The effect of temperature increase on the development of *Rhodnius prolixus* and the course of *Trypanosoma cruzi* metacyclogenesis

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Abstract

The increase in the global land temperature, expected under predictions of climate change, can directly affect the transmission of some infectious diseases, including Chagas disease, an anthropozoonosis caused by *Trypanosoma cruzi* and transmitted by arthropod vectors of the subfamily Triatominae. This work seeks to study the effects of temperature on the development of the life cycle, fertility and fecundity of the insect vector *Rhodnius prolixus* and on the metacyclogenesis of *T. cruzi*. All of the variables were subjected to 3 temperatures: 26˚C, 28˚C and 30˚C. Hatching time was evaluated, along with time to fifth instar, time to adult, fecundity studied using the e-value, and egg viability during the first 3 reproductive cycles. In addition, the amounts of metacyclic trypomastigotes of the TcI and TcII DTUs in *R. prolixus* were evaluated from days 2 to 20 at two-day intervals and from weeks 6 to 8 post-infection. Decreases were observed in time to hatching (15–10 days on average) and in time to fifth instar (70–60 days on average) and transition to adult (100–85 days on average). No significant differences in egg viability were observed in any of the reproductive cycles evaluated, but an increase in fecundity was observed at 30˚C during the third reproductive cycle. At 30˚C, there was also an increase in the number of infective forms and a decrease in the time at which metacyclic trypomastigotes were detected in the rectal ampulla of the insects for both TcI and TcII. According to these results, the expected temperature increase under climate change would cause an increase in the number of insects and a greater probability of infection of the parasite, which affects the transmission of Chagas disease.
Author summary

Chagas disease is an anthropozoonosis caused by the flagellated protozoan *Trypanosoma cruzi* and mainly transmitted through the infected faeces of insects of the subfamily Triatominae. Because these insects are sensitive to climatic conditions, it is expected that disease transmission may be affected by the increase in global land temperature, predicted under climate change. Therefore, we wanted to evaluate the effect of temperature increase on the development, viability of eggs and fertility of *R. prolixus*, the most important vector insect in Colombia, and on the development of the parasite within this insect. We observed a decrease in the development time of *R. prolixus* and an increase in the number of infectious forms of *T. cruzi* in the insect as the temperature increased. These results suggest that if the temperature increases as expected, there may be an increase in the number of insects that can transmit the disease, as well as an increase in the likelihood of infection due to the increase in the number of infectious forms. Our data contributes to the understanding of the possible effects of the expected temperature increase under climate change on Chagas disease transmission and can be used to make predictive models that can more accurately predict the future of Chagas disease.

Introduction

According to climate prediction models, it is expected that by the end of the 21st century, the global terrestrial temperature will increase between 0.3°C and 4.8°C with respect to the average temperature observed between 1986 and 2005 [1]. This increase can affect the transmission of infectious disease agents transmitted by vectors because both insects and vertebrate reservoirs are sensitive to climatic conditions [2]. The alteration in transmission occurs because the temperature and rainfall expected under the effect of climate change can affect the range, proliferation, viability and maturation rates of vectors, pathogens and reservoirs [3]. Several studies have reported variations in the transmission of diseases such as malaria [4], dengue [5,6] and leishmaniasis [7,8]. In the case of malaria, it has been observed that the time taken by *Plasmodium*, within the vector mosquito, to pass from gametocytes to infective sporozoites decreases when the temperature of the environment increases [9]. Likewise, a decrease in the development time of the insect vector has been reported, which leads to an increase in the number of insects per season and consequently to a higher transmission rate [10]. In some countries, such as Colombia, Venezuela, Guyana and Peru, there has been a resurgence or intensification of endemic and epidemic malaria that correlates with the phenomenon of the El Niño Southern Oscillation (ENSO) [11, 12]. Likewise, changes in the distribution of leishmaniasis and Dengue virus vectors that may lead to an increase in the risk of transmission of these diseases have been predicted [5, 7].

Another parasitic disease transmitted by vectors that could be affected by climate change is Chagas disease, an anthropozoonosis caused by the flagellated protozoan *Trypanosoma cruzi*, which is transmitted to humans and other mammals mainly through the infected faeces of insects of the subfamily Triatominae [13]. The disease is a serious public health problem; it is estimated that more than eight million people are infected in Latin America [14]. Despite efforts to control vectors in various regions of Latin America, the World Health Organization estimates that 5,274 new cases occur annually due to vector transmission in Colombia, 933 in Honduras and 873 in Venezuela, countries in which *Rhodnius prolixus* is the main transmitting vector of the disease [15].
More than 150 species of triatomines have been reported in Latin America, of which 10 are considered primary vectors of the parasite because they colonize houses, while another 20 are considered secondary vectors because they invade human habitations from their peri-domestic or wild habitat. [16] R. prolixus is present in both domestic and wild transmission cycles, which is why it is imperative to design very specific vector control strategies [17]. The life cycle of this vector includes 5 nymphal stages and an adult stage, all obligate haematophagous. None of the nymphs can develop into another stage without having fed at least once. In addition, adults cannot produce eggs without ingesting blood, as it is essential for this process [18]. R. prolixus is considered the most important vector of T. cruzi transmission in Colombia, Venezuela and most Central American countries [13,19]. Also, different outbreaks of oral transmission have been reported in Colombia and Venezuela incriminating R. prolixus [20, 21, 22].

The effect of temperature over R. prolixus life cycle has been barely studied. It is well known that under laboratory conditions, the temperature range for eclosion and molting of R. prolixus was reported to be between 16–34˚C [23]. No development was observed at 15˚C and 35˚C [24, 25]. Development is generally studied at constant temperatures between 25 and 28˚C and about 70% humidity or unspecified conditions of ambient temperature and humidity. However, temperature and humidity in insect habitats may differ considerably and vary according to circadian and seasonal patterns. R. prolixus is found mainly in Colombia and Venezuela from 0 to 2,600 m above sea-level, in regions with annual median temperatures from 11 to 29˚C and 250 to 2,000 mm annual precipitation [26, 27, 28].

The biological cycle of T. cruzi in the vertebrate host begins with contact of the faeces of a triatomine infected with metacyclic trypomastigotes. The parasite penetrates the cells by forming a vacuole bound to lysosomes, then escapes from the vacuole and differentiates into amastigotes, which reproduce in the cytoplasm by binary fission. Subsequently, amastigotes differentiate into metacyclic trypomastigotes that are mobile. These latter forms of the parasite lyse the host cell, disperse and can infect other cells. When insects ingest blood from an infected mammal, trypomastigotes in the blood differentiate into epimastigotes, and some develop into spheromastigotes. The epimastigotes divide in the midgut by binary fission and migrate to rectal ampulla, where they are transformed into metacyclic trypomastigotes (a process called metacyclogenesis), which are eliminated in the faeces of the triatomine [29]. T. cruzi, in turn, has high genetic variability, which has been subdivided by international consensus into six Discrete Typing Units (DTUs), TcI—TcVI, including a DTU associated with anthropogenic bats and called TcBat [30, 31, 32]. In the northern countries of the southern cone, TcI and TcII are the most frequent DTUs in humans, vectors and reservoirs [30]. In Colombia, it is observed that TcI and TcII are the most frequent DTUs in Rhodnius prolixus [17].

Efforts to predict the future of transmission of Chagas disease under the effects of climate change have been scarce and have focused, above all, on trying to establish possible changes in the distribution of vectors through mathematical models [33, 34, 35, 36, 37, 38]. In general, the results obtained have been varied and seem to depend on the vector species, which is why more studies are needed on the physiology of the parasite and the vector to enable more reliable predictions. The objective of this work is to study the effects of temperature, expected under predictions of temperature increase on the development of the life cycle of the insect vector R. prolixus and on the metacyclogenesis of T. cruzi under controlled laboratory conditions.

**Materials and methods**

**Study design**

The T. cruzi scenarios of transmission have been previously studied by our group in Maní, Casanare (Colombia) and is characterized by dense forests of Palms. Infestation rates of 100%
in a transect of 120 studied palms were reported [39]. Insect were collected manually in the the axils of *A. butyracea* palms and transported to our laboratory in the same day of capture.

Specimens of *R. prolixus* collected in *Attalea butyracea* palms were kept in incubators that simulated the same average annual humidity and temperature conditions of the axils of the palms where they were collected (26˚C, 80% RH and photoperiod 12:12). Temperature was measured using an "EXTECH RHT 20: Humidity / Temperature Datalogger" in different palms. Previous field reports stated that the temperature in palms axils is 26˚C-27˚C according to Urbano et al. [40]. On the other hand, these devices were also used to measure and control the three temperatures (26, 28, 30˚C) in the incubators, during the whole time of the study.

The triatomines maintained in the laboratory were fed chicken (*Gallus gallus*) blood once every 15 days. Veterinary medical services of Universidad de los Andes–Biological Sciences bioterium, (mice (*Mus musculus*) blood-chicken (*Gallus gallus*) blood).

**Temperature regimes**

Taking into account the minimum temperature increase expected under climate change and the average temperature of the collection site, three temperatures were chosen for monitoring parasitemia in mice (*Mus musculus*) and metacyclogenesis in insect vectors: 26˚C, 28˚C and 30˚C. These temperatures were adjusted in closed incubators provided with constant air flow, a constant relative humidity of 80% and a 12:12 light: dark photoperiod.

**Parasites**

Taking into account that in Colombia, the most prevalent DTUs of *T. cruzi* are TcI and, less frequently, TcII, blood trypomastigotes of the strains MHOM/CO/04/MG (TcI) and MHOM/BR/53/Y (TcII) were used to conduct the experiments. These reference strains were previously characterized by 24 microsatellite markers and 10 mitochondrial markers (mMLST) and were maintained in successive passages from mouse to mouse every 15 days, following the recommendations of the Institutional Committee for the Care and Use of Laboratory Animals of the University of the Andes.

**Effects of temperature on the life cycle of *R. prolixus***

For each temperature, 4 groups of 30 newly oviposited eggs were formed. Each group was kept in a plastic container (Diameter 10.5 cm, Height 17 cm) with a filter paper base containing faeces of uninfected insects to ensure the presence of the microbiota in the insects to be studied. For each group, the mortality by stage, the average time to hatching, change to fifth instar and change to adult were recorded.

**Effects of temperature on fecundity and egg viability**

For each temperature, 20 females and 20 virgin males obtained from the study of the life cycle were taken. The sample size was estimated for each experiment, taking into account the variability and the average of experimental data previously reported [41, 42]. The weights of the females were recorded before and after feeding as reported elsewhere [41], and reproductive pairs were formed, verifying copulation by observing the "spermatophore casing". This procedure was performed during 3 reproductive cycles, taking each cycle as the 21 days after feeding. The eggs oviposited per cycle were transferred to a Petri box with a filter paper base and were kept there to observe hatching. The *e*-value was calculated, as indicative of the capacity of a female to use the blood ingested towards egg production [43], as was the hatching rate for
each pair in each reproductive cycle. The e-value was calculated as follows:

\[
e - value = \frac{\text{Total number of eggs laid 21 days post meal}}{\text{Initial female weight} \times \text{blood meal weight}} \times 1000
\]

Inoculation of ICR-CD-1 mice

Sixty mice were inoculated intraperitoneally with 0.2 mL of infected blood with each of the selected \( T. cruzi \) strains. Fifteen days post-inoculation, the mice were anaesthetized intraperitoneally with pentobarbital and were then exsanguinated by cardiac puncture. This procedure was previously approved by the research ethics committee and CICUAL of the University of the Andes in ruling 318 of 2014. The concentration of parasites in the blood and their viability were verified by light microscopy using a Neubauer chamber. If necessary, the infected blood was diluted in pathogen-free mouse blood to a concentration of \( 1 \times 10^6 \) parasites/ml.

Effects of temperature on the metacyclogenesis of \( T. cruzi \) in \( R. prolixus \)

For each of the selected strains, 186 fifth-instar nymphs of \( R. prolixus \) were fed heparinized mouse blood at a concentration of \( 1 \times 10^6 \) parasites/mL using an artificial feeder. After feeding, these nymphs were separated into groups of 62 individuals and were subjected to each of the temperatures. (Fig 1)

To establish the possible differences in the time at which metacyclic trypomastigotes were observed in the rectal ampulla, 2 nymphs were dissected per temperature and strain, from the second post-infection day and at intervals of 2 days until day 20. Similarly, to determine if there were differences between temperatures in the number of infectious forms in the rectal ampulla, 2 nymphs were dissected daily per temperature and strain from week 6 to week 8 post-infection. To perform the dissection, a cut was made in the last abdominal segment of the insect, and the rectal ampulla was carefully separated from the rest of the intestinal tract. The contents of the ampulla were macerated and resuspended in 100 mL of physiological saline.

Fig 1. Methodology used to study the effect of temperature on the metacyclogenesis of \( T. cruzi \) in \( R. prolixus \). (A) Inoculation of mice with strains of the parasite. (B) Artificial nymph feeding. (C) Analysis of trypomastigotes in the rectal ampulla and count in the Neubauer chamber.

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(0.9%) at room temperature. Ten microlitres of this solution was used to quantify the number of metacyclic trypomastigotes in the Neubauer chamber.

Ethics statement

All procedures with animals were conducted according to the Guide for the care and use of laboratory animals (8 ed)–National Research Council EEUU and the Institutional Animal Care and Use Committee Guidebook of OLAW. The Universidad de los Andes APRAC, in ruling 318 of 2014, approve all animals protocols used in the present work.

Statistical analysis

All of the statistical analyses were performed in GraphPad Prism 7 software. First, the normality of the data was evaluated using the Shapiro-Wilk test; when the data were not normal, non-parametric tests were used. The Kruskal-Wallis test was used to evaluate the effects of temperature on the life cycle, fecundity and viability of *R. prolixus* eggs. Dunn’s multiple comparisons post hoc test was used to determine which temperatures were responsible for the significant difference found. In addition, a Chi-square test was perform to establish if there were differences in the mortality of the insects subjected to different temperatures. A Kruskal-Wallis test was performed to evaluate if there were statistically significant differences between the temperatures evaluated and the DTUs used. Two tests were necessary, before 20 days post-infection (DPI) and after 35 DPI. Additionally, followed by an analysis of multiple comparisons to determine the groups that showed these differences.

Results

Effects of temperature on the life cycle of *R. prolixus*

The time to hatching, change to fifth instar nymph and change to adult were significantly different between temperatures (Kruskal-Wallis test, *P* < 0.0001). In evaluating hatching time and time to adult, it was found that each temperature differed significantly from the other (Dunn’s test, *P* < 0.0001). However, for the time required to change to fifth instar, significant differences were only found between 26˚C and other temperatures (Dunn’s test, *P* < 0.0001), but not between 28˚C and 30˚C. Compared with 26˚C, the control temperature, the development time from egg to adult was reduced by 13% when the insects were kept at 30˚C and by 9% when the eggs were kept at 28˚C (Fig 2).

No significant differences were found in insect mortality between temperatures (Chi-square *P* = 0.67). In general, mortality was low for all temperatures (6.6% for 30˚C, 5.83% for 28˚C and 4.16% for 26˚C) in comparison with previous studies [44], and was introduced especially

![Fig 2. Effects of temperature on the life cycle of *R. prolixus*. (A) Hatching time by temperature. (B) Time needed to change to fifth instar nymph by temperature. (C) Time to change to adult by temperature. * = P < 0.05, ** = P < 0.01, *** = P < 0.001.](https://doi.org/10.1371/journal.pntd.0006735.g002)
during the change from fifth instar to adult, representing 75% of the mortality at 30˚C, 83.3% at 28˚C and 60% at 26˚C.

Effect of temperature on the fertility and fecundity of females

The *e-value*, as an indicator of fecundity of the females, was not significantly different between reproductive cycles for the insects subjected to 26˚C (Kruskal-Wallis test, *P* = 0.1080) and 28˚C (Kruskal-Wallis test, *P* = 0.7409). At 30˚C, the reproductive cycle did affect the *e-value* (Kruskal-Wallis test, *P* = 0.0001), with the first cycle being shorter and significantly different from the second (Dunn’s multiple comparisons test, *P* = 0.0083) and third cycles (Dunn’s multiple comparisons test, *P* = 0.0001). No significant differences were observed in the *e-values* between temperatures during the first (Kruskal-Wallis test, *P* = 0.2367) and second reproductive cycles (Kruskal-Wallis test, *P* = 0.3118), but there were differences during the third (Kruskal-Wallis test, *P* = 0.0043), with 30˚C significantly different from 26˚C (Dunn’s multiple comparisons test, *P* = 0.0092) and 28˚C (Dunn’s multiple comparisons test, *P* = 0.0092) (Fig 3A).

However, the hatching rate was not affected by either the reproductive cycles (Kruskal-Wallis test, 26˚P = 0.0533, 28˚ P = 0.1687 and 30˚C P = 0.2032) or the temperatures to which the eggs were subjected (Kruskal-Wallis test, 26˚ P = 0.3689, 28˚ P = 0.2511 and 30˚C P = 0.1333) (Fig 3B).

Effects of temperature on the metacyclogenesis of *T. cruzi* in *R. prolixus*

For TcI, the appearance of metacyclic trypomastigotes was observed in the rectal ampulla on the sixth day post-infection at 30˚C, while at 26˚C and 28˚C, it was observed on days 10 and 14, respectively. (Fig 4A). Similarly, for TcII, the appearance of infective forms was observed on day 16 post-infection at 30˚C, while at 28˚C and at 26˚C, metacyclic trypomastigotes were not observed during the first 20 days post-infection. (Fig 4B). All the data evaluated here exhibited a non-parametric distribution. Then, a Kruskal-Wallis test was used to evaluate the difference between temperatures and DTUs. A first analysis carried out up to 20 DPI showed that there was no statistically significant difference in the concentration of parasites between the temperatures analyzed for each DTUs, and the same when making a comparison between both DTUs. Otherwise it happened with the analysis carried out after the 35 DPI, where a statistically significant difference of the concentration of parasites was observed between the temperatures analyzed for both DTUs (*p* < 0.0001). When performing the multiple comparisons test, the temperature of 30˚C presented a clear difference with respect to 26˚ and 28˚, given by

![](https://doi.org/10.1371/journal.pntd.0006735.g003)
an increase in the concentration of parasites. These results were maintained for both DTUs. An analysis between DTUs showed a higher concentration of parasites in the insects infected with the TcI DTU with respect to TcII. These results were verified statistically ($p < 0.0001$), and were maintained throughout the three temperatures.

From weeks 6 to 8 after inoculation, there were significant increases in the numbers of infective forms of TcI and TcII found in the rectal ampulla of insects subjected to 30˚C compared with insects subjected to 26˚C and 28˚C (Fig 4C and 4D). In general, a much greater amount of metacyclic trypomastigotes of the TcI strain compared with the TcII strain was observed. (Fig 4)

**Discussion**

In this study, the effects of increasing temperature, as expected under predictions of climate change, on the life cycle, fecundity and viability of *R. prolixus* eggs and on the development of *T. cruzi* in *R. prolixus* were evaluated. Our results indicate that increasing temperature from 26˚C to 30˚C has effects on the time of development (Fig 2), the fecundity of the insect (Fig 3A) and the development of the parasite (Fig 4). However, no significant effect on egg viability was observed (Fig 3B).

The negative relationship between the temperature and the development time of *R. prolixus* obtained was in agreement with findings previously reported by Clark [45] and Luz et al [46]. This faster development of insects can be a result of increased metabolic rate caused by the increase in temperature [47], a relationship already demonstrated for *R. prolixus* [48]. In Fig 2B and 2C, showing the results for changes to fifth instar and to adult, one can see that the times to such changes in some insects are equal to those for other temperatures. These observations could be explained by the frequency of feeding and the amount of blood ingested by each
specific individual. Although a food source was offered weekly, it was observed that some insects did not feed or did not ingest enough blood to change, which generates an increase in the life cycle duration [18].

The temperature at which the greatest mortality was observed was 30˚C; however, the percentage was similar to that reported by Arevalo et al [41] and was less than observed by Gomes et al [49] for *R. prolixus* under laboratory conditions. In general, the greatest mortality occurred at the change to fifth instar, which has been reported by other investigators for *R. prolixus* [41, 49, 46] and other triatomine species, such as *R. robustus* [50], *T. infestans* [51] and *Meccus picturatus* [52]. Despite of our results, acclimation capability to temperature must be considered. This is a long-step process that might take several years and in our study we were not able to consider that variable. This is relevant in the light of previous studies that show the three main sensitive parameters for Chagas disease transmission (mortality rate, density of vectors and bite rate) [53]. We explored two of them (mortality and density). However, in future studies the bite rate must be considered because with the increase of temperature, there will occur a higher metabolic rate and in consequence a higher bite rate which could drive the increase in the transmission of *T. cruzi*. Therefore, in the future bite rate should be studied.

Although the presence of Spermatophore casings was checked to verify copulation, the number of copulations per couple was not taken into account. Therefore, it is possible that both the differences observed in the e-value between reproductive cycles for 30˚C and the low hatching rate observed during the first reproductive cycle at this temperature can be explained by a difference in the number of copulations. For *T. brasiliensis*, it is known that females that have multiple copulations produce more eggs with a greater percentage of fertility than females that copulate only once [54]. However, it would be interesting to study fluctuations in fecundity during more reproductive cycles to establish if the differences observed in the third reproductive cycle are maintained or if they are only the result of the number of copulations. If these differences were maintained between reproductive cycles, it would be possible to think that at 30˚C, the number of eggs laid per female would increase, and as a consequence, there would be a greater number of insects per reproductive cycle that would be available to transmit *T. cruzi*. This has been studied by Schilman and Lazzari in 2004 [55], where they found that females oviposit across a range of temperatures from 22 to 33˚C with a peak at 25–26˚C in accordance with our findings. Nevertheless, they cannot discern whether *R. prolixus* females actively choose certain oviposition substrates according to temperature or whether they oviposit where they find themselves. These results, both in the life cycle and in the fecundity and fertility of the insects, suggest that the expected temperature increase under climate change could increase the insect density in the palms of *A. butyacea*. This possibility is not very different from field reports that state that the density of insects in these palms is higher in summer seasons than in rainy seasons, when the temperature decreases [39].

The time at which infectious forms of both TcI and TcII are observed in the rectal ampulla of *R. prolixus* is shorter at 30˚C than at the other temperatures. These results are consistent with previous data reported for *Triatoma infestans*, where metacyclic trypomastigotes were observed to be faster at 28˚C than at 20˚C [56]. Likewise, the number of infective forms observed for TcI from week 6 to week 8 was greater at 30˚C than at the other temperatures. This relationship between the temperature and reproduction of the parasite has been previously reported under *in vitro* conditions for the epimastigote stage [43, 57]; however, it must be taken into account that under *in vitro* conditions, the parasite is not subject to the immune factors and the microbiota of the insect [58]; therefore, it is difficult to extrapolate and compare the *in vitro* results with the results obtained in this study.

The amount of metacyclic trypomastigotes observed in the rectal ampulla of *R. prolixus* was markedly higher for TcI than for TcII. This difference is likely due to the presence of
trypanolytic factors in the haemolymph of \textit{R. prolixus} that differentially affect TcII but not TcI, which is the main reason why this species of vector is considered to lack the capacity to transmit said DTU \cite{59, 60}. However, it is interesting to note that the temperature may be affecting this interaction, since at 30˚C, an increase in the number of infective forms was observed for TcII (Fig 4B–4D). Therefore, in the future, it would be important to evaluate whether this increase in the number of infective forms, due to temperature, could enhance the capacity of \textit{R. prolixus} to transmit \textit{T. cruzi} II.

In conclusion, this study showed that as temperature increases (26 to 30˚C), there is a more rapid appearance and an increase in the number of infective forms of \textit{T. cruzi} in \textit{R. prolixus}, along with a significant decrease in the development time of said vector. These results could suggest that under the effects of climate change, the probability of infection with \textit{T. cruzi} could increase. However, it is necessary to study the effects of more climatic and ecological factors and the effects of such factors on parasite-vector interactions to predict the future of Chagas disease with better accuracy.

**Supporting information**

S1 Database. Includes the life cycle, hatching rate, fertility and parasite counting for each temperature.

(XLSX)

**Author Contributions**

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**References**

1. IPCC: Climate Change 2014: Synthesis Report. Contribution of Working Groups I, II and III to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change [Core Writing Team, R.K. Pachauri and L.A. Meyer (eds.)] Geneva: IPCC; 2014. Available: http://www.ipcc.ch/pdf/assessmentreport/ar5/syr/SYR_AR5_FINAL_full_wcover.pdf. Accessed 18 Junio 2017.

2. Githeko AK, Lindsay SW, Confalonieri UE, Patz JA: Climate change and vector-borne diseases: a regional analysis. Bulletin of the World Health Organization. 2000; 78:9.

3. McMichael AJ, Haines A. Global climate change: the potential effects on health. BMJ: British Medical Journal. 1997; 315(7111):805–809. PMID: 9345177
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4. Yu M, Mengersen K, Dale P, Ye X, Guo Y, Turner L, Wang X, Bi Y, Mcbride WJH, Mackenzie JS, Tong S. Projecting future transmission of malaria under climate change scenarios: Challenges and research needs. Crit Rev Environ Sci Technol. 2014; 45(7): 777–811. https://doi.org/10.1080/10643389.2013.852392

5. Morin CW, Comrie AC, Ernst K. EHP–Climate and Dengue Transmission: Evidence and Implications. Environ Health Perspect. 2013; 121(11–12): 1264–1272. https://doi.org/10.1289/ehp.1306556 PMID: 24058050

6. Butterworth MK, Morin CW, Comrie AC. An analysis of the potential impact of climate change on dengue transmission in the southeastern United States. Environ Health Perspect. 2017; 125:579–585. https://doi.org/10.1289/EHP218 PMID: 27713106

7. González C, Wang O, Strutz SE, González-Salazar C, Sánchez-Cordero V, Sarkar S. Climate change and risk of leishmaniasis in North America: Predictions from ecological niche models of vector and reservoir species. PLoS Negl Trop Dis. 2010; 4(1). https://doi.org/10.1371/journal.pntd.0000585 PMID: 20098485

8. Peterson AT, Campbell LP, Moo-Llanes DA, Travi B, Gonzalez C, Ferro MC, Ferreira GEM, Bandão-Filho SP, Cuppollo E, Ramsey J, Leffer AMC, Pech-May A, Shaw JJ. Influences of climate change on the potential distribution of Lutzomyia longipalpis sensu lato (Psychodidae: Phlebotominae). Int. J. Parasitol. 2017; 47(10–11): 667–674 https://doi.org/10.1016/j.ijparasit.2017.04.007 PMID: 28868326

9. Stratman-Thomas WK. The Influence of Temperature on Plasmodium Vivax. Am J Trop Med Hyg. 1940; S1-20(5): 703–715. https://doi.org/10.4269/ajtmh.1940.s1-20.703

10. Bayoh MN, Lindsay SW. Effect of temperature on the development of the aquatic stages of Anopheles gambiae sensu stricto (Diptera: Culicidae). Bull. Entomol. Res. 2003; 93(5): 375–381. https://doi.org/10.1017/S0007469103000490

11. Bayoh MN, Lindsay SW. Effect of temperature on the development of the aquatic stages of Anopheles gambiae sensu stricto (Diptera: Culicidae). Bull. Entomol. Res. 2003; 93(5): 375–381. https://doi.org/10.1017/S0007469103000490

12. Zell R. Global climate change and the emergence/re-emergence of infectious diseases. Int. J. Med. Microbiol. 2004; 293, (Suppl. 37): 16–26. https://doi.org/10.1016/S1433-1128(04)80005-6

13. Guhl F. Chagas disease in Andean countries. Mem. Inst. Oswaldo Cruz. 2007; 102(Suppl. I): 29–37.

14. World Health Organization. La enfermedad de Chagas (Tripanosomiasis Americana). 2017. http://www.who.int/tdr/publications/research_priorities/en/

15. World Health Organization: Research Priorities for Chagas Disease, Human African Trypanosomiasis and Leishmaniasis. 2012, World Health Organization, http://www.who.int/tdr/publications/research_priorities/en/

16. Vallejo GA, Guhl F, Schaub GA. Triatomaene-Trypanosoma cruzi/T. rangeli: Vector-parasite interactions. Acta Trop. 2009; 110(2–3): 137–147. https://doi.org/10.1016/j.actatropica.2008.10.001 PMID: 18992212

17. Guhl F, Ramírez JD. Retrospective molecular integrated epidemiology of Chagas disease in Colombia. Infect Genet Evol. 2013; 20: 148–154. https://doi.org/10.1016/j.meegid.2013.08.028 PMID: 24012949

18. Buxton MA. The biology of a blood-sucking bug, Rhodnius prolixus. Ecological Entomology. 1930; 78(2):227–256. https://doi.org/10.1111/j.1365-2311.1930.tb00385.x/full

19. Moncayo A, Silveira AC. Current epidemiological trends of Chagas disease in Latin America and future challenges: Epidemiology, surveillance, and health policies. Mem. Inst. Oswaldo Cruz. 2009; 104(1): 17–30.

20. Hernández C, Vera MJ, Cucunuá Z, et al. High-Resolution Molecular Typing of Trypanosoma cruzi in 2 Large Outbreaks of Acute Chagas Disease in Colombia. J Infect Dis. 2016; 214(8):1252–1255. https://doi.org/10.1093/infdis/jiw360 PMID: 27511897

21. Herrera L, D’Andrea PS, Xavier SCC, Mangia RH, Fernandez O, Jansen AM. Trypanosoma cruzi infection in wild mammals of the National Park ‘Serra da Capivara’and its surroundings (Piauí, Brazil), an area endemic for Chagas disease. Trans R Soc Trop Med Hyg, 2005; 99(5), 379–388. https://doi.org/10.1016/j.trstmh.2004.07.006 PMID: 15780345

22. Roque ALR, Xavier SC, da Rocha MG, Duarte ACM, D’Andrea PS, Jansen AM. Trypanosoma cruzi transmission cycle among wild and domestic mammals in three areas of orally transmitted Chagas disease outbreaks. Am J Trop Med Hyg. 2008; 79(5):742–9. PMID: 18981516

23. Gómez Nuñez JC, Fernández J. La colonia de Rhodnius prolixus en el Instituto Venezolano de Investigaciones Científicas. Bol Dir Malaria y San Amb. 1963; 3: 132137.

24. Galliard H. Recherches sur les réduvides hématophages Rhodnius et Triatoma. IV—Organes génitaux femelles. Ann Parasitol Hum Comp. 1935; 13: 497–527.

25. Okasha AYK. Effects of high temperature in Rhodnius prolixus (Stål). Nature. 1964; 204: 1221–1222.
26. Carcavallo RU, Tonn RJ, Carrasco B. Distribución de triatominos en Venezuela, (Hemiptera, Reduviidae). Actualización por entidades y zonas biogeográficas. Bol. Dir. Malaria. San. Amb. 1977; 17 (1):53–65.

27. Carcavallo RU, Tonn RJ, Ortega R, Betancourt P, Carraquero B. Notas sobre la biología, ecología y distribución geográfica del *Rhodnius prolixus* Stal, 1859 (Hemiptera, Reduviidae) Bol Dir Malaria San Amb. 1978; 18: 175–198.

28. Lent H, Wygodzinsky P. Revision of Triatominae (Hemiptera, Reduviidae), and their significance as vectors of Chagas’ disease. Bull. Am. Mus. Nat. Hist. 1979; 163(3):123–520.

29. Rassi Jr, Rassi A, Marín-Neto JA. Chagas Disease. The Lancet. 2010; 375(9723):17–23. https://doi.org/10.1016/S0140-6736(10)60061-X

30. Zingales B, Miles MA, Campbell DA, Tibayrenc M, Macedo AM, Teixeira MMG, Schijman AG, Llewellyn MS, Lages-Silva E, Machado CR, Andrade SG, Sturm NR. The revised *Trypanosoma cruzi* subspecific nomenclature: rationale, epidemiological relevance and research applications. Infect Genet Evol. 2012 Mar; 12(2):240–53. https://doi.org/10.1016/j.meegid.2011.12.009 PMID: 22226704

31. Marcili A, Lima L, Cavazzana M, Junqueira AC, Veludo HH, Da Silva FM, Campaner M, Paiva F, Nunes VL, Teixeira MM. A new genotype of *Trypanosoma cruzi* associated with bats evidenced by phylogenetic analyses using SSU rDNA, cytochrome b and Histone H2B genes and genotyping based on ITS1 rDNA. Parasitology. 2009; 136(6):641–55. https://doi.org/10.1017/S0031182009005861 PMID: 19368741

32. Ramirez JD, Tapia-Calle G, Muñoz-Cruz G, Poveda C, Rendón LM, Hincapié E, & Guhl F. Trypanosome species in neo-tropical bats, biological, evolutionary and epidemiological implications. Infect. Genet. Evol. 2014; 22: 250–256. https://doi.org/10.1016/j.meegid.2013.06.022 PMID: 23831017

33. Medone P, Ceccarelli S, Parham PE, Figuera A, Rabinovich JE. The impact of climate change on the geographical distribution of two vectors of Chagas disease: implications for the force of infection. Philos Trans R Soc Lond B Biol Sci. 2015; 370(1665). https://doi.org/10.1098/rstb.2013.0560 PMID: 25688019

34. Escolar LE, Romero-Alvarez D, Leon R, Lepe-Lopez MA, Craft ME, Borbor-Cordova MJ, Svenning JC. Declining prevalence of Disease Vectors Under Climate Change. Sci Rep. 2016; 6(39150). https://doi.org/10.1038/srep39150 PMID: 27982119

35. Costa J, Dornak LL, Almeida CE, Peterson AT. Distribution potential of the *Triatoma brasiliensis* species complex at present and under scenarios of future climate condition. Parasit Vectors. 2014; 7(238). https://doi.org/10.1186/1756-3305-7-238 PMID: 24886587

36. Garza M, Feria Arroyo TP, Casillas EA, Sanchez-Cordero V, Rivaldi CL, Sarka S. Projected future distributions of vectors of *Trypanosoma cruzi* in North America under climate change scenarios. PLoS Negl Trop Dis. 2014; 8(5):e2818. https://doi.org/10.1371/journal.pntd.0002818 PMID: 24831117

37. Cordovez JM, Rendon LM, Gonzalez C, Guhl F. Using the basic reproductive number to assess the effects of climate change in the risk of Chagas disease transmission in Colombia. Acta Trop. 2014; 129:74–82. https://doi.org/10.1016/j.actatropica.2013.10.003

38. Tapia-Garay V, Figueroa DP, Maldonado A, et al. Assessing the risk zones of Chagas’ disease in Chile, in a world marked by global climatic change. Mem Inst Oswaldo Cruz. 2018; 113(3):283–288. https://doi.org/10.1590/0074-02760170186 PMID: 29211105

39. Rendón LM, Guhl F, Cordovez JM, Erazo D. New scenarios of *Trypanosoma cruzi* transmission in the Orinoco region of Colombia. Memorias do Instituto Oswaldo Cruz. 2015; 110(3):283–288. https://doi.org/10.1590/0074-02760140403 PMID: 25830543

40. Urbano P, Poveda C, Molina J. Effect of the physiology of *Attalea butyracea* (Arecidae) on population density and age distribution of *Rhodnius prolixus* (Triatominae). Parasites & Vectors. 2015; 8:199. https://doi.org/10.1186/s13071-015-0813-6 PMID: 25889617

41. Ariávalo A, Carranza JC, Guhl F, Clavijo JA, Vallejo GA. Comparación del ciclo de vida de *Rhodnius colombiensis* (Moreno, Jurberg & Galvão, 1999) y *Rhodnius prolixus* Stal, 1872 (Hemiptera, Reduviidae, Triatominae) en condiciones de laboratorio. Biomédica. 2007; 27: 119–129.

42. Rueda NK. Evaluación de la competencia vectorial de Rhodnius colombiensis, R. pallescens, R. prolbus, R. robustus y T. maculata a través de la metaciglicogénesis en vivo de Trypanosoma cruzi I y T. cruzi II (Tesis de maestría).

43. Fellet MR, Lorenzo MG, Elliot SL, Carrasco D, Guameri AA. Effects of infection by *Trypanosoma cruzi* and *Trypanosoma rangeli* on the reproductive performance of the vector *Rhodnius prolixus*. PLoS ONE. 2014; 9(8):26–32. https://doi.org/10.1371/journal.pone.0109255 PMID: 25136800

44. Chaves LF, Hernandez MJ, Revilla TA, Rodriguez DJ, Rabinovich JE. Mortality profiles of *Rhodnius prolixus* (Heteroptera: Reduviidae), vector of Chagas disease. Acta Trop. 2004 Oct; 92(2):119–25. https://doi.org/10.1016/j.actatropica.2004.06.003 PMID: 15350963
45. Clark N. The effect of temperature and humidity upon the eggs of the bug, Rhodnius prolixus (Heteroptera, Reduviidae). J Anim Ecol. 1935; 4: 82–87.

46. Luz C, Fargues J, Grunewald J. Development of Rhodnius prolixus (Hemiptera, Reduviidae) under Constant and Cyclic Conditions of Temperature and Humidity. Mem. Inst. Oswaldo Cruz. 1999; 94(3): 403–409. https://doi.org/10.1590/S0074-02761999000300022 PMID: 10348991

47. Ratte HT. Temperature and Insect Development. In: Hoffmann KH, editor. Environmental Physiology and Biochemistry of Insects. Berlin: Springer Berlin; 1984. 33–66. https://doi.org/10.1007/978-3-642-70020-0_2

48. Okasha A. Effects of Sub-Lethal High Temperature on An Insect, Rhodnius Prolixus (Stål.) III. Metabolic changes and their bearing on the cessation and delay of moulting. J. Exp. Biol. 1968; 48: 475–486.

49. Gomes JE, Azambuja P, Garcia ES. Comparative studies on the growth and reproductive performances of Rhodnius prolixus reared on different blood sources. Mem Inst Oswaldo Cruz. 1990; 85: 299–304. PMID: 2134703

50. Rocha DS, Jurberg J, Carcavallo Presgrave OAF, Cunha V, Galvão C. Influência da temperatura e umidade no desenvolvimento ninhal de Rhodnius robustus. Rev. Saúde Pública. 2001; 35(4): 400–406. https://doi.org/10.1590/S0034-89102001000400011

51. Rabinovich JE. Vital statistics of Triatominae (Hemiptera, Reduviidae) under laboratory conditions. I. Triatoma infestans Klug. J Med Entomol. 1972; 9(4):351–370. https://doi.org/10.1093/jmedent/9.4.351 PMID: 4559948

52. Martínez-Ibarra JA, Novelo M, Hernández MR, Grant Y. Influence of the blood meal source on the biology of Meccus picturatus Usinger 1939 (Hemiptera: Reduviidae: Triatominae) under laboratory conditions. Mem. Inst. Oswaldo Cruz. 2003; 98(2): 227–232. https://doi.org/10.1590/S0074-02762003000200011 PMID: 12764439

53. Canals M, Cáceres D, Alvarado S, Canals A, Cattan PE. Modeling Chagas disease in Chile: From vector to congenital transmission. Biosystems. 2017 Jun–Jul; 156–157:63–71. https://doi.org/10.1016/j.biosystems.2017.04.004 PMID: 28435013

54. Daflon-Teixeira NF, Carvalho-Costa FA, Chiang RG, Lima MM. Influence of blood meal and mating in reproduction patterns of Triatoma brasiliensis females (Hemiptera: Reduviidae), under laboratory conditions. Mem. Inst. Oswaldo Cruz. 2009; 104(7): 1031–1034. https://doi.org/10.1590/S0074-02762009000700016 PMID: 20027473

55. Schilman PE, Lazzari CR. Temperature preference in Rhodnius prolixus, effects and possible consequences. Acta Trop. 2004 Mar; 90(1):115–22. PMID: 14739030

56. Asin S, Catala S. Development of Trypanosoma cruzi in Triatoma infestans: influence of temperature and blood consumption. J Parasitol. 1995; 81(1):1–7. PMID: 7876960

57. Elliot SL, Rodrigues JdeO, Lorenzo MG, Martins-Filho OA, Guarneri AA. Trypanosoma cruzi, Etiologic Agent of Chagas Disease, Is Virulent to Its Triatomine Vector Rhodnius prolixus in a Temperature-Dependent Manner. PLoS Negl Trop Dis. 2015; 20: 9(3):e0003646. https://doi.org/10.1371/journal.pntd.0003646

58. Kollien AH, & Schaub GA. The development of Trypanosoma cruzi in triatominae. Parasitol. Today. 2000; 16(9):381–387. https://doi.org/10.1016/S0169-4758(00)01724-5 PMID: 10951597

59. Zabala D, Carranza JC, Urrea DA, Guhl F, Jaramillo N, Teixeira MM, Vallejo GA. Respuesta inmune diferencial de triatominos contra Trypanosoma cruzi y T. rangeli. Rev. Asoc. Col. Cienc. (Col.), 2011; 23:134–143.

60. Cruz-Saaavedra L, Muñoz M, León C, Patarroyo MA, Arevalo G, et al. Purification of Trypanosoma cruzi metacyclic trypomastigotes by ion exchange chromatography in sepharose-DEAE, a novel methodology for host-pathogen interaction studies. J. Micrcol. Methods 2017; 142: 27–32.