductory section. Please add a brief paragraph at the end of the introductory section of the article that states the aims of the article (i.e. what key point does the article put forward and why is this important to discuss?). This will provide a framework so that the reader can approach the next sections of the article with these aims in mind.

#### Authors' response

We have added the following to the end of the Introduction:

"In the context of the London Declaration, models will be needed to perform economic evaluations to investigate what changes in policy are required to meet the 2020 NTD goals in the most cost-effective way. The aim of this paper is to (a) highlight why this area of research should be addressed with dynamic, rather than the more commonly used static transmission models, and (b) to outline what we feel are key research needs for this area." <u>Editor's comment 3</u>. Concluding remarks. Please include in the Concluding Remarks section a brief summary of the key points you raised in the "Research Needs" section.

# Authors' response

We have incorporated the following into the Concluding remarks.

"In order for this to happen effectively, a wide range of research needs should be addressed including, but not limited to: i) more empirical studies and greater availability of programmatic data to further parameterise/develop these dynamic transmission models, and to improve estimated health effects, ii) the advancement of economic evaluation frameworks and collection of detailed cost data that account for integrated NTD control programmes, and iii) rigorous model validation and comparison."

<u>Editor's comment 4</u>. Highlights. I have a suggested revision to the manuscript's highlights below, mainly aimed at highlighting the "Research Needs" section.

Highlights

- New strategies are needed to reach the WHO NTD control and elimination goals.
- Economic evaluations using mathematical models could improve resource allocation.
- Dynamic models are needed to account for nonlinearities in transmission.
- Challenges and research needs to address this gap are outlined.

#### Authors' response

Thank you for these excellent edits. We have incorporated them into the manuscript highlights.

# **Referees Comments**

### Reviewer #1:

The authors observed that dynamic transmission models were used in only 16/49 modelbased economic evaluation studies in the field of NTDs, argue that this is a fundamental research gap, and present an agenda for research. I am not convinced that this paper needs to be published now in this journal, for the following reasons:

1.1. In 2012, PloS-NTDs publishd a series of research agenda's on helminthic diseases (as subgroup of NTDs), including one on modelling (Basanez et al). At this point, I would rather want to see new research than yet another (and not particulary renewing) review / research agenda.

### Authors' response 1.1

We feel it is important to highlight that the PLoS-NTDs review to which the Reviewer refers is only on helminthiases, where as in this paper we cover all of the NTDs considered by the London Declaration (including leprosy, trachoma, sleeping sickness, Chagas disease, and visceral leishmaniasis). Furthermore, this paper focuses on the use of models for economic evaluations (a relatively minor aspect of the PLoS NTDs review) which we believe, especially in the context of the London Declaration, will have an increasingly important role – and more so than in the past – for decision making and resource prioritisation, including investigating the most cost-effective changes to policy to meet the World Health Organization's 2020 elimination goals.

We have endeavoured to make our assertions clearer by making changes to the Introduction and Conclusions sections of the manuscript, and by explicitly setting out the goals of the paper. Following the Editor's suggestion, we have also changed the manuscript into an Opinion piece, rather than a Review.

1.2. The review component of this paper is very small and superficial. The authors observed that dynamic transmission models were used in only 16/49 model-based economic evaluation studies in the field of NTDs; other studies used static models. The authors acknowledge that the use of static models may not always have been the wrong choice (depending on both the intervention and the parasite under investigation), but fail to evaluate whether the choice of models has been appropriate or not.

# Authors' response 1.2

We agree with the Reviewer that the review component is relatively minor and not an integral part of the manuscript. Therefore, after discussions with the Editor, and following her recommendation, we have changed the article to an Opinion piece. In this new context, we believe that the level of detail given on the methodologies used by the articles identified by our systematic review is sufficient to support our assertion that dynamic transmission models are seldom used for the economic analysis of interventions against NTDs.

1.3. The authors present the lack of economic evaluations using mathematical models as a major, even fundamental, research gap. I would argue that there is a research gap on almost everything when it comes to the neglected diseases. I am not convinced that economic evaluations in general or economic evaluations with dynamical models are more neglected than other research aspects (at least, not by the "evidence" presented here). The more important research questions may simply be on different aspects. For me, the lack of

empirical studies in general is the more fundamental gap. Although I do agree, that mathematical models are not used to their maximum potential to inform both new research projects and control policies.

## Authors' response 1.3

We agree with the Reviewer that research on NTDs in general remains relatively overlooked and underfunded, albeit to a lesser extent now than in the past. Yet we firmly believe that to maintain the unprecedented commitment and drive by the global health community to rid the world's most impoverished people of these diseases, it is crucial that resources are most effectively deployed and that wasteful, ineffective, or ill-suited intervention strategies are avoided. The coupling of economic evaluations to dynamic transmission models provides an essential analytical framework to achieve this broad strategic goal. We do agree with the Reviewer that the lack of empirical studies is indeed a pressing research gap, and have made greater reference to this in the text (including in the Concluding remarks).

# Empirical and programmatic data

"A key impediment to the evaluation of NTD interventions, both in terms of economic and public-health outcomes, is the lack of longitudinal data from large-scale, long-term control programmes [18] which are made available to modellers and analysts for model parameterization and impact evaluation. This relative paucity of empirical studies is in itself another major research gap for NTDs."

Concluding remarks

•••••

"In order for this to happen effectively, a wide range of research needs should be addressed including, but not limited to: i) more empirical studies and greater availability of programmatic data to further parameterise/develop these dynamic transmission models, and to improve estimated health effects, ii) the advancement of economic evaluation frameworks and collection of detailed cost data that account for integrated NTD control programmes, and iii) rigorous model validation and comparison."

#### Reviewer #2:

This is a well written paper that argues for dynamic modelling for NTD. This is the same argument as used for vaccination programmes (see work by W. John Edmunds et al), but for a different audience. Although the paper makes a good case, it is largely from the point of view of the modellers (of which I am one). I think that I missed a novel insight, or good evidence that "proper" modelling would change the situation dramatically. As far as I can see, the case needs to be made for more analysis and data against just intervening. I have listed some points below that the authors might wish to consider when re-writing.

#### Authors' response

We thank the referee for these appreciative comments.

Comment 2.1) P4 L3 - "public health resources" - perhaps a hyphen to distinguish between public-health resources and public health-resources. Also it might be worth including a

sentence or two about the relationship between political and economic objectives and the role of public health. Most of the current impetus to control NTD is from international perspective (World Bank, WHO, BMGF etc), and this perspective matters when considering the economics of control. They might, for example, be more interested in making an impact quickly rather than spending a long time doing the research to find the most cost-effective intervention for a particular situation. Modelling costs, not only in terms of direct resources, but also in terms of potential delays to decision making - is it worth this cost from the perspective of the funders?

#### Authors' response 2.1

We have added the hyphen as suggested to allay any ambiguity and discussion regarding relationship between political and economic objectives.

"The use of mathematical models to perform economic evaluations of new health care interventions is an important tool for deciding how to allocate public-health resources [9]. Accurately parameterised mathematical models have the advantage of enabling investigation of the cost effectiveness of various strategies in a range of different epidemiological and programmatic scenarios, and can therefore help optimise the use of resources. This is particularly important for NTDs, which are most prevalent in resource-poor settings and historically have suffered from a lack of funding (for control implementation and research), visibility, and political advocacy. Modellers must work effectively with funders and politicians who will often want to demonstrate impact quickly rather than delaying decision making while comprehensive evaluations are undertaken. Consequently, models must be flexible and versatile enough to capture the effects and accompanying uncertainties of past and ongoing intervention activities."

Comment 2.2) - Static models are fine if they are only considering a one-time intervention in a narrow time window. The failure comes when they don't consider the time effects - future incidence is dependent on current incidence (which is dependent on current prevalence). Inclusion of the time dimension in the discussion of static infection models would be welcome.

#### Authors' response 2.2

#### We agree and have added more discussion regarding time dimension into the manuscript.

"However, and critically, dynamic transmission models capture the so-called 'herd effect' (or indirect effects) of interventions targeting particular population groups. Such interventions include vaccination campaigns and school-based mass drug administration (MDA) programmes, whereby all individuals within a population can benefit from the intervention, albeit in a differential manner, regardless of whether they are within the target group or not. Figure 1 illustrates this concept by showing that a school-based MDA programme treating children for *Ascaris lumbricoides* can have a notable indirect (herd) benefit for the untreated adults whose worm burden is also reduced over time, due to population-wide reductions in transmission. The impact of these indirect herd effects will accumulate over time as reductions in transmission affect reductions in incidence which, in turn, manifest more slowly in reduced population levels of infection. Consequently, the extent that static models underestimate the impact of transmission reducing interventions will increase over

time. By the same token, static models may provide a reasonable approximation to dynamic models over a relatively short time frame."

"Although static models are suitable for the evaluation of interventions that do not markedly affect the targeted parasites transmission cycle—such as when there is a large untargeted reservoir of parasites or vectors, when evaluating over a short time horizon—they notably underestimate the impact of, and thus are inappropriate for evaluating, interventions that reduce levels of infection transmission in a population, such as MDA or vaccination campaigns. This is particularly important when investigating elimination strategies."

Comment 2.3) - Dynamic models generally have greater inherent uncertainty because of the difficulties of parameterization and structure. This model uncertainty is probably a better measure of real uncertainty than that which can be gained from static models. The paper would be improved by explicitly mentioning uncertainty in decision making based on models - the models should not only get the expectation right, they should get the risk of different outcomes right as well.

#### Authors' response 2.3

We agree with the Reviewer on this very important point and we have incorporated it in the revised manuscript.

#### Model validation and comparisons

"By explicitly considering transmission, dynamic models have greater inherent uncertainty than their static counterparts, in both parameterization and structure. This makes their validation even more important. However, adequately reflecting uncertainty in our understanding of infections and their transmission dynamics has an important role in model-based decision making; policy makers should be presented with both the expectation and the accompanying uncertainty associated with epidemiological and economic projections alike."

Comment 2.4)-Elimination. The London Declaration et al. have generally confused the concept of elimination. Some of what is written here presumes the original definition "Reduction to zero of the incidence of a specified disease (/infection) in a defined geographical area as a result of deliberate efforts; continued intervention measures are required". However, the current 2020 goals are written as "elimination as a public health problem" which is defined as control in the original definition. What is written here is conflating the confusion. Precision of these ideas will become an increasingly important issue as incidence is reduced. I think that you should at least define here what you mean, and that should include the reference the Dahlem conference that defined it. I actually think that this confusion over the term elimination is exactly what models are good at avoiding.

#### Authors' response 2.4

We have added into the manuscript and the glossary the definition of elimination as articulated at the Dahlem conference (along with the reference). We have also removed reference to the London Declaration in this section to avoid confusion.

"Second, dynamic models can be used to evaluate the possibility of infection elimination under specific intervention strategies or intervention combinations (i.e. the reduction to zero of the incidence of infection caused by a specific agent in a defined geographic area as a result of deliberate efforts; continued measures to prevent re-establishment of transmission are required { Added Suggested reference}. For this purpose, dynamic models should generally be stochastic, capturing the reality of random chance in demographic and transmission processes. This is because such random variation becomes increasingly important in determining whether a disease is eliminated as infection is decreased to very low levels, a process known as 'stochastic fade-out'. By contrast, in (non-stochastic) deterministic models, all events occur in a pre-specified way depending on the parameter values and initial conditions of the model, ignoring the effects of chance. Stochastic models are particularly relevant to NTDs as goals shift from control to elimination because new interventions are likely to be more expensive than current tools and may only be deemed cost effective if they lead to elimination faster, avoiding future costs [24]."

Comment 2.5)-You have argued (quite well) above that a static model is almost always wrong. Static models are always wrong for vaccination, and will seriously under-estimate the impact for any other intervention.

## Authors' response 2.5

We agree with the Reviewer that it is important to state explicitly that static models will underestimate the impact of interventions which reduce transmission in a population and are being evaluated over a considerable timespan. As the Reviewer suggests, this is particularly relevant to vaccination campaigns but also to MDA programmes. We have modified that text in several places to emphasize this important point.

"Consequently, although static models are suitable for the evaluation of interventions that do not markedly affect the targeted parasite's transmission cycle, such as those that relieve disease symptoms and sequelae but do not treat infection, or those that target a very small fraction of the vector, host, or parasite population (leaving a large infectious reservoir), they are rarely suitable for evaluating interventions that reduce levels of infection transmission in a population. In such circumstances, static models will underestimate the impact of an intervention"

"Although static models are suitable for the evaluation of interventions that do not markedly affect the targeted parasite's transmission cycle—such as when there is a large untargeted reservoir of parasites or vectors or when evaluating over a very narrow time horizon—they notably underestimate the impact of, and are thus inappropriate for evaluating, interventions that reduce levels of infection transmission in a population, such as MDA or vaccination campaigns. This is particularly important when investigating elimination strategies."

Comment 2.6) - Have transmission breakpoints ever been demonstrated, or are they only a theoretical construct? Are they actually useful in control terms? More important, I suggest, is the modeling of potential patterns of resurgence, and the surveillance required to detect them.

#### Authors' response 2.6

To demonstrate a transmission breakpoint one would have to first suppress, with an effective intervention strategy, a parasite population to extremely low but detectable levels and then, withholding the intervention, observe the residual parasite population decline to local extinction. By these criteria, transmission breakpoints have not been demonstrated and consequently remain largely theoretical. Yet they remain an integral part of deterministic transmission models (much less so in stochastic models) which could otherwise not strictly be used to make predictions on elimination, although empirical programmatic thresholds are sometimes used as an alternative given the theoretical and unproven nature of the breakpoints. In practice, theoretical breakpoints are not used as an explicit goal for intervention campaigns and we agree with the Reviewer on the more important practical issue of modelling patterns of potential resurgence and the necessary surveillance to detect this after cessation of an intervention. We have added this to the research needs section.

# Transmission breakpoints and surveillance strategies

"Transmission breakpoints theoretically arise due to the operation of positive density dependencies (Box 1) and refer to the density......"

"Dynamic models can also be useful in investigating patterns of potential infection resurgence following cessation of control and in designing cost-effective surveillance strategies to detect this. This area will become increasing important as progress towards the 2020 NTD goals is made."

Comment 2.7) Research Needs - personally I think that the validation of models is a step that is too often ignored and has a negative impact on the use of models. There needs to be more funding for the relatively boring task of showing why a particular model should be believed. However, validation is usually held back by lack of appropriate data.

#### Authors' response 2.7

We agree with the Reviewer's salient remark that model validation using appropriate data is an important research need and we have added this point into the research needs section of the revised manuscript.

#### Model validation and comparisons

"More funding and resources need to be made available for model validation; before undertaking detailed evaluations of competing intervention strategies, one must be able to demonstrate that a model adequately describes epidemiological patterns both in endemic parasite (pathogen) populations and, crucially, in populations perturbed by control activities. This process is often held back by a lack of appropriate longitudinal data from sentinel populations. Indeed, the gathering of such data is often not prioritized highly enough despite being essential to robust policy-relevant modelbased evaluations.

Comment 2.8)- this relates back to the previous point. Why is this an "important research gap"? The prevalence of all NTD has been dramatically reduced - have models been important in this process? As a modeler, I can see the strong philosophical argument, but in

the case of NTD I am not sure that the use of modeling has been validated.

## Authors' response 2.8

We agree with the Reviewer that hitherto modelling has not played a hugely prominent role in influencing NTD control strategies (with some notable exceptions such as in onchocerciasis control) and in this sense its relevance to policy has not been truly accomplished. However, in the context of the London Declaration, and as tools to investigate what changes to policy and resource allocation need to be made to meet the demanding 2020 NTD goals, we believe that models will play a more substantial and important role than in the past. Recognising this need, the B&MGF is considering the establishment of an NTD Modelling Consortium which would fuel the development, refinement and rigorous comparison and validation of dynamic models for the diseases under the London Declaration.

We have made changes to the Introduction and Conclusions sections of the manuscript to clarify this argument and the goals of the paper (see Authors' response 2.9).

Comment 2.9) Better to avoid hyperbole ("vital" is used three times, "crucial" twice, "critical" three times). Usually what is meant is "potentially important". I don't get a sense from the paper of where the authors believe modelling should sit in the process of decision making. What are they arguing for? This comes across as a bit of a general whinge rather than clearly providing evidence that using static models gets the answer wrong, and that this has had a bad outcome.

#### Authors' response 2.9

We thank the Reviewer for his/her stylistic suggestions and caution against hyperbole. We have amended the following sentences to somewhat temper the vivacity of our arguments in the revised manuscript:

"Thus, the microparasite population would be less likely to recover after cessation of MDA, highlighting the potential importance of accounting for these effects when evaluating elimination strategies."

"Comprehensive costs of NTD interventions should be incorporated into the modelling analyses, in particular the relative cost of alternative/complementary strategies, which may change with scale and can be influenced by other related intervention activities"

"Moreover, multiple perspectives should be investigated, not just from the health care providers (i.e. the control programmes and non-governmental organization (NGO) donors), but also that from the pharmaceutical companies that donate or discount drugs, as different intervention strategies will influence the costs associated with these parties in different ways."

"Accounting for the herd effect is potentially important to the validity, robustness and policy relevance of conclusions drawn from cost-effectiveness evaluations of interventions against transmissible diseases in general, and of NTDs in particular [14, 15, 17]."

"Programmatic data should also be used to validate current and future model projections and, with individual data, to understand better the impact of individual

# heterogeneities in responses to interventions (such as individual drug responses) on population-level benefits and cost-effectiveness."

At the suggestion of the Editor, the article has now changed from a review to an opinion piece, allowing us greater freedom to make our central argument more explicitly; namely, that static models will not be appropriate to address the current research questions in the context of the WHO's 2020 control and elimination targets and the London Declaration

#### Aim

"In the context of the London Declaration, models will be needed to perform economic evaluations to investigate what changes in policy are required to meet the WHO 2020 goals in the most cost-effective way. The aim of this paper is to (a) highlight why this area of research should be addressed with dynamic, rather than the more commonly used static transmission models, and (b) to outline what we feel are key research needs for this area."

### **Concluding Remarks**

"The lack of economic evaluations using dynamic transmission models demonstrates an important research gap and methodological shortcoming in the evaluation of NTD control and elimination interventions. This is particularly important in the context of the London Declaration, where dynamic models will be nesseccary to investigate what changes to policy are required to meet the 2020 NTD control and elimination goals. Despite being easier to develop, and more commonly used, static infection models are generally ill-suited for this undertaking, potentially generating inaccurate results leading to poor policy decisions on intervention strategies.

We believe that reaching the 2020 NTD goals will not only depend on the continuing and burgeoning implementation of existing intervention strategies, and the development of new more effective intervention tools, but also on the concomitant development of judiciously formulated and well parameterized dynamic transmission models with which to undertake economic evaluations. Such evaluations will inform policy by determining the optimum way in which intervention tools should be deployed in fundamentally resource-poor settings.

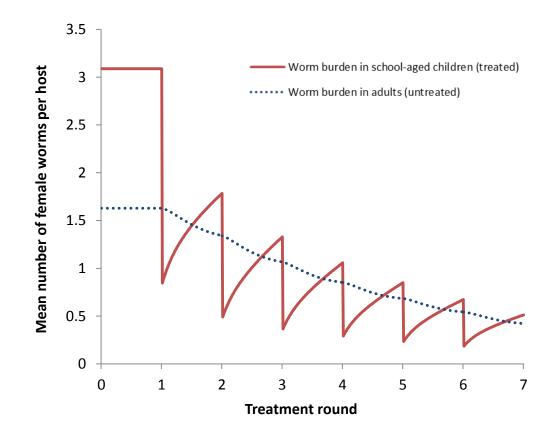
In order for this to happen effectively, a wide range of research needs should be addressed including, but not limited to: i) more empirical studies and greater availability of programmatic data to further parameterise/develop these dynamic transmission models, and to improve estimated health effects, ii) the advancement of economic evaluation frameworks and collection of detailed cost data that account for integrated NTD control programmes, and iii) rigorous model validation and comparison." Turner *et al* Neglected tools for neglected diseases: mathematical models in economic evaluations

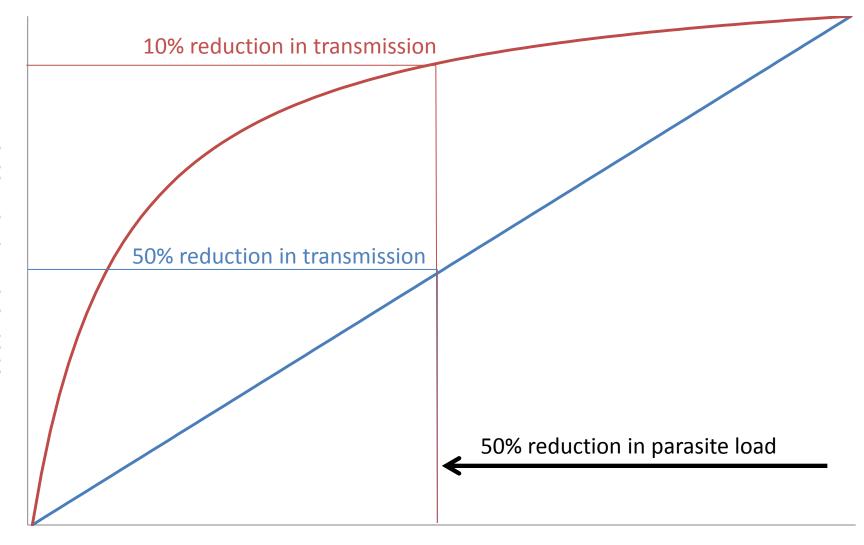
# Highlights

• New strategies are needed to reach the WHO NTD control and elimination goals.

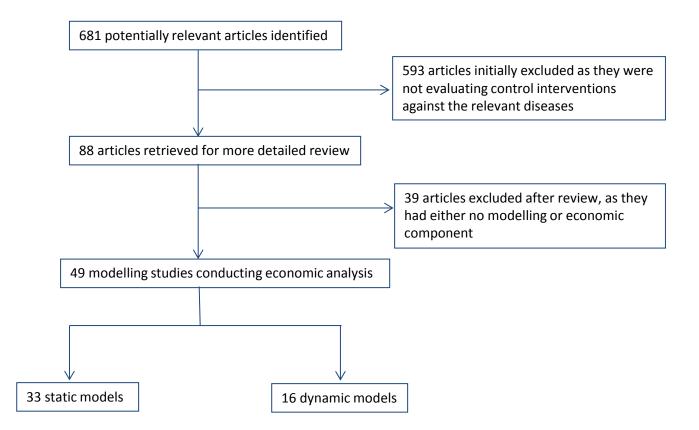
• Economic evaluations using mathematical models could improve resource allocation.

- Dynamic models are needed to account for nonlinearities in transmission.
- Challenges and research needs to address this gap are outlined.





**Parasite Density** 



Original Figure File 1 Click here to download Original Figure File: Figure 1\_data.xlsx Original Figure File 2 Click here to download Original Figure File: Figure 2\_data.xlsx