

ORIGINAL RESEARCH

WHF IASC Roadmap on Chagas Disease

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Background: Chagas Disease is a neglected tropical disease caused by the protozoan *Trypanosoma cruzi*, with some of the most serious manifestations affecting the cardiovascular system. It is a chronic, stigmatizing condition, closely associated with poverty and affecting close to 6 million people globally. Although historically the disease was limited to endemic areas of Latin America recent years have seen an increasing global spread. In addition to the morbidity and mortality associated with the disease, the social and economic burdens on individuals and society are substantial. Often called the 'silent killer', Chagas disease is characterized by a long, asymptomatic phase in affected individuals. Approximately 30% then go on develop chronic Chagas cardiomyopathy and other serious cardiac complications such as stroke, rhythm disturbances and severe heart failure.

Methods: In a collaboration of the World Health Federation (WHF) and the Inter-American Society of Cardiology (IASC) a writing group consisting of 20 diverse experts on Chagas disease (CD) was convened. The group provided up to date expert knowledge based on their area of expertise. An extensive review of the literature describing obstacles to diagnosis and treatment

of CD along with proposed solutions was conducted. A survey was sent to all WHF Members and, using snowball sampling to widen the consultation, to a variety of health care professionals working in the CD global health community. The results were analyzed, open comments were reviewed and consolidated, and the findings were incorporated into this document, thus ensuring a consensus representation.

Results: The WHF IASC Roadmap on Chagas Disease offers a comprehensive summary of current knowledge on prevention, diagnosis and management of the disease. In providing an analysis of 'roadblocks' in access to comprehensive care for Chagas disease patients, the document serves as a framework from which strategies for implementation such as national plans can be formulated. Several dimensions are considered in the analysis: healthcare system capabilities, governance, financing, community awareness and advocacy.

Conclusion: The WHF IASC Roadmap proposes strategies and evidence-based solutions for healthcare professionals, health authorities and governments to help overcome the barriers to comprehensive care for Chagas disease patients. This roadmap describes an ideal patient care pathway, and explores the roadblocks along the way, offering potential solutions based on available research and examples in practice. It represents a call to action to decision-makers and health care professionals to step up efforts to eradicate Chagas disease.

Keywords: Chagas disease; cardiomyopathy; neglected tropical disease; heart failure

Introduction

In 2014, the World Heart Federation (WHF) launched an initiative to develop a series of Roadmaps, which identify potential roadblocks on the pathway to effective prevention, detection, and management of cardiovascular disease (CVD), along with evidence-based solutions to overcome them. The resulting documents act as a tool to help turn strategic intent into action plans for integrating the latest knowledge and evidence into national plans for optimal management of cardiac diseases.

The Roadmap publications have become the cornerstone of WHF activities as implementation resources, and they guide initiatives to support heart health globally, translating science into policy and influencing agencies, governments, and policymakers alike. They aim to provide a framework for countries that wish to develop or update national initiatives and programmes tackling cardiac diseases.

Despite more than a century having elapsed since the discovery of CD it remains a major public health concern with significant social and economic burdens in both Latin America and increasingly on a global scale. CD, like other neglected tropical diseases (NTD), is a chronic, stigmatizing condition, closely associated with poverty. Despite being infectious in origin, the predominant and most serious chronic manifestations of CD affect the cardiovascular system. In keeping with WHF's mission to deliver cardiac health for all as a



Figure 1: An integrated approach to patient care.

fundamental human right and crucial element of global health justice, we have made the elimination of this disease one of our priorities.

The WHF Roadmap on CD is a document created for all stakeholders to provide an integrated approach to patient care (**Figure 1**). Its goal is to present a framework for prevention and control efforts at a national, regional, and global level that balances the feasibility, acceptance, and accessibility of solutions presented for local implementation.

The Roadmap on CD sheds light on the barriers to patients accessing care, and proposes evidence-based practical solutions for healthcare professionals, health authorities and governments to help overcome these barriers. As part of the WHF Roadmap series, it complements existing Roadmaps on rheumatic heart disease [1], tobacco control [2], hypertension [3], the use of secondary prevention for CVD [4], atrial fibrillation [5], heart failure, and prevention of cardiovascular disease among people living with diabetes [6]. **Figure 2** outlines the design and methodology of the WHF Roadmap series.

Expert writing group

In 2019, the WHF and the Inter-American Society of Cardiology (IASC) convened a Roadmap writing group in joint partnership, consisting of 20 diverse CD experts: clinicians, allied health professionals, health systems experts, and researchers representing many of the important stakeholders working on this disease. Considering CD not only as an infectious disease but as a chronic cardiac disease requiring comprehensive care and management across the lifespan, this Roadmap provides an essential framework for all involved in the planning, organization, patient management and implementation of approaches to CD. It includes an integrated approach from a broad group of stakeholders including healthcare professionals, patients, academic and research institutions, and policy-makers. Recommendations are identified from the standpoint of CD experts as well as those living with CD.

Methodology

To ensure a best practice approach and a consensus document, the ‘WHF IASC Chagas Disease Roadmap’ was developed through the review of published guidelines and research papers. An extensive review of the literature describing obstacles to diagnosis and treatment of CD along with proposed solutions was conducted. A survey was sent to all WHF Members and, using snowball sampling to widen the consultation, to a variety of health care professionals working in the CD global health community. The results were analyzed, open comments were reviewed and consolidated, and the findings were incorporated into this document, thus ensuring a consensus representation (**Figure 2**). The ‘WHF IASC Chagas Disease Roadmap’ can be used as a springboard to initiate a *call for action* and prescribe measurable steps towards a common goal, at national and international levels.

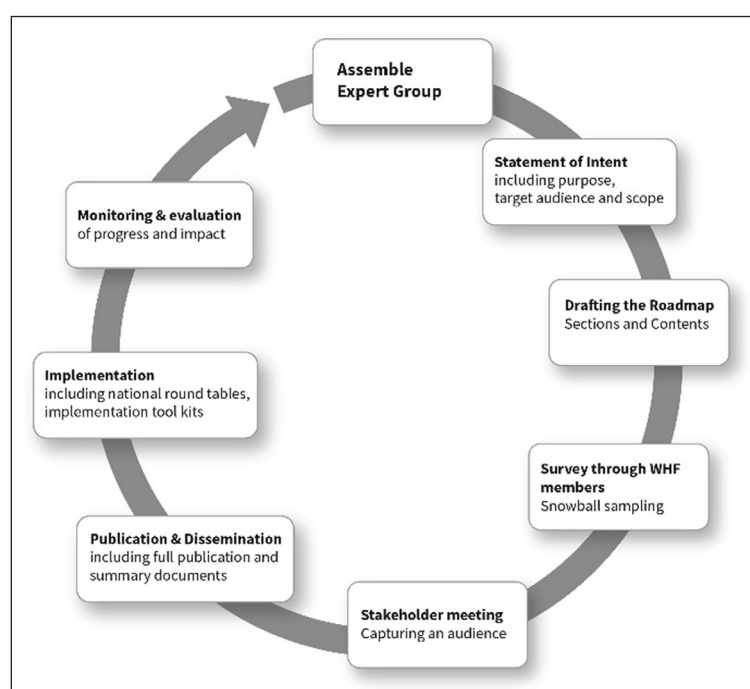


Figure 2: WHF Roadmaps design and methodology.

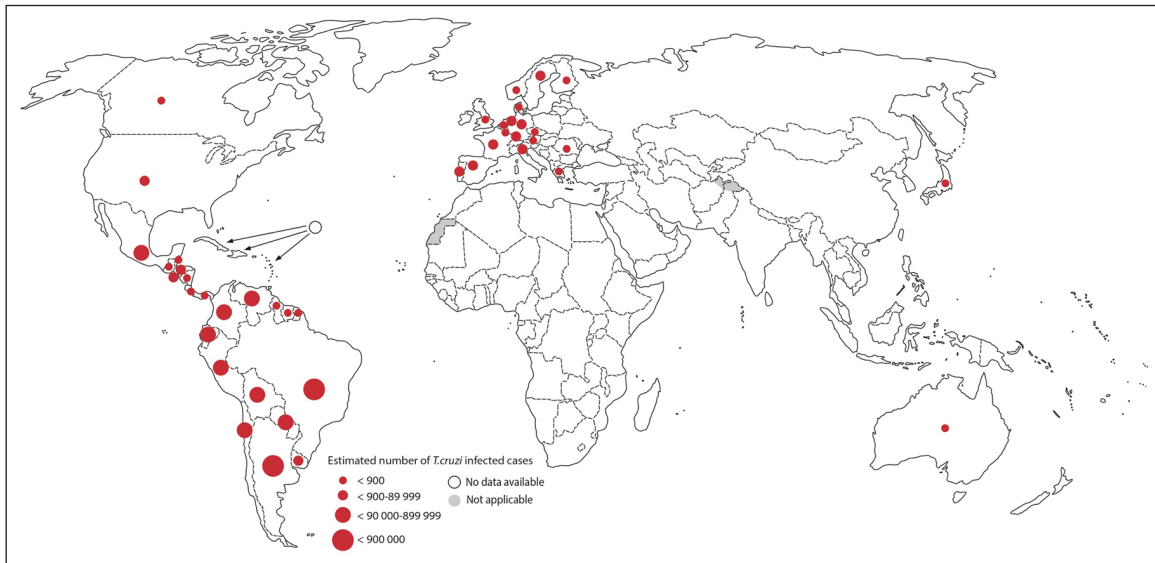


Figure 3: Global distribution of Chagas disease cases, based on official estimates, 2006–2015 [9].

What is Chagas Disease?

CD is a multi-systemic disorder that can affect the cardiovascular, digestive and central nervous systems [7]. CD is caused by *Trypanosoma cruzi*, a hemoflagellate parasite that is transmitted through various species of hematophagous reduviid insects ('kissing bugs') whose habitat ranges from Argentina and Chile to the southern half of the United States. The parasite can also be transmitted transplacentally, as well as through infected blood transfusions or organ donations, laboratory accidents, needle sharing among intravenous drug users (IVDU), and orally through food and drink contaminated with triatomines or their feces.

CD is endemic to all continental Latin American countries. The most recent estimates from WHO (2015) indicate a prevalence of 5.7 million in endemic countries, mostly concentrated in Argentina (1,505,235 cases, prevalence 3.6%), Brazil (1,156,821, 0.6%), Mexico (876,458, 0.7%) Bolivia (607,186, the highest prevalence, 6.1%), Colombia (437,960, 0.9%) and Venezuela (310,000, 1.1%) [8].

An estimated 10–14,000 patients with Chagas die each year, and given its poor prognosis, chronic Chagas cardiomyopathy (CCC) is associated with substantial morbidity that correlates with an increased economic burden for individuals and communities [9, 10]. Because CD often manifests 15–30 years after childhood infection [7], the impact on earning potential for affected individuals is high, which is catastrophic for socially disadvantaged individuals and their families, among whom CD is disproportionately concentrated.

Estimates of CD prevalence have decreased markedly in the last decades, from 17 million in 1980 to less than 6 million in 2010, which has been attributed to coordinated programmes aiming to interrupt transmission of CD [11]. Indeed, as vector and blood transfusion transmission has been interrupted in many countries (Uruguay, Chile, Brazil, Paraguay, Honduras, Nicaragua, Belize, and some states of Argentina, Bolivia, Peru, Colombia, El Salvador, Guatemala, Panamá, Costa Rica), the number of new cases has dropped substantially [12]. However, there are still some areas where vector-based transmission occurs, possibly related to the existence of high-density vector infestation and delayed implementation of vector-control interventions. Most of these new vector-transmitted cases have been reported in Bolivia and the Gran Chaco region, as well as in Central American countries such as Guatemala and El Salvador [8]. **Figure 3** shows the global distribution of CD cases, based on official estimates, 2006–2015.

In parallel to this trend of decreasing incidence, several additional phenomena have impacted the changing epidemiological profile of CD over the last two decades:

Globalization: Because of political and economic pressures, large numbers of infected individuals have migrated from endemic countries to non-endemic areas, including Europe, Japan, Australia, and the United States. CD has thus become a global health concern. It is estimated that more than 300,000 people with CD live in the USA [13], and another 42,000 reside in Spain [14]. In these countries, physicians are generally unaware of the disease and therefore do not recognize (**Figure 3**) or treat it, and health system responses to care for affected people have been slow to materialize. In addition, the frequently tenuous social and economic conditions of migrants complicate access to care.

Aging: In most places where vector transmission has been interrupted, the majority of individuals with CD are now adults or older persons, and CCC often co-exists with other cardiac risk factors and diseases such as diabetes, coronary artery disease and hypertensive cardiomyopathy [15]. In Brazil, the burden of morbidity and mortality from CD is highest among males, the elderly, and in those Brazilian states encompassing important endemic areas for vector transmission in the past [16]. The natural history of CD in the elderly has not been completely described, although we know that Chagas cardiomyopathy remains a strong predictor of higher risk of death [15].

Congenital transmission: While other routes of transmission are decreasing in importance due to successful interventions interrupting vector and blood transfusion routes, congenital transmission has become proportionately more relevant. It now accounts for about one-third of new infections [8], representing the major mode of transmission in non-endemic areas.

Oral transmission: Oral transmission is now more frequently recognized, especially in the Amazon region and the subtropical Andes (Venezuela, Colombia and Ecuador) [17]. It is now the primary mechanism of acute cases in the Brazilian Amazon and Venezuela [18, 19], and is characterized by higher mortality during the acute phase than the vector-mediated acute disease [20].

Underreporting: Most estimates of the burden of CD are based on official mortality figures. However, since CD is a neglected disease that is often not recognized by physicians, it is not included as the cause of death in many cases [21], leading to marked under-recognition of the disease burden [22].

Natural History of the Disease

Most CD patients remain asymptomatic throughout life. As shown in **Figure 4**, approximately 30% progress to clinical forms of the disease, often after a silent phase of many years, and can go on to develop severe clinical complications, mainly cardiovascular, that may lead to incapacity and death [23].

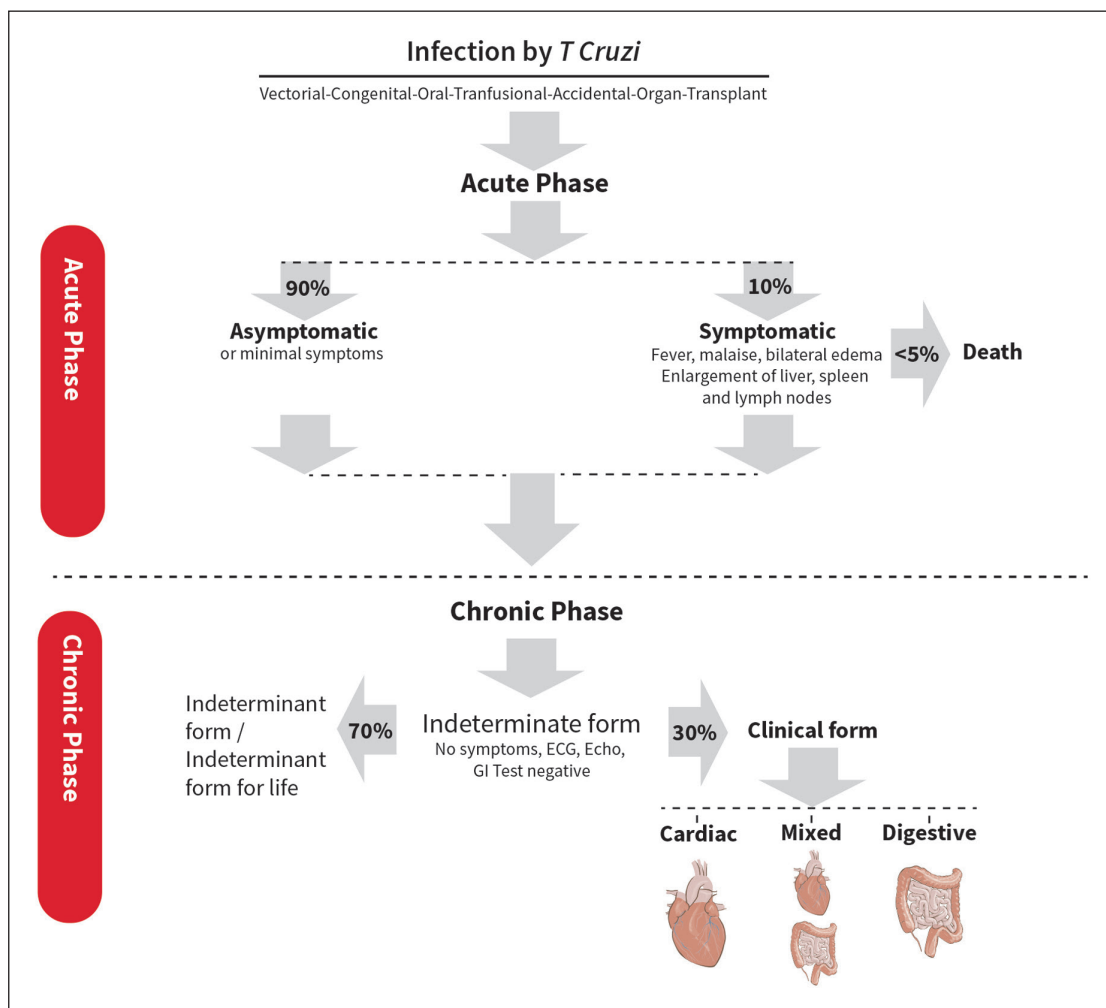


Figure 4: Phases of Chagas disease.

Following an incubation period that varies from seven to fifteen days for vector-based transmission and around thirty to forty days in the case of transfusion-related transmission, the infection passes through two distinct phases. The initial acute phase is characterized by high levels of parasitemia, in which the parasite assumes a trypomastigote form and invades the liver, gut, spleen, lymphatic ganglia, central nervous system, skeletal and cardiac muscles. After the acute phase, *T. cruzi* assumes the dividing form (amastigote), unleashing a local inflammatory reaction [8]. The acute phase generally lasts for one to two months and is followed by an asymptomatic indeterminate phase, during which time no clinical manifestations are observed. In the third of patients that go on to develop chronic CD, the parasite and the immune response cause damage to end organs (**Figure 4**) [24].

Clinical manifestations

Acute Chagas

It is likely that many cases are asymptomatic, or that acutely infected individuals have nonspecific symptoms that do not prompt them to seek medical care, including: fever, fatigue, rash, anorexia, headache, body aches, diarrhea, and vomiting. Among patients who have a clinical evaluation during this phase, marked parasitemia is noted, and clinical signs include hepatosplenomegaly, generalized or local edema (in limbs or face), and lymphadenopathy. In vector-transmitted CD, some specific signs may be present: inflammation at the inoculation site (inoculation chancre), and Romaña's sign, a unilateral bi-palpebral painless edema [25, 26, 27]. Severe acute disease occurs in less than 1–5% of vector-transmitted cases, and may present hemorrhagic manifestations, jaundice, myocarditis, pericardial effusion, tachycardia, arrhythmias, atrioventricular block, and, in a small percentage, meningoencephalitis [28]. Severe acute disease also carries a risk of mortality between 0.2–0.5%. The acute phase of orally transmitted CD is associated with higher risk of a severe presentation, as is also the case in immunosuppressed patients, such as patients taking chemotherapy or those with advanced HIV infection [29]. In the case of vertical transmission, the majority of affected newborns remain asymptomatic; however, at least 10% present with hepatosplenomegaly, sepsis, respiratory failure, low birth weight, or premature delivery [24]. Apart from these specific exceptions, in the majority of cases, symptoms related to the acute phase resolve spontaneously and patients remain chronically infected if untreated. A high index of suspicion for CD is therefore necessary to be able to make an early diagnosis and initiate treatment in order to avoid progression to the chronic stage of the disease, which results in end organ damage.

Reactivated Chagas disease

Pharmacological immunosuppression or HIV/AIDS, particularly with CD4 counts <200, increases the risk of reactivation in patients with chronic *T. cruzi* infection [30, 31]. The overall observed prevalence of reactivation in the absence of prophylactic treatment is 28% in transplant patients and 36–40% in people co-infected with HIV/AIDs [32].

In immunocompromised patients, the most frequent manifestations of acute or reactivated CD are prolonged febrile syndrome and neurological manifestations (meningoencephalitis and/or cerebral granuloma). Also frequent are cardiac manifestations (myocarditis, arrhythmias, and cardiac insufficiency). Dermatologic lesions may be observed in transplant patients, including acute panniculitis in the arms, legs and abdomen [33, 34].

Indeterminate phase

After resolution of the initial acute illness, patients generally pass into a phase of CD in which there are no end organ manifestations of the illness in the setting of positive serology. While this often asymptomatic stage persists for most infected patients, some will pass on to the chronic stage of the disease.

Chronic Chagas disease

Cardiac manifestations (chronic Chagas cardiomyopathy)

The 30% of infected patients who progress from the indeterminate phase of the disease develop manifest damage to organs, particularly the heart or viscera [23]. Patients may suffer sudden cardiac death, thromboembolic phenomena, syncope, and congestive heart failure (CHF). Signs and symptoms of cardiac involvement primarily include electrical and mechanical alterations; sinus bradycardia, atrial and ventricular arrhythmias; atrioventricular and intraventricular conduction disorders, such as right bundle-branch block and/or left anterior fascicular block [35]; and ST-T changes. Cardiac imaging demonstrates regional wall-motion abnormalities, apical aneurysms, mural thrombi with embolic potential, and dilated cardiomyopathy with reduced LVEF [36] (**Figure 5**).

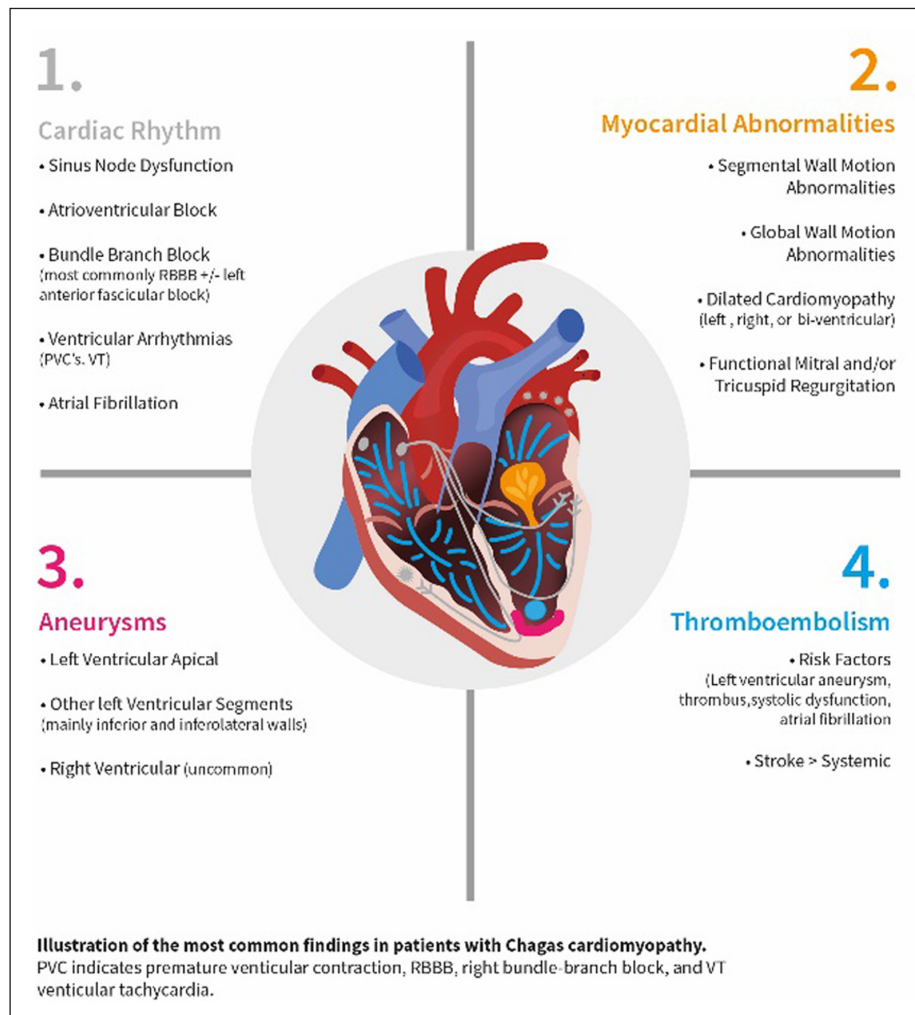


Figure 5: Most common findings in patients with Chagas cardiomyopathy [23].

The extent of cardiac involvement in the chronic phase of the disease appears to be the result of the parasite-activated immune response, but parasite persistence during the chronic stage of infection is critical. The immune response elicited in the acute phase and maintained during the chronic one seems to be influenced by variables such as parasite load during the acute phase, parasite strain, the magnitude of the immune response, and the presence or absence of reinfection [7].

CCC has a worse prognosis than other etiologies, with about 10% of patients progressing to terminal CHF, and is also associated with higher rates of hospital readmissions and mortality, regardless of age and in the absence of other comorbidities [7, 37, 38, 39, 40].

Cardiac mortality among CCC patients is mainly due to the high prevalence of life-threatening ventricular arrhythmias, manifesting as cardiac arrest and sudden death [23]. Additionally, the association of atrial fibrillation and apical aneurysms, along with a hypercoagulable state from *T. cruzi* infection provokes higher rates of embolic events compared to other heart failure etiologies [35, 36].

Although different clinical scoring systems, imaging modalities (Echo, MRI), and several biomarkers including NT-proBNP and Hs-cTnT have been associated with CCC stages of severity and subsequent mortality, the development of better predictors of disease progression and prognosis is still needed [41].

Gastrointestinal manifestations

Some patients, especially those infected with strains of the parasite found in the southern countries of Latin America (Brazil, Bolivia, Argentina) can present with the digestive form of the disease. This form, which involves denervation of the autonomic plexuses of the digestive tract, leads to disturbances in absorption, motility, and secretion. Motor incoordination and subsequent dilatation result in megaviscera, involving mainly the esophagus and the colon [7]. Symptoms of megaesophagus include those typical for achalasia, such as dysphagia (retention of food in the esophagus), and symptoms of megacolon include constipation, often profound, with rare volvulus requiring surgical correction.

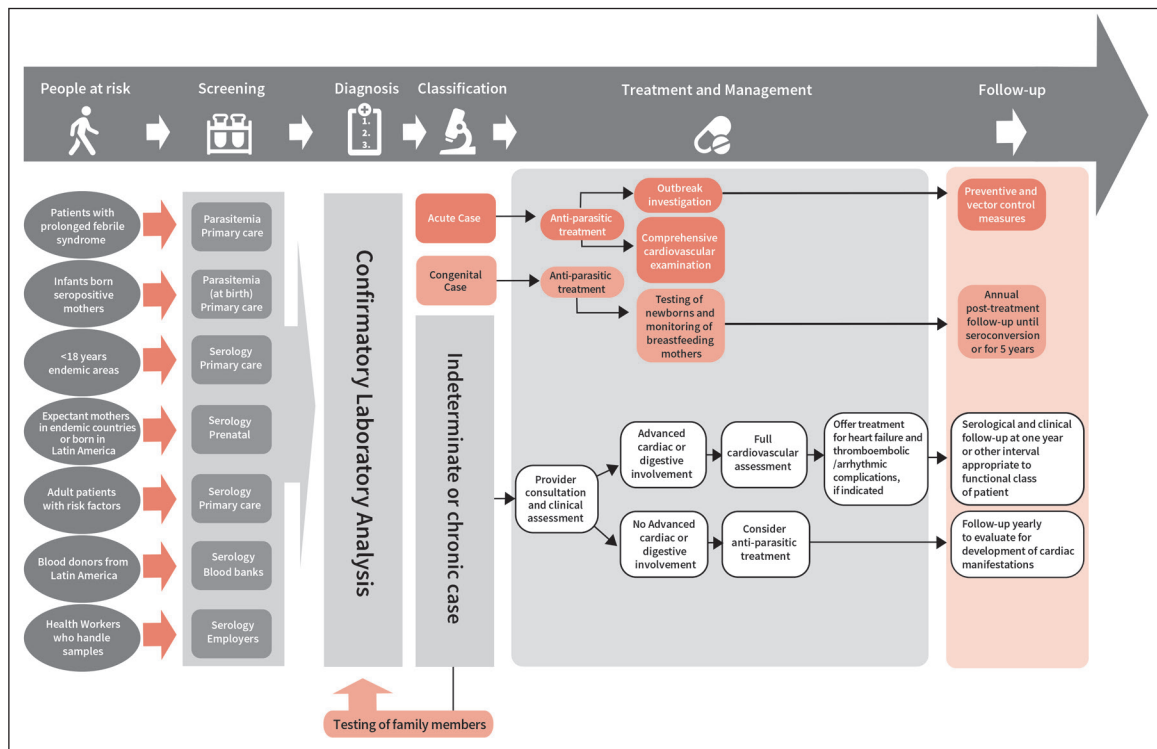


Figure 6: Ideal patient care pathway, Chagas disease.

Central nervous system involvement in chronic CD

In chronic CD, central nervous system (CNS) involvement is rare but associated with a poor prognosis. Most cases of CNS involvement are due to CD reactivation, especially in immuno-suppressed patients.

Patient pathway of care

Figure 6 shows an ideal patient journey for individuals affected by Chagas disease and represents a minimal standard for comprehensive care through all levels of interventions. In the following sections of the document, as we move along the steps of the journey and levels of intervention, we will endeavour to highlight gaps and roadblocks in the current standard of care, make recommendations and propose evidence-based solutions.

Prevention of Chagas Disease: Primary, secondary and tertiary interventions

Primary prevention

Preventing vector-based transmission

Vector control measures have achieved substantial success at eliminating transmission by domiciliary vectors in several countries, and as a result have been highly effective at reducing new vector-mediated cases of CD (Figure 7). Nevertheless, there are some remaining foci, notably in Central America and the Chaco region, where vector-based transmission persists. Reasons for this include reduced efficacy of pesticides in houses made of natural materials, and cases of pesticide resistance [42, 43]. The elimination of sylvatic vectors is virtually impossible and in some cases, they may occupy the niches left vacant by the elimination of domiciliary vectors. The re-emergence of vectors is a potential issue in areas where surveillance is weak and spraying programmes are sporadically enforced. In addition to this, political commitment to maintaining surveillance may wane once an area is certified as free of transmission [44]. Vector-based transmission of *T. cruzi* is closely associated with socioeconomic conditions, and is strongly associated with housing constructed of natural materials (mud, adobe, thatch).

The most sustainable way to interrupt vector-based transmission in remaining foci of infestation is through housing improvement and community development programmes that are supplemented by regular, systematic pesticide applications [43]. Successful control strategies will address both domestic and peri-domestic structures and raise awareness in the affected communities about the reasons for these efforts [43, 45, 46]. Strengthening of vector surveillance in endemic areas to allow for timely reporting of infestations, ideally as a collaborative effort between communities, health systems, and vector control teams, can help prevent re-emergence of vector-based transmission in areas where this modality of transmission has been interrupted.

| Level of prevention | Primary prevention | Secondary prevention | Tertiary prevention |
|---|--|--|--|
| Target population | At risk | Early/asymptomatic disease | Established disease |
| Main Objective | Prevent the transmission of the disease | Eliminate infection or prevent disease progression | To reduce morbidity and mortality and improve quality of life |
| Main strategies | <ul style="list-style-type: none"> Prevention of transmission by vector Prevention of transmission by transfusion Prevention of transmission by organ transplantation Prevention of congenital transmission Prevention of accidental transmission | <ul style="list-style-type: none"> Screening Case finding Antiparasitic treatment Periodic health examinations | <ul style="list-style-type: none"> Continuum of care Early detection of complications Clinical management and treatment |
| Psychological support, assess risk of reinfection, family screening, follow up | | | |
| Main interventions | <ul style="list-style-type: none"> Improvements of housing socioeconomic development Domiciliary infestation and environmental management Ongoing surveillance and screening Screening and treatment of females of childbearing age Promotion of safe food handling | <ul style="list-style-type: none"> Etiological treatment Improved access to health services Timely detection and treatment of congenitally infected infants | <ul style="list-style-type: none"> Comprehensive medical management of cardiac and digestive manifestations Establish systems of palliative care |
| | No Disease | Indeterminate form | Chronic complications |
| Disease progression | | | |

Figure 7: Levels of prevention and main interventions.

Preventing congenital transmission

Congenital transmission is most effectively interrupted by screening all women of childbearing age and treating seropositive women with anti-parasitic agents prior to pregnancy. Several observational studies indicate that anti-trypanosomal treatment of women is dramatically effective in preventing congenital transmission in future pregnancies [47, 48, 49]. While contraindicated during pregnancy, treatment of females of childbearing age is now recommended in major guidelines [50, 51]. Although new national and regional initiatives aim at reducing congenital transmission, many gaps remain. Healthcare personnel are often unfamiliar with the risk of congenital transmission [52]. Screening of pregnant women is far from universal, and totally absent in some affected countries [53, 54, 55], despite the demonstrated cost-effectiveness of these programmes in both endemic and non-endemic settings [53, 55, 56]. The regional initiative Eliminating Mother to Child Transmission Plus, launched by PAHO in 2017 [57], works to strengthen health systems in participating countries to interrupt congenital transmission of CD along with other congenitally transmitted infectious diseases, although the recommendations in this document are far from being widely adopted in the region.

For pregnant women, a universal serological screening protocol during prenatal visits should be established at the primary care level and integrated into existing perinatal care structures. Clear guidelines and procedures for verified cases of infected newborns should be developed, including serological monitoring for infection of any babies born to infected mothers who do not have overt parasitemia at birth. Decentralized models of care, such as outreach primary care should also include local medical facilities for serologic testing and clinical follow-up of infected adults, children and pregnant mothers, in a timely manner.

Preventing oral transmission

Orally acquired CD, an important syndrome in the Amazon region, is associated with a potentially virulent acute phase [20]. Because oral transmission occurs primarily through food, especially fruit contaminated with triatomine feces, implementation of safe food-handling practices combined with health promotion activities on the prevention of oral transmission are needed to prevent continued oral CD outbreaks. Promotion of safe food-handling practices among households and food vendors in areas at risk is essential, especially for high-risk foods such as sugar cane and acai [58, 59].

Preventing transmission via blood transfusion, organ transplant, and laboratory accidents

There has been growing acceptance of the need for screening of blood donations, in both endemic and non-endemic countries over the past 30 years, resulting in a significant decrease in known transfusion-related diseases. Nevertheless, screening remains only partially implemented and is essential for preventing transfusion and transplant transmission [60, 61, 62]. Prevention of accidental transmission in laboratory settings is best achieved through rigorous maintenance of safe laboratory practices in those institutions where *T. cruzi* and/or infected triatomines or blood and tissue samples are handled [63, 64, 65].

CD blood bank testing protocols should be in place with specific technical guidelines, collection protocols, reporting, and counselling and follow-up of positive cases. Special attention should be given to confirming inconclusive results with a combination of tests, given that the estimated prevalence of CD has been reported to be 13.30% in the stratum of donors with inconclusive serology at screening, and up to two false-negative results have also been reported, depending on the technique used for testing [32, 66].

Roadblocks to primary prevention:

- Discontinuation and interruptions of vector control interventions threaten to stall remarkable progress towards the elimination of vector-transmitted disease.
- There is inadequate focus on continued surveillance and eradication.
- Ongoing congenital transmission is facilitated by the lack of widespread adoption of screening recommendations to achieve early diagnosis and treatment.

Secondary prevention: Diagnosis and treatment

Timely diagnosis and treatment is critical in order to improve treatment success rates and prevent progression to the chronic phase of the disease. As part of a comprehensive approach, the diagnosis and treatment process should involve screening for chronic manifestations of the disease at the point of care. A thorough clinical review, including medical history and physical examination, should seek to elicit clinical signs and symptoms not only of acute CD but also cardiac and gastrointestinal chronic manifestations. As a minimum, an ECG should be performed to detect cardiac involvement in all patients with positive serology.

In order to vastly improve currently inadequate diagnosis rates, all individuals from endemic areas should be screened for CD on at least one occasion. In endemic regions, pregnant women as well as newborns from infected mothers should be tested. Blood donors should always be tested.

In non-endemic settings where screening is not universally offered, healthcare providers should consider the following risk factors when determining who to test:

- Having lived in an endemic area.
- Having lived in a rural area of an endemic country, particularly in housing made of natural materials (mud, adobe, thatch, palm leaves).
- Having a family member with CD, especially a sibling with congenital CD.
- Having been born to an infected mother (if the answer is no, it is important to ask for the epidemiological background of grandmothers and other family members, as well as a history of cardiac disease).
- Having been in contact with kissing bugs, particularly having seen them in the home or recalling bites.
- Having received blood transfusions, or having undergone major surgery (during which a blood transfusion may have occurred).
- Having received organ transplants in endemic areas.
- Having a history of intravenous drug use in an endemic country.

Immunocompromised patients with *T. cruzi* infection must be under clinical and laboratory monitoring for acute reactivation. Any clinical signs such as fever, neurological findings or acute myocarditis must be considered as warning signs of infection reactivation [67].

Diagnosis of *T. cruzi* infection in the general population

In areas with endemic vector-borne transmission, acute CD is generally a disease of childhood, although there are notable exceptions such as orally transmitted infection. Given the non-specific clinical manifestations, the diagnosis of acute CD is rarely made. However, if suspected, detection of the parasite via direct observation in the blood and/or molecular diagnosis are the first line techniques due to the high level of parasitemia in this phase. Serological testing is the second option, since antibodies may not yet be detectable.

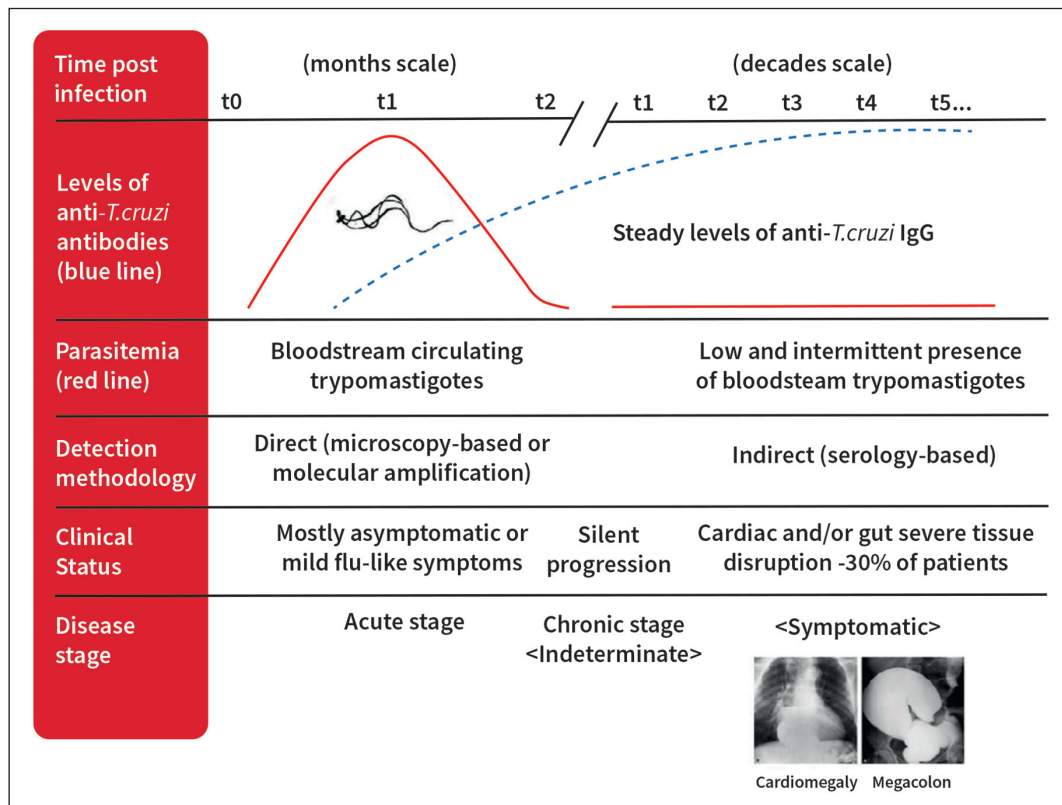


Figure 8: Timescale of Chagas disease diagnosis and clinical status progression [68].

Regardless of the mode of transmission, without treatment CD exhibits a trend of parasitemia that peaks around 30 days after infection (**Figure 8**). It then declines steadily until around 90 days after infection, when the parasite becomes undetectable by direct microscopy and specific antibodies against *T. cruzi* are present in blood.

During the chronic phase of the infection, which is lifelong in the absence of successful antitrypanosomal treatment, parasitemia remains below microscopically detectable levels, and diagnosis relies on serologic assays to demonstrate IgG antibodies against the parasite. To date, no single assay is sufficiently sensitive and specific to define infection. Positive results by the screening assay must be confirmed by a second test; hence, the diagnostic process is cumbersome. Additionally, the performance of diagnostic tools is highly variable [69, 70]. Simplifying diagnosis with high quality rapid testing that does not require external confirmation is critical to timely diagnosis, ideally utilizing a test that performs acceptably with patients at risk for infection with a variety of *Trypanosoma* strains.

There are currently a host of other barriers that are severely restricting access to timely and accurate diagnosis for at risk populations [71, 72]. The awareness of CD in the population and among healthcare providers is low, and there is a lack of knowledge on who to screen as well as a lack of clarity on the appropriate tests and interpretation of results [73]. There is an underappreciation of the importance of early diagnosis and treatment, especially at the primary healthcare level, and this represents a missed opportunity for treatment in the early phases, when treatment success rates are higher, and before severe organ damage occurs.

Diagnosis of *T. cruzi* in special populations

Diagnosis in newborns is made by direct visualization of trypomastigotes in the blood. This method is highly specific and confirms congenital infection, but it has low sensitivity that can be influenced by low parasitemia or an inexperienced technician [74]. Usually, at least, two blood samples are needed, one in the perinatal period and the second up to one month after birth. If positive, the newborn should be treated. If negative, a serologic test is required after the tenth month of age, when maternally transferred IgG antibodies have disappeared. In many regions, the required schedule of testing can make follow-up difficult, due to barriers to healthcare as well as migratory flows of populations [55, 68, 75]. This may account in part for the estimation that one in two of all congenital infections with CD are missed [76]. New molecular diagnostic tools such as PCR, which is used in the United States for the diagnosis of congenital CD, may eventually facilitate less cumbersome diagnosis of such cases if they can be successfully adapted to clinical settings [24, 77, 78, 79, 80].

In patients who are immunosuppressed, detection of the parasite by a direct parasitological test or exponential increase of parasitic load on quantitative PCR must be considered a reactivation, and the affected individual should receive anti-parasitic treatment immediately to avoid severe morbidity and/or mortality [81, 82].

Anti-trypanosomal treatment during the acute phase

Anti-parasitic treatment should be provided as soon as possible following detection of acute *T. cruzi* infection. Treatment during the acute phase is highly effective, producing serological cure, reducing potentially severe clinical manifestations of the acute phase, and preventing progression to chronic CD [83, 84].

The efficacy of treatment in the acute phase is almost immediately demonstrable by negative parasitemia with direct or indirect parasitological testing. In addition, antibodies disappear completely (sero-negativization) in at least 65% of cases, with some studies demonstrating sero-negativity in almost 100% of cases within 18 months of follow-up after treatment. This effect is independent of patient age. The absence of parasitemia, demonstrated by direct methods such as Strout or micro method, and negative PCR results always precede the reduction of antibodies [85, 86, 87]. Parasitic negativization occurs in the majority of treated newborns. Long-term follow-up studies are necessary to assess the relationship of treatment to rates of subsequent clinical events in infants [88, 89, 90, 91, 92].

Anti-trypanosomal treatment during the chronic phase

Anti-parasitic treatment during the chronic phase has an acceptable safety profile and is better tolerated in children than adults. However, around 17–35% of adult patients suspend the treatment due to side effects [93, 94]. The most frequent and serious are cutaneous, neurologic, hepatic, and hematologic. Anti-parasitic therapy is contraindicated in pregnancy, but may be offered during breastfeeding if indicated [95, 96]. Other contraindications include renal or hepatic insufficiency and advanced cardiomyopathy (**Figure 9**).

In the absence of a gold standard, serological tests to confirm cure and molecular tests to demonstrate failure are the best available tools to assess response to antiparasitic treatment in the chronic phase. While positive PCR during the first 24 months after treatment is indicative of therapeutic failure, sensitivity is variable [97, 98]. Post-treatment sero-negativization can take several years, depending on: (i) the age of the individual at time of treatment; (ii) the time elapsed between treatment and follow-up, and (iii) the region where the individual was infected [98, 99, 100]. Although complete sero-negativization can be obtained within five years in more than 70% of children treated, this rate only reaches about 30% in adult patients after roughly 20 years of follow-up (**Figure 10**) [48, 101, 102, 103, 104].

Although several observational studies show that trypanocidal treatment prevents morbidity and mortality [102, 103, 104], a randomized control trial of adults with established cardiac disease did not demonstrate clinical benefit after six years of follow-up [93]. In the absence of randomized trial evidence addressing the reduction of clinical events associated with anti-parasitic therapy, the efficacy of anti-parasitic therapy in indeterminate phase adults remains unknown, and demonstrates the need for further research to develop medications with better efficacy in this patient group.

Recent clinical trials indicate a high ($\approx 80\%$) rate of negativization by PCR in adults undergoing treatment with benznidazole [105, 106, 107]. In a recently concluded study, this rate was maintained after only two weeks of treatment with the standard dose, and no adverse events occurred [108]. However, new chemical entities (posaconazole and fosravuconazole) did not successfully demonstrate anti-parasitic efficacy in recent research [105, 106, 107].

Because of the high rate of side effects associated with current anti-parasitic therapy, treatment must be accessible in healthcare facilities near affected communities, and integrated into the local primary care infrastructure. This should be coupled with culturally sensitive educational health promotion programs about the importance of treatment and information on possible side effects. Community-led patient support groups also have an important role to play in providing information regarding CD and treatments. Healthcare staff should be adequately trained to deliver appropriate treatment and manage side effects, in keeping with locally adapted guidelines. Comprehensive follow-up of patients should also involve assisting them in navigating the healthcare system and addressing social or environmental hurdles that could interfere with treatment adherence (**Figure 10**).

Improved regimens of benznidazole and nifurtimox along with more research into their efficacy in adults, as well as development of new drugs with improved safety profiles and tolerability, are an important future step. However, providers serving affected populations should also receive tools and training to provide optimal treatment with the current drug regimens according to PAHO and national guidelines. Another crucial need is the development of biomarkers to help clinicians assess treatment response. Testing and treatment need to be included in public and private insurance plans, and should be available at no cost to low-income patients in facilities that are near affected communities.

Key treatment interventions

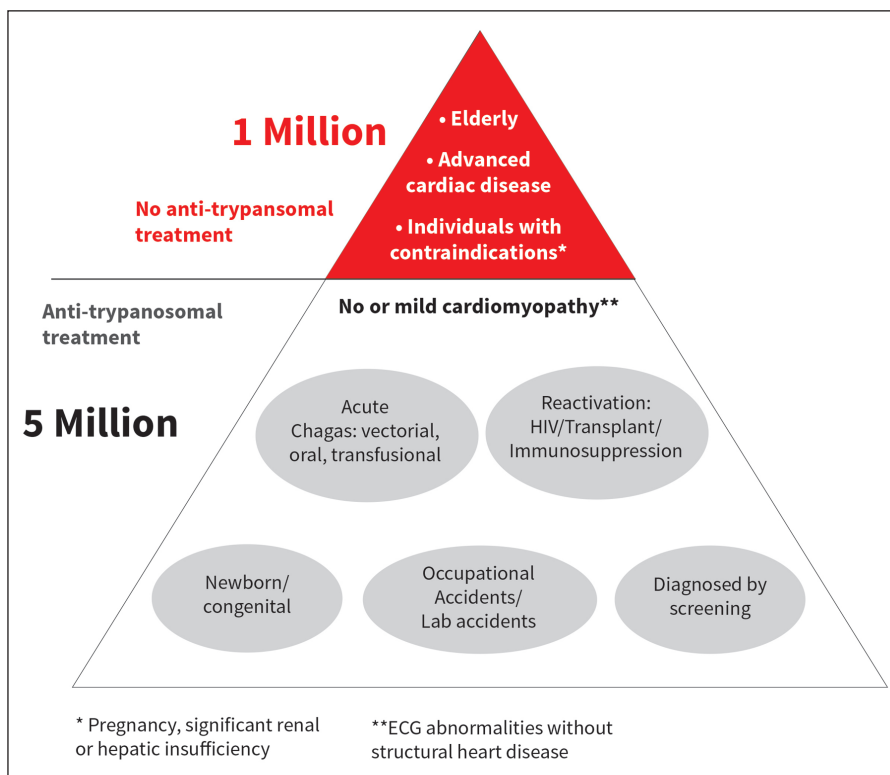


Figure 9: Indications and contraindications for anti-trypanosomal treatment and number of people in both categories globally.

| Category | PAHO recommendation | Recommendation level | Strength of evidence |
|--|--|----------------------|----------------------|
| Acute infection | Administer anti-trypanosomal treatment | Strong | Moderate |
| Congenital infection | Administer anti-trypanosomal treatment | Strong | Moderate |
| Children with chronic infection | Administer anti-trypanosomal treatment | Strong | Moderate |
| Females of childbearing age (15-44) | Administer anti-trypanosomal treatment | Strong | Moderate |
| Adults with chronic infection, no organ involvement | Offer anti-trypanosomal treatment | Conditional | Low |
| Adults with chronic infection, moderate to severe organ involvement | Do not offer any anti-trypanosomal treatment | Conditional | Moderate |
| Reactivation in immunocompromised patients* | Administer anti-trypanosomal treatment | Strong | Low |
| Cases of laboratory or surgical accidents | Administer anti-trypanosomal treatment | Strong | Low |

Figure 10: Anti-trypanosomal Treatment Recommendation. Adapted from Guidelines for the diagnosis and treatment of Chagas disease. Washington, DC: PAHO; 2019.

Roadblocks to secondary prevention:

- Inadequate screening programmes hampering early diagnosis and treatment.
- Inadequate knowledge of the diagnostic process and treatment recommendations in the healthcare community.
- Underappreciation by healthcare professionals of the importance of early diagnosis and treatment, and of screening for early stages of end organ damage.
- Cumbersome diagnostic process, with suboptimal test performance that requires confirmatory testing. Poor availability at the point of care.
- Inadequate investment in R&D for effective anti-parasitic agents with acceptable side effect profiles.
- Prohibitive cost of medical care and other out-of-pocket expenses.

In the words of one patient who spent over 10 years seeking treatment, 'I thought I had a doctor, a clinic who could help me, but Chagas taught me this is not always so. Because nobody, I mean nobody knew what CD is. My doctor didn't know, and he sent me to a specialist, and every week I went to a different doctor, but nobody knew what to do with me. And after a year I gave up.' [109].

Tertiary prevention

Tertiary prevention involves activities that mitigate or reduce morbidity and mortality from cardiac and other complications caused by CD, and improve the quality of life of affected people. Often, after a long asymptomatic phase, CCC is the most important clinical manifestation of the disease, resulting in the main burden of CD morbidity and mortality [110].

Clinical management and treatment of chronic Chagas cardiomyopathy

When the acute phase of CD is symptomatic, treatment has two purposes: clinical stabilization of the patient, and control of infection [7]. Management of the clinical consequences of myocarditis caused by *Trypanosoma cruzi* does not differ from other types of infectious myocarditis except for the use of anti-trypanosomal treatment [111]. Approximately 30% of individuals with positive serology without abnormal cardiac function at baseline will eventually develop myocardial dysfunction [23]. Compared to non-infected patients with ventricular dysfunction of other etiologies, CD patients have higher rates of conduction abnormalities, lethal arrhythmia, thromboembolic events, and ultimately mortality [39]. In addition, CCC has a worse prognosis in terms of impact on the quality of life, with higher rates of hospitalizations as a result of these complications [39]. The few studies that have evaluated guideline directed medical therapy in CCC have been compromised by small sample sizes, and most are non-randomized. However, this panel of medical therapy is accepted as appropriate, and includes beta-blockers, angiotensin-converting enzyme inhibitors/angiotensin receptor blockers, angiotensin receptor-neprilysin inhibitors (ARNI), and mineralocorticoid receptor blockers [112]. Because of the unique pathophysiology of CCC, new protocols specifically evaluating therapies in this population are critical, and include PARACHUTE-HF (sacubitril-valsartan) [113] and COACH (colchicine) [114].

Diuretic therapy and digoxin use, while not studied in Chagas cardiomyopathy patients specifically, are appropriate medical therapies to improve functional class and control symptoms in Chagas heart failure. There are theoretical concerns about digoxin effects on an already damaged conduction system, so it should be used with caution. Amiodarone is frequently used to suppress ventricular tachyarrhythmias as well, also in the absence of randomized trial data establishing a benefit. Stroke, systemic, and pulmonary embolism are common in CD, more so than in heart failure of other etiologies. The incidence of ischemic stroke in patients with CCC has been reported to be as high as 2.67 events/100 person-years [115]. Apical aneurysms, found in up to 50% of symptomatic CCC patients, and higher rates of atrial fibrillation in CCC contribute to this elevated thromboembolic risk. A score of at least four points on the well-known Sousa Scale confers a stroke risk of 4.4% per year, which theoretically would overcome the risks of use of warfarin or a direct anticoagulant [116]. Despite the logic of initiating anticoagulation in these individuals, there are still no published randomized clinical trials to confirm the benefit of anticoagulants in CCC in a primary prevention setting, particularly that acknowledge the unusual mechanisms of stroke that this disease confers.

Because of the higher rate of arrhythmia in CCC, electrophysiologic interventions are a cornerstone of therapy for this disease. Amiodarone is widely used, although studies are conflicting in regard to its effect

on mortality. Device therapies used in CCC include implantable devices such as implantable cardioverter defibrillators (ICDs) and cardiac resynchronization therapy (CRT). Although use of CRT is strongly supported by trial evidence in heart failure in general, there have been no randomized trials to date that have evaluated these therapies in CCC. Importantly, observational studies have not ubiquitously shown a benefit, underscoring the substantial need for performing studies of both ICDs and CRT in CCC patients, as well as the usefulness of amiodarone versus ICDs in CCC [117, 118]. Any study of ICD therapy would ideally address the higher risk among Chagas patients for lethal arrhythmias at ejection fractions that are higher than those that usually confer risk in heart failure of other etiologies.

In the past, CCC has been considered a potential contraindication to transplantation because of concern about reactivation of infection. Nevertheless, in spite of frequent but easily treatable acute reactivation events after transplant, acceptable survival rates have been demonstrated. Because of these results, CCC has not been considered a contraindication for transplantation since the 1990s [119].

Evaluation and treatment plan for CCC

Figure 11 delineates the ideal treatment algorithm for a patient with positive serology for CD. Initial cardiac evaluation can be performed at the primary care level, but when ECG changes are present, evaluation by a cardiologist is appropriate. In the presence of significant structural heart disease or arrhythmia, evaluation by a heart failure/cardiomyopathy specialist with experience with Chagas disease is critical, to make sure that appropriate emphasis is placed on risk stratification for stroke and arrhythmia. This is particularly important in non-endemic countries where the disease is not well understood. In the presence of clinical deterioration, patients should be assessed in a tertiary care centre with advanced heart failure treatment options.

Many critical barriers exist to timely, appropriate care for CCC. Low provider awareness, including the frequent under-recognition of CD as the cause of cardiomyopathy, can result in substandard management. Additionally, many imaging modalities, complementary exams and necessary devices (pacemakers, cardioverter-defibrillators) are not available in low-resource settings [120]. The coexistence of advanced chronic CD with a range of other comorbidities, including diabetes, hypertension, depression, or coinfection with HIV, means that CD may be overshadowed by other more 'urgent' medical needs [121, 122, 123].

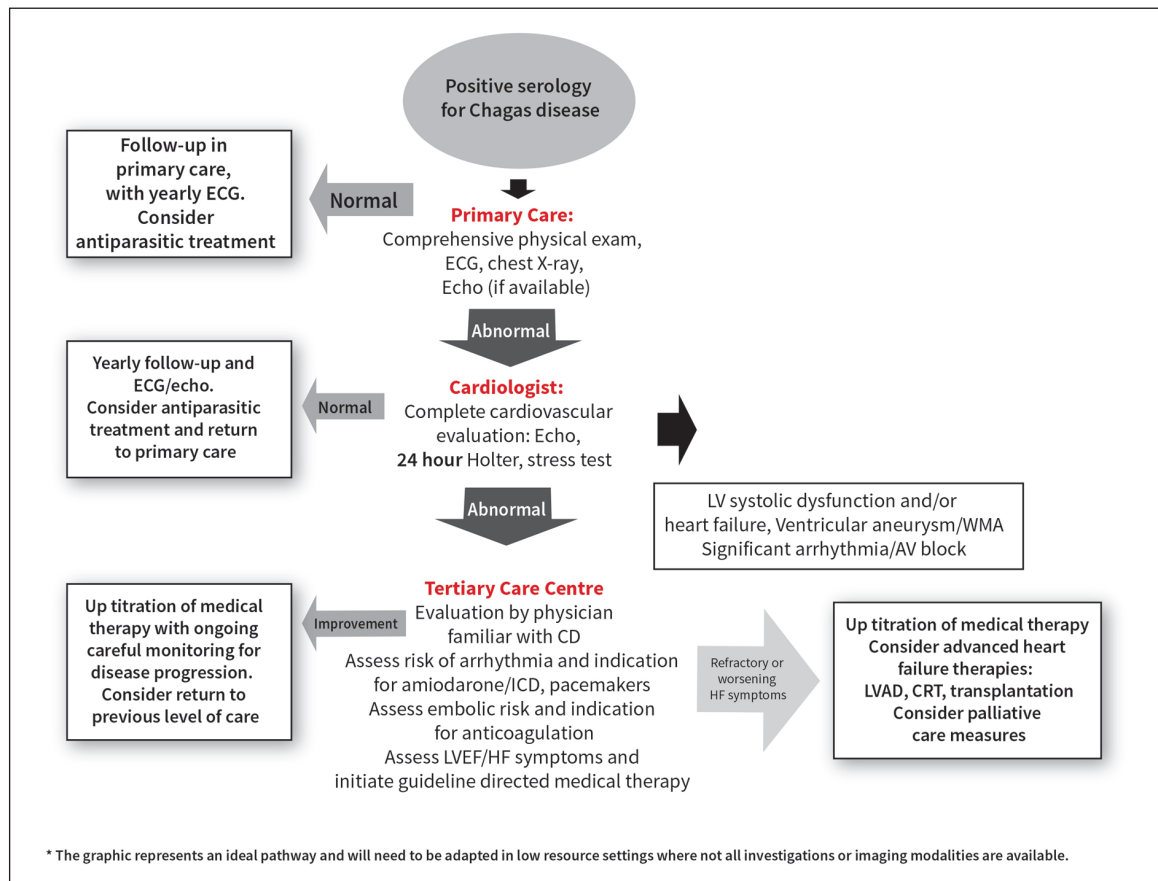


Figure 11: Algorithm for cardiovascular evaluation in patients with positive serology for CD.

Clear, rapid referral pathways to tertiary care centres are needed from the primary care level in order to access necessary imaging and investigations, as well as cardiologists and other specialists. Interventions which increase the availability of specialized care to patients in non-urban settings (e.g. telemedicine, mobile clinics) are useful to increase patient access. Ongoing monitoring of patients should be systematically coordinated locally with patients' primary healthcare service.

In terms of treatment for individuals with moderate to severe cardiomyopathy, R&D for new treatments capable of reversing or negating organ damage secondary to *T. cruzi* infection are urgently needed, thereby acknowledging that Chagas cardiomyopathy is a distinctive and more frequently lethal form of cardiomyopathy.

Roadblocks to tertiary care:

- Therapies for advanced disease are costly and frequently only available in urban centres, which are often far from where patients reside.
- Inadequate research has been conducted to evaluate which individuals will progress to clinically significant disease.
- There has not been adequate research to prove that accepted heart failure therapies are effective in CCC, and how to address the higher risk of thromboembolic phenomenon and ventricular arrhythmias in this disease.
- New therapies for advanced CCC that halt disease progression are needed.

Chagas Disease: Key Issues

Inadequate R & D

While significant progress has been made in the understanding of CD in the century since Carlos Chagas' landmark research, there are still only two available anti-trypanosomal drugs, both having unsatisfactory efficacy and safety profiles. Funding for R&D to improve treatments is strikingly low, given the 75 million people at risk in Latin America alone and the significant financial and social burden of the disease. Less than \$1 million USD, representing only 0.04% of R&D funding dedicated to neglected diseases, was spent on the development of new drugs for CD in 2007. Although there has been some improvement in the last decade, there is still much work to be done [124, 125]. For a disease of global public health importance affecting millions, renewed impetus to develop improved treatments is well overdue. A substantial increase in R&D investment, preferably via product development partnerships (PDPs), will be critical to addressing the need for better pharmacologic therapy. PDPs success in providing patients in resource-poor settings with important therapeutic improvements against malaria demonstrate the efficiency of such collaboration [126].

Although there have been some new developments in R&D from recent clinical trials, there is an urgent need to review and challenge the current evidence and define clinical research priorities for the immediate future. This would ensure that appropriate, efficacious drugs for CD are developed, but will require a focused and collaborative effort from the entire CD research community [127].

Additional anti-trypanosomal drugs are currently under development, and there are also CD vaccine (immunotherapeutic) candidates at the pre-clinical stage [128]. In this respect, a PDP is exploring an approach that links therapeutic vaccination to pharmacotherapy [129].

Scaling up access to diagnostic testing and treatment

The last decades have seen significant changes in approaches to the treatment of CD. Anti-trypanosomal treatments were previously reserved for acute cases and children, but improved knowledge of the pathophysiology of CD and a subsequent shift in thinking has meant that recommendations for testing and treatment now cover a much larger population group, including adults in the indeterminate phase. While randomized trial evidence is not yet available to assess efficacy of anti-parasitic therapy in indeterminate phase adults, recommendations suggest consideration of therapy in this population based on expert consensus. The prioritization of public health resources to focus on vector control programmes as the most cost-effective mechanism to reduce the burden of CD was a rational approach, and continuation of these programmes remains critical to maintaining success. However, it is now time to scale efforts up to provide easier access to diagnostic testing and medical therapy for those with established CD.

Recent PAHO guidelines recommend more widespread treatment of adults in the indeterminate phase of the illness, but to date the 'treatment gap' remains vast. With the exception of Spain, non-endemic countries do not have an organized approach to screening or treatment, largely due to a lack of awareness that individuals with CD now reside outside of endemic. Additionally, despite the widespread

acceptance of the benefits of prenatal screening, such programmes are not widespread across the Latin American region.

Several examples of local multi-stakeholder initiatives involving national and local governments provide successful models to scale up diagnosis and treatment in the adult population. In collaboration with the Spanish organization IS Global, the Chagas Platform in Bolivia provides an example of scaling up diagnosis and treatment in adults [130]. Under the umbrella of the National Healthcare System's Chagas National Programme (ChNP), the platform created several specialized centres for CD. The centres used clinical protocols, in which healthcare workers were trained and which involved robust data collection. The Platform demonstrates the possibility of dramatically scaling up access and provides a robust model for national adoption in Bolivia. The 4D (Diagnose, Design, Deliver, Demonstrate) approach utilized in Colombia provides another example of a multi-stakeholder pilot project which delivers a comprehensive model of care [131]. This model is also based on a collaborative approach with government and stakeholders, and aims to deliver a model of care which is needs based and data driven, with associated health care system and health care worker capacity building to ensure sustainability of the project. The Precede-Proceed model upscaling prenatal screening in Guatemala has also demonstrated that a data driven comprehensive approach involving professionals from all aspects of care can improve diagnosis/treatment rates [132].

Future regional and multinational efforts to address the shortfall in diagnosis and treatment should build upon these local and regional efforts, with strong educational components for both physicians and public health officials that promote the cost-efficiency of such interventions while highlighting the moral imperative of addressing the needs of this generally indigent and marginalized patient population.

Why do only some patients develop cardiac disease?

A critical knowledge gap exists as to why only 30% of infected individuals will progress to CCC [23]. Several host and parasite factors have been evaluated to understand this phenomenon. Thus far, the main factors identified as being related to cardiac damage are: Immune response to antigens of the parasite leading to fibrosing inflammation (T CD8 lymphocyte response), direct damage to myocytes by the presence of the parasite, damage of the neuronal cardiac system, autoantibodies against neuro-receptors, microvascular abnormalities, non-specific damage due presence of eosinophils and neutrophils, and oxidative stress. Moreover, different *T. cruzi* strains may be associated with different levels of cardiac toxicity [133, 134]. While these factors likely act at least simultaneously, if not synergistically, there is growing evidence that parasite persistence is a necessary factor for disease progression.

A genetic susceptibility to develop CCC has been proposed, which may result from polymorphisms in genes related to the IFN- γ axis that can lead to variations in the intensity of the immune response involved in the pathogenesis of the disease [135]. Further research is necessary to reveal the profile of patients who would benefit the most from trypanocidal treatment.

Why are there poor cure rates in chronic patients?

Without a reliable measure of parasitological eradication, it is difficult to know true cure rates of chronically infected Chagas patients, hence serologic cure is a surrogate endpoint for treatment success. The reasons that there are much lower rates of sero-negativization in a treated infected adult relative to treated children remain unknown. Identification of biomarkers that more accurately measure efficacy could revolutionize clinical research and practice, and are a current subject of research initiatives [136]. Although potentially these would be improved indicators of 'cure', there are still surrogate endpoints in the absence of data that demonstrate a correlation with reduction in clinical event rates.

Age has been identified as an important predictor of cure; anti-parasitic treated children have higher rates and a prompter response to therapy than adults [98]. In addition, studies suggest a differential response to therapy depending on country of origin, suggesting variations in susceptibility to anti-parasitic agents depending on the parasite strain.

Host and parasite biology may also have additional implications for the response to anti-parasitic therapy. Research to date has also hypothesized that the quality of T-cell responses and immune-regulatory mechanisms might determine the pattern of cellular responses and the severity of disease in chronic *T. cruzi* infection. The quality of T-cell responses might be a key factor, not only in disease evolution but also in chemotherapy responsiveness [137]. This may also explain the differing response rate to anti-parasitic treatment among children in comparison with adults, as children have a T-cell profile associated with a more robust clinical response [138].

Additionally, animal model research demonstrates that a small proportion of trypomastigotes are dormant at any given time during infection, and are likely to be protected from anti-parasitic compounds in this state [139].

Psychological aspects in comprehensive care of CD

“The patient with Chagas seeks something more than a tablet.” –Nilce Mendoza, an individual affected by CD residing in Spain [140]

Diagnosis of CD produces a range of reactions in individuals, from scepticism to fear and anxiety [141, 142]. Those affected may feel understandable doubts about their diagnosis, given the lack of noticeable symptoms, and may not remember having been exposed to triatomines or other risk factors. Particularly in non-endemic areas, many newly diagnosed individuals have never heard of CD and encounter health care professionals who are similarly unaware of the disease and may be dismissive of its importance [143]. Often, individuals feel devastated by the knowledge that they have a potentially life-threatening disease, which can lead to depression, especially in those who suffer debilitating chronic symptoms [144]. For many, thoughts of CD have to be pushed to the background in light of a host of other social, economic, and emotional challenges involved in the daily struggle to survive. Further, affected individuals who have immigrated to non-endemic countries may be isolated from their traditional support networks and could find it difficult to engage with healthcare providers in host countries because of political, linguistic and cultural barriers [109]. Finally, health systems rarely offer support for the emotional and social challenges of living with CD, which are exemplified below in the words of an affected person:

“[...] I have family with Chagas. And of course, when they said I have Chagas, and at any moment I could die, of course I became sad. I thought a lot about the disease. I worried a lot.” [109]

Patient and community involvement and empowerment in tackling CD

“It’s a fatal disease, and yet you don’t hear anything about it, it’s like a phantom disease that is killing people but nobody knows it exists, until they tell you, you have it. You always hear about diabetes, cancer, but [Chagas] disease is something that’s never heard anywhere, not even in the media.” –Sara, 60, El Salvador [109]

CD is largely a hidden disease, not only due to its long asymptomatic phase but also because it primarily (but not exclusively) affects politically and economically marginalized people [145]. This, combined with extremely low awareness of the disease among both providers and the general public, poses challenges for the organization and empowerment of people with CD. Nonetheless, patient organizations have a long history of being actively engaged in efforts to raise awareness of the disease, increase access to healthcare, and provide social and emotional support for affected people. For instance, the Pernambuco Association of Chagas and Heart Disease Patients in Brazil has worked since 1987, not only in educating community members but also providing social support to patients, operating in close collaboration with local healthcare services [146]. Patient organizations have emerged in endemic countries from Argentina to Mexico, and also in non-endemic countries. In 2010, the International Federation of Associations of People Affected by Chagas was formed and now consists of over 20 patient organizations. The road ahead for these patient organizations remains arduous, as the public profile of the disease, political commitment of governments, and access to political power of affected people all remain low.

Improving access to resources for clinical decision support?

Clinical management resources and clinical guidelines have been written and updated in different regions and settings, but require further updating and a broader discussion of comprehensive CD care.

To date, three types of guidelines have been published:

1. Regional guidelines. The most recent regional clinical guideline is the one published by the Pan-American Health Organization/WHO (PAHO/WHO) [32]. This document focuses on evidence-based recommendations for the diagnosis and treatment of CD in adult and paediatric patients. Whom to screen, management of cardiomyopathy, and how to implement the recommendations are not addressed.

2. Endemic country national guidelines. Several Latin American countries have developed clinical guidelines for screening and treatment [147, 148, 149, 150]. In general, these documents contain consistent guidelines and recommendations about prevention, epidemiological surveillance, vector control, and medical therapy. In particular, the Brazilian Ministry of Health has developed a very comprehensive Clinical Protocol (**Protocolo Clínico e Diretrizes Terapêuticas em Doença de Chagas**) [151], which includes a comprehensive consultation of key stakeholders and health professionals in order to help implement Brazil's Consensus Guidelines [43].
3. Non-endemic country guidelines. Several guidelines, consensus documents, and review articles have been published during the last decade in non-endemic countries to address specific clinical topics [23, 67, 152, 153, 154], related to CD and catering to clinicians with no previous knowledge of the disease. However, these guidelines do not address whom to screen.

To address the need for a more comprehensive, up-to-date, and broadly applicable guideline document, the Chagas Coalition has formed a CD Guidelines Review Board. The aim of this committee is to address the gaps in prior guideline documents and to update existing recommendations based on new evidence. This document will use PAHO guidelines as a general framework, harmonizing when appropriate individual country guidelines to a general standard. The document also aims to be comprehensive in addressing all levels of CD care, incorporating recommendations from a diverse group of clinicians from a variety of specialties and communities.

Roadblocks on the Roadmap to Chagas disease prevention, diagnosis and treatment

Figure 12 overcoming the roadblocks to appropriate and timely prevention, diagnosis, and treatment for CD cannot be separated from the overarching goal of providing universal health coverage (UHC). Access to comprehensive, community-centred health services is the cornerstone of UHC and also to delivering appropriate care to individuals affected by CD. In addition, strategies to address the social determinants of health should underpin any comprehensive strategy on CD. At the policy level, there is a need for strengthening stewardship and governance to ensure that CD is embedded in government programmes and Ministry of Health policies. Financing policies should acknowledge that prevention and early diagnosis are cost-efficient programmes to reduce the burden of disease. **Figure 12** summarizes the main roadblocks at four levels of intervention: prevention, diagnosis, treatment, and management of the clinical complications of the disease in its chronic stage.

Four factors common to all four steps on the pathway of care are discussed below: governance and advocacy, healthcare financing, the interaction between patients and caregivers, and information technology and registries.

Factors influencing all four levels of care

Governance and advocacy

Since the term 'Neglected Tropical Diseases' was first coined in 2003, global advocacy actions addressing CD as one of this diverse group of diseases have been gaining momentum and are a vital component in addressing the unmet needs of patients with CD. As a disease surrounded by stigma and affecting marginalized populations, CD has been slow to gain the attention of governments and policy makers. Its inclusion within this group of neglected diseases, with an organized and concerted advocacy strategy, has meant an unprecedented rise in awareness over the last 10 years. The London Declaration of 2012 was the first global multi-stakeholder commitment on NTDs, bringing together civil society organisations, development agencies and private enterprise with an ambitious statement to control or eliminate 10 NTDs by 2020 [155]. This declaration ran in parallel to the second WHO 2020 Roadmap on NTDs, which is due to be updated in 2020 [156]. Chagas patient groups have also played an important role in recent years, by giving a voice to CD patients. Prompted by the International Federation of Associations of People Affected by Chagas (FINDECHAGAS), the 72nd World Health Assembly in Geneva marked the adoption of April 14th as World Chagas Day in 2019. The formation of the Chagas Coalition in 2012, which is a collaborative alliance of stakeholders working on the disease, has added cohesion to individual actors' advocacy efforts and is also a valuable vehicle for pooling knowledge and technical expertise on CD.

| Level of intervention | Roadblock | Potential Solutions |
|--|--|--|
| Prevention | Lack of awareness and prioritization of CD by policy-makers and limited understanding of disease burden and cost to society | Increase advocacy by all stakeholders, NGO's and patient advocacy groups. |
| | Inadequate data on disease burden hinders effective policy making and design of effective healthcare systems interventions. | Establish CD (acute and chronic cases) as a reportable disease, with clear protocols for follow up |
| | Inadequate proportion of the at risk population screened | Establish and disseminate clear recommendations on which at risk populations to screen and work with local and national health systems to implement these recommendations |
| | Lack of integration within community centered healthcare | Integrate CD into existing primary care, reproductive health, and maternal child health care programs to ensure streamlined delivery of services and maximise cost cutting efficiencies |
| | Poor level of disease awareness among healthcare workers and physicians | Raise awareness and knowledge about the disease among healthcare professionals and physicians through education and training |
| | Poor levels of awareness among general population due to lack of information as well as social and cultural barriers | IEC (information, education, communication) activities in at risk populations |
| Diagnosis | Inadequate access to diagnosis at point of care | Integrate services in primary care and strengthen local healthcare infrastructure. Develop effective point of care diagnostic technologies |
| | Unsatisfactory performance of diagnostic tools and high incidence of inconclusive results | Development of improved diagnostic tools to simplify diagnostic process |
| | Lack of understanding by healthcare workers of the diagnostic pathway | Increase education for healthcare workers to better understand diagnostic processes |
| | Out of pocket expenses for diagnostic tests | Ensuring diagnostic tests are covered by health insurance programs Minimize associated out-of-pocket expenses for patients (travel, missed employment) |
| Treatment | Lack of therapeutic options with acceptable side effect profile and efficacy | Increased R&D to develop new molecules/treatments with improved efficacy and better side effect profiles. Continue research in to reformulations of existing molecules, and shorter treatment regimes. |
| | Stockouts and shortages of medications in some settings | Registration of medications in all affected countries and improve procurement practices to ensure supply stability |
| | Out of pocket expenses for treatments | Ensure treatments are free of charge |
| | Lack of healthcare worker and physician knowledge on treatment and side effects | Improve healthcare worker education and training on indications for treatment and how to manage side effects Design and disseminate clinical guidelines that are locally adapted |
| | Low levels of patient awareness on the importance of adherence to treatment and side effects, leading to discontinuation | Increase culturally-sensitive patient education and support Ensure adequate healthcare system infrastructure and staff to support follow up during treatment |
| Diagnosis and treatment of the clinical complications the disease | Lack of tertiary cardiac care facilities outside urban centers | Ensure rapid referral systems for cardiologist review and to tertiary centers Increase availability of complementary tests in non-urban areas for both cardiovascular and gastrointestinal complications of the disease (eg ECG, Echo) potentially via mobile or telemedical programs |
| | Low level of understanding among healthcare workers and physicians regarding chronic manifestations of the disease, including early identification of cardiomyopathy | Disseminate clinical guidelines which outline diagnosis, treatment and management of CCC Increase healthcare professional and physician education and training on chronic disease management, treatments and tertiary interventions |
| | Insufficient understanding of which patients are at risk of disease progression | Promote research initiatives focused on clinical and diagnostic predictors of disease evolution |
| | Insufficient evidence to guide appropriate medical therapy for CCC that acknowledges the CD as a distinct entity from other forms of heart failure | Develop innovative treatments for the chronic phase of the disease, increase funding for research to ascertain which medical therapies are appropriate for this population. |

Figure 12: Roadblocks and proposed solutions at different levels of interventions.

Advocacy is a key component to enable the embedding of CD interventions and programmes in Ministries of Health and other public institutions. Such a structure favours the sustainability of health programmes and promotes better national coordination and financing. NGOs and other stakeholders also have an important role to play in encouraging this 'buy in' from governments. Where government participation is absent, advocacy is needed to raise CD in political agendas.

Financing

Fragmentation is a common characteristic of most Latin American healthcare systems, leading to inefficiencies in financing and increased expenditure. Healthcare budgets are also all experiencing increased strain due to aging populations, and the increasing prevalence of non-communicable diseases. While increasing health-care spending on any individual disease may cause acute strain on healthcare budgets, it is clear that improving preventative efforts on CD as well as many other diseases, can effectively reduce this spending over time. In addition to these direct costs, CD has a substantial impact on worker productivity, by causing premature disability and death. It has been reported to represent a large economic burden [157], especially in endemic countries, with a global lifetime estimated burden of over \$ 188 billion USD [10]. Therefore, financial investment in health care interventions to tackle CD should be framed in terms of the long-term savings to the economy as well as the healthcare system. Efforts to streamline expenditure that is already in place for CD, and maximising cross-cutting efficiencies by integrating interventions into existing health care infrastructure, are necessary and important measures to better use available funds.

Individuals affected by CD and their caregivers

CD is rarely discussed in mass media or health education campaigns, despite the fact affected people suffer stigma and even exclusion from jobs [145]. In addition to this, in non-endemic countries, a lack of documentation of Chagas patients may cause affected individuals to be reticent to draw attention to themselves or their medical needs. The marginalisation and stigmatisation of the disease exerts a significant toll, both practically in terms of access to care as well as psychologically.

Both patients and caregivers benefit from support in navigating the social and emotional impacts of the disease. This should ideally involve social workers and mental health professionals as well as community based support groups [146]. The geographic distance of facilities from patients' communities can also be a major barrier in accessing care. Solutions such as providing transportation to referral centres or employing mobile clinics can help bridge this gap.

“Before, I couldn't go [to the doctor], because I didn't know how to drive. I had to wait for someone to take me; I depended on someone giving me a ride as a favour. And then I didn't have money to pay for the appointment or the ride, or sometimes for lack of time, and I've had to neglect other tasks so I could go to the doctor.” -Renata, 36, Mexico [109]

Many of the populations most affected by CD are migrant populations. Given this characteristic, these groups may be indirectly or overtly excluded from healthcare systems in host countries and may face linguistic, political, and cultural barriers that hamper their efforts to access services and also to organize and advocate for the right to healthcare. Information about CD and its treatment options should be accessible to individuals affected by the disease and their caregivers, available in their preferred language, and written in a straightforward, culturally appropriate manner without unnecessary medical jargon.

Registries, information technology and digital health

Registration of patients with CD is the starting point not only for clinical follow-up and as a mechanism for improving treatment adherence, but also as the foundation for healthcare planning and population health initiatives, allowing for the allocation of resources and evaluating the impact of healthcare interventions over time. In some cases, national registries represent less than 1% of the total expected cases based on estimates of prevalence [71].

Up-to-date, interlinked information systems are also a key element of an integrated approach to patient care within the healthcare system, offering an important opportunity to make clinical information accessible to all health care professionals and health authorities involved in the pathway of care. Furthermore, such a system has the potential to eventually provide information that can be used by the health sector and related sectors to develop truly integrated approaches to health in all policies.

Health professionals working in remote and resource-limited places will benefit from the development of digital and web-based tools to access expert advice through teleconsultations on diagnostic tools [158], such as electrocardiogram [159] and echocardiogram [160]. Mobile health interventions, such as the use of text messaging, can also be useful for both health professionals in the care of CD patients and for improving the adherence of patients to clinical care [161].

Platforms for e-learning are also a valuable tool to expand access to medical education, online access to guidelines and medical updates.

In the future, cognitive computing combined with artificial intelligence offers the prospect of readily accessible tools for patient self-assessment, including symptom status and side effects of therapy. Such tools may also be beneficial in monitoring adherence to treatment and improving patient health literacy and awareness of the disease [162].

Conclusions

CD is a complex but entirely preventable and treatable disease. The barriers affecting access to diagnosis, treatment and care are complex, and a strategic and comprehensive approach is required to address these roadblocks in the various settings where they exist.

This Roadmap provides an example of an ideal patient care pathway for CD (Figure 6), and explores the roadblocks along this pathway, considering potential solutions based on available research and evidence in practice. To move from prescribed global recommendations to local and national implementation, a number of specific actions are required to plan, design, and implement change.

The challenge remains of how to move from recommendations to practice implementation, and in that sense, it is crucial to adapt these recommendations to each particular framework at the national level. This means considering the particularities of the healthcare system and policy environment, and identifying specific barriers and potential strategies on a regional, national and local level.

The WHF implementation framework in Figure 13 offers a step-by-step approach to specific action areas and highlights the importance of an integrated approach across multiple care settings. Moving from a global roadmap initiative to a national call for action requires involvement and concerted action from national ministries of health, health care system decision-makers, and health care professionals. Patients are a pivotal part of these efforts, with the support of their families and caregivers, and civil society has an important role to play as well.

Bringing key leaders and stakeholders together for national roundtable discussions to consider a unified CD agenda based on national and global needs should be considered a necessary first step.

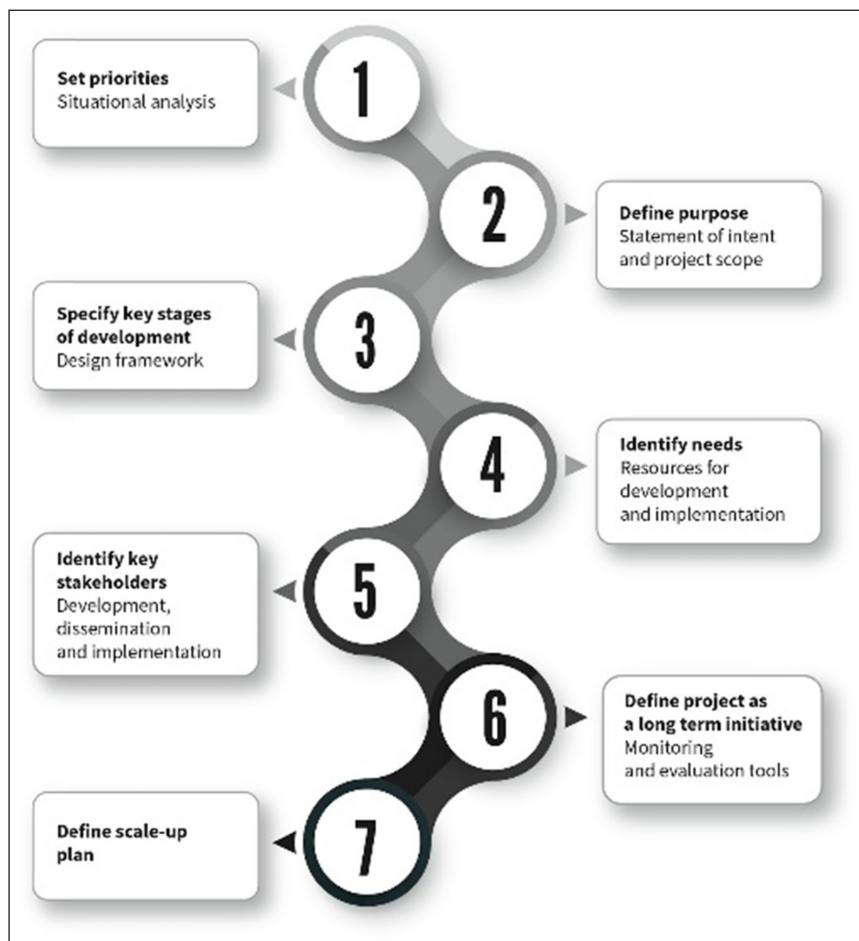


Figure 13: WHF Implementation framework.

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Competing Interests

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References

1. **Palafox B, Mocumbi A, Krishna Kumar K, Sulafa A.** The WHF Roadmap for Reducing CV Morbidity and Mortality Through Prevention and Control of RHD: A World Heart Federation Roadmap. *Global Heart*, 2017; 12(1): 47–62. DOI: <https://doi.org/10.1016/j.gheart.2016.12.001>
2. **Grainger Gasser A, Welch C, Arora M,** et al. Reducing Cardiovascular Mortality Through Tobacco Control: A World Heart Federation Roadmap. *Global Heart*. 2015; 10: 123–133. DOI: <https://doi.org/10.1016/j.gheart.2015.04.007>
3. **Adler A, Prabhakaran D, Bovet P,** et al. Reducing Cardiovascular Mortality Through Prevention and Management of Raised Blood Pressure: A World Heart Federation Roadmap. *Global Heart*; 2015; 111–122: 10. DOI: <https://doi.org/10.1016/j.gheart.2015.04.006>
4. **Perel P, Avezum A, Huffman M,** et al. Reducing Cardiovascular Mortality Through Tobacco Control: A World Heart Federation Roadmap. *Global Heart*. 2015; 10: 99–110. DOI: <https://doi.org/10.1016/j.gheart.2015.04.003>
5. **Murphy A,** et al. The World Heart Federation Roadmap for Nonvalvular Atrial Fibrillation. *Global Heart*. 2017; 12(4): 273–284. DOI: <https://doi.org/10.1016/j.gheart.2017.01.015>
6. **Mitchell S,** et al. A Roadmap on the Prevention of Cardiovascular Disease Among People Living With Diabetes. *Global Heart*. 2019; 14(3): 215–240. DOI: <https://doi.org/10.1016/j.gheart.2019.07.009>
7. **Rassi A Jr., Rassi A, Marin-Neto J.** Chagas Disease. *The Lancet*. 2010; 375(9723): 1388–1402. DOI: [https://doi.org/10.1016/S0140-6736\(10\)60061-X](https://doi.org/10.1016/S0140-6736(10)60061-X)
8. Chagas Disease in Latin America: An epidemiological update based on 2010 estimates. *Weekly Epidemiological Record*. 2015; 90(6): 33–43.
9. **WHO Department of Control of Neglected Tropical Diseases.** Integrating neglected tropical diseases in global health and development. *Fourth WHO report on neglected tropical diseases*. 2017.
10. **Lee BY, Bacon KM, Bottazzi ME, Hotez PJ.** Global economic burden of CD: A computational simulation model. *Lancet Infect Dis*. 2013; 13: 342–348. DOI: [https://doi.org/10.1016/S1473-3099\(13\)70002-1](https://doi.org/10.1016/S1473-3099(13)70002-1)
11. **Moncayo A, Silveira A.** Current epidemiological trends for CD in Latin America and future challenges in epidemiology, surveillance and health policy. *Mem Inst Oswaldo Cruz*. 2009; 104(1): 17–30. DOI: <https://doi.org/10.1590/S0074-02762009000900005>
12. **Chuit R, Meiss R, Salvatella R, Altchek J, Freilij H.** Epidemiology of Chagas Disease. 2019: 351. DOI: https://doi.org/10.1007/978-3-030-00054-7_4
13. **Manne-Goehler J, Umeh CA, Montgomery SP, Wirtz VJ.** Estimating the Burden of CD in the United States. *PLoS Neg Trop Dis*. 2016; 10(11). DOI: <https://doi.org/10.1371/journal.pntd.0005033>
14. **Gascon J, Bern C, Pinazo M.** CD in Spain, the United States and other non-endemic countries. *Acta Trop*. 2010; 115(1–3): 22–27. DOI: <https://doi.org/10.1016/j.actatropica.2009.07.019>
15. **Lima-Costa M, Peixoto S, Ribeiro A.** CD and mortality in old age as an emerging issue: 10 year follow-up of the Bambuí population-based cohort study (Brazil). *Int J Cardiol*. 2010; 145(2): 362–363. DOI: <https://doi.org/10.1016/j.ijcard.2010.02.036>
16. **Martins-Melo F, Carneiro M, Ribeiro A, Bezerra J, Werneck G.** Burden of CD in Brazil, 1990–2016: findings from the Global Burden of Disease Study 2016. *Int J Parasitol*. 2019; 49(3–4): 301–310. DOI: <https://doi.org/10.1016/j.ijpara.2018.11.008>
17. **Organización Panamericana de la Salud.** Guía para vigilancia, prevención, control y manejo clínico de la enfermedad de Chagas aguda transmitida por alimentos. Rio de Janeiro, 2009.
18. **Silva-Dos-Santos D, Barreto-de-Albuquerque J, Guerra B, Moreira O, Berbert L, Ramos M, Mascarenhas B, Britto C, Morrot A, Serra Villa-Verde D, Garzoni L, Savino W, Cotta-de-Almeida**

- V, de Meis J.** Unraveling CD transmission through the oral route: Gateways to *Trypanosoma cruzi* infection and target tissues. *PLoS Negl Trop Dis.* 2017; 11(4). DOI: <https://doi.org/10.1371/journal.pntd.0005507>
19. **Altcheh JM, Freilij H.** Chagas Disease: A Clinical Approach. *Springer International Publishing*; 2019. DOI: <https://doi.org/10.1007/978-3-030-00054-7>
 20. **Shikanai-Yasuda MA, Carvalho NB.** Oral Transmission of CD. *Clinical Infectious Diseases.* 2012; 54(6): 845–852. DOI: <https://doi.org/10.1093/cid/cir956>
 21. **Capuani L, Bierrenbach A, Pereira Alencar A, Mendrone A Jr., Ferreira J, Custer B, Ribeiro A, Cerdeira SE.** Mortality among blood donors seropositive and seronegative for CD (1996–2000) in São Paulo, Brazil: A death certificate linkage study. *PLoS Negl Trop Dis.* 2017; 11(5). DOI: <https://doi.org/10.1371/journal.pntd.0005542>
 22. **Stanaway J, Roth G.** The burden of CD: estimates and challenges. *Global Heart.* 2015; 10(3): 139–144. DOI: <https://doi.org/10.1016/j.gheart.2015.06.001>
 23. **Nunes M, Beaton A, Acquatella H,** et al. Chagas cardiomyopathy: An update of current clinical knowledge and management: A scientific statement from the American Heart Association. *Circulation,* 138(12): e169–e209. DOI: <https://doi.org/10.1161/CIR.0000000000000599>
 24. **Carlier Y, Torrico F, Sosa-Estani S,** et al. Congenital CD: recommendations for diagnosis, treatment and control of newborns, siblings and pregnant women. *PLoS Negl Trop Dis.* 2011; 5(10): e1250. DOI: <https://doi.org/10.1371/journal.pntd.0001250>
 25. **Prata A.** Clinical and epidemiological aspects of CD. *Lancet Infect Dis.* 2001; 1: 92–100. DOI: [https://doi.org/10.1016/S1473-3099\(01\)00065-2](https://doi.org/10.1016/S1473-3099(01)00065-2)
 26. **Laranja F, Dias E, Nobrega G, Miranda A.** Chagas' disease. A clinical, epidemiologic, and pathologic study. *Circulation.* 1956; 14: 1035–1060. DOI: <https://doi.org/10.1161/01.CIR.14.6.1035>
 27. **Pinto A, Valente S, Valente V, Ferreira A Jr., Coura J.** Acute phase of CD in the Brazilian Amazon region: Study of 233 cases from Pará, Amapá and Maranhão observed between 1988 and 2005. *Rev Soc Bras Med Trop.* 2008; 41: 602. DOI: <https://doi.org/10.1590/S0037-86822008000600011>
 28. **Echeverría L, Morillo C.** American Trypanosomiasis (Chagas Disease). *Infectious Disease Clinics of North America.* 2019; 33(1): 119–134. DOI: <https://doi.org/10.1016/j.idc.2018.10.015>
 29. **Lewis MD, Francisco AF, Jayawardhana S, Langston H, Taylor MC, Kelly JM.** Imaging the development of chronic CD after oral transmission. *Scientific Reports.* 2008; 8(1). DOI: <https://doi.org/10.1038/s41598-018-29564-7>
 30. **Almeida E, Lima J, Lages-Silva E, Guariento M, Aoki F,** et al. Chagas' disease and HIV co-infection in patients without effective antiretroviral therapy: prevalence, clinical presentation and natural history. *Trans R Soc Trop Med Hyg.* 2010; 104: 447–452. DOI: <https://doi.org/10.1016/j.trstmh.2010.02.004>
 31. **Pierrotti L, Carvalho N, Amorin J, Pascual J, Kotton C, López-Vélez R.** CD Recommendations for Solid-Organ Transplant Recipients and Donors. *Transplantation,* 2008; 102(2S): S1–S7. DOI: <https://doi.org/10.1097/TP.00000000000002019>
 32. **Organización Panamericana de la Salud.** *Guía para el diagnóstico y el tratamiento de la enfermedad de Chagas.* Washington, DC: OPS; 2018.
 33. **Pinazo M, Espinosa G, Cortes-Lletget C,** et al. Immunosuppression and CD: A management challenge. *PLoS Negl Trop Dis.* 2013; 7(1): e1965. DOI: <https://doi.org/10.1371/journal.pntd.0001965>
 34. **Lattes R, Lasala MB.** CD in the immunosuppressed patient. *Clinical Microbiology and Infection.* 2014; 20(4): 300–309. DOI: <https://doi.org/10.1111/1469-0691.12585>
 35. **Rojas L, Glisic M, Pletsch-Borba L,** et al. Electrocardiographic abnormalities in CD in the general population: A systematic review and meta-analysis. *PLoS Negl Trop Dis.* 2018; 12(6). DOI: <https://doi.org/10.1371/journal.pntd.0006567>
 36. **Acquatella H, Asch F, Barbosa M,** et al. Recommendations for Multimodality Cardiac Imaging in Patients with CD: A Report from the American Society of Echocardiography in Collaboration With the InterAmerican Association of Echocardiography (ECOSIAC) and the Cardiovascular Imaging Department. *Journal of the American Society of Echocardiography.* 2018; 31(1): 3–25. DOI: <https://doi.org/10.1016/j.echo.2017.10.019>
 37. Presence of parasite. *Am. J. Trop. Med. Hyg.* 2004; 70(2): 210–220.
 38. **Schijman AG,** et al. *Trypanosoma Cruzi* Dna In Cardiac Lesions of Argentinean Patients with End-Stage Chronic Chagas Heart Disease. *The American Journal of Tropical Medicine and Hygiene.* 2004; 70(2). DOI: <https://doi.org/10.4269/ajtmh.2004.70.210>
 39. **Shen L, Ramirez F, Martinez F, Bodanese L, Echeverría L, Gómez E,** et al. Contemporary Characteristics and Outcomes in Chagasic Heart Failure Compared with Other Nonischemic and

- Ischemic Cardiomyopathy. *Circ Heart Fail.* 2017; 10(11). DOI: <https://doi.org/10.1161/CIRCHEARTFAILURE.117.004361>
40. **Bocchi E, Bestetti R, Scanavacca M, Cunha Neto E, Issa V.** Chronic Chagas Heart Disease Management: From Etiology to Cardiomyopathy Treatment. *J Am Coll Cardiol.* 2017; 70(12): 1510–1524. DOI: <https://doi.org/10.1016/j.jacc.2017.08.004>
 41. **Echeverría L, Rojas L, Calvo L, et al.** Profiles of cardiovascular biomarkers according to severity stages of Chagas cardiomyopathy. *Int J Cardiol.* 2017; 227: 577–582. DOI: <https://doi.org/10.1016/j.ijcard.2016.10.098>
 42. **Gurevitz JM, Gaspe MS, Enriquez GF, Vassena CV, Alvarado-Otegui JA, Provecho YM, Mougabure Cueto GA, Picollo MI, Kitron U, Gürtler RE.** Unexpected Failures to Control CD Vectors With Pyrethroid Spraying in Northern Argentina. *Journal of Medical Entomology.* 2012; 49(6): 1379–1386. DOI: <https://doi.org/10.1603/ME11157>
 43. **Gürtler RE, Kitron U, Cecere C, Segura EL, Cohen JE.** Sustainable vector control and management of CD in the Gran Chaco, Argentina. *Proceedings of the National Academy of Sciences.* 2007; 104(41): 16194–16199. DOI: <https://doi.org/10.1073/pnas.0700863104>
 44. **Abad-Franch F, Diotaiuti L, Gurgel-Gonçalves R, Gürtler R.** Certifying the interruption of CD transmission by native vectors: cui bono? *Mem Inst Oswaldo Cruz.* 2013; 108(2): 251–254. DOI: <https://doi.org/10.1590/0074-0276108022013022>
 45. **Samuels A, Clark E, Galdos-Cardenas G, et al.** Epidemiology of and impact of insecticide spraying on CD in communities in the Bolivian Chaco. *PLoS Negl Trop Dis.* 2013; 7(8). DOI: <https://doi.org/10.1371/journal.pntd.0002358>
 46. **Rojas-de-Arias A.** CD prevention through improved housing using an ecosystem approach to health. *Cadernos de Saúde Pública,* 2001; 17: S89–S97. DOI: <https://doi.org/10.1590/S0102-311X2001000700017>
 47. **Sosa-Estani S, Cura E, Velazquez E, Yampotis C, Segura E.** Etiological treatment of young women infected with *Trypanosoma cruzi*, and prevention of congenital transmission. *Revista da Sociedade Brasileira de Medicina Tropical.* 2009; 42(5): 484–487. DOI: <https://doi.org/10.1590/S0037-86822009000500002>
 48. **Fabbro D, Danesi E, Olivera V, Codebó, M, Denner S, Heredia C, et al.** Trypanocide Treatment of Women Infected with *Trypanosoma cruzi* and Its Effect on Preventing Congenital Chagas. *PLOS Neglected Tropical Diseases.* 2014; 8(11): e3312. DOI: <https://doi.org/10.1371/journal.pntd.0003312>
 49. **Moscatelli G, Moroni S, García-Bournissen F, Ballering G, Bisio M, Freilij H, et al.** Prevention of congenital Chagas through treatment of girls and women of childbearing age. *Memórias do Instituto Oswaldo Cruz.* 2015; 110(4): 507–509. DOI: <https://doi.org/10.1590/0074-02760140347>
 50. **Dr. Mario Fatala Chaben, Instituto Nacional de Parasitología.** Pautas para la atención al paciente infectado con *Trypanosoma cruzi* (Enfermedad de Chagas). Buenos Aires: Ministerio de Salud; 2015.
 51. **Dias J, Ramos A Jr., Gontijo E, Luquetti A, Shikanai-Yasuda M, Coura J, et al.** Second Brazilian Consensus on CD. *Revista da Sociedade Brasileira de Medicina Tropical.* 2016; 49(3): 60. DOI: <https://doi.org/10.1590/0037-8682-0504-2016>
 52. **Verani J, Montgomery S, Schulkin J, Anderson B, Jones J.** Survey of obstetrician-gynecologists in the United States about CD. *Am J Trop Med Hyg.* 2010; 83(4): 891–895. DOI: <https://doi.org/10.4269/ajtmh.2010.09-0543>
 53. **Sicuri E, Muñoz J, Pinazo M, Posada E, Sanchez J, Alonso P, Gascon J.** 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): 110–117. DOI: <https://doi.org/10.1016/j.actatropica.2011.02.012>
 54. **Edwards MS, Stimpert KK, Bialek SR, Montgomery SP.** Evaluation and Management of Congenital CD in the United States. *Journal of the Pediatric Infectious Diseases Society.* 2019; 8: 461–469. DOI: <https://doi.org/10.1093/jpids/piz018>
 55. **Picado A, Cruz I, Redard-Jacot M, et al.** The burden of congenital CD and implementation of molecular diagnostic tools in Latin America. *BMJ Global Health.* 2018; 3. DOI: <https://doi.org/10.1136/bmjgh-2018-001069>
 56. **Stillwaggon E, Perez-Zetune V, Bialek S, Montgomery S.** Congenital CD in the United States: Cost Savings through Maternal Screening. *Am J Trop Med Hyg.* 2018; 98(6): 1733–1742. DOI: <https://doi.org/10.4269/ajtmh.17-0818>
 57. **Pan American Health Organization.** *EMTCT Plus, Framework for elimination of mother-to child transmission of HIV, Syphilis, Hepatitis B, and Chagas.* Washington, DC: PAHO; 2017.

58. **Ferreira R, Cabral M, Martins R**, et al. Detection and genotyping of *Trypanosoma cruzi* from açai products commercialized in Rio de Janeiro and Pará, Brazil. *Parasit Vectors*. 2018; 11(1): 233. DOI: <https://doi.org/10.1186/s13071-018-2699-6>
59. **Nóbrega A, García M, Tatto E, Obara M, Costa E, Sobel J, Araujo W**. Oral Transmission of CD by Consumption of Acai Palm Fruit, Brazil. *Emerging Infectious Diseases*. 2009; 15(4): 653–655. DOI: <https://doi.org/10.3201/eid1504.081450>
60. **Casadei D**. Chagas' disease and solid organ transplantation. *Transplant Proc*. 2010; 42(9): 3354–3359. DOI: <https://doi.org/10.1016/j.transproceed.2010.09.019>
61. **Organización Panamericana de la Salud**. *Suministro de sangre para transfusiones en los países de Latinoamérica y el Caribe 2012 y 2013*. Washington, DC: OPS; 2015.
62. Reporte del grupo de trabajo científico sobre la enfermedad de Chagas. Buenos Aires: Programa Especial de Investigaciones y Enseñanzas sobre Enfermedades Tropicales (TDR). 2005.
63. **Requena-Méndez A, Albajar-Viñas P, Angheben A, Chiodini P, Gascón J, Muñoz J**. Health policies to control CD transmission in European countries. *PLoS Negl Trop Dis*. 2014; 8(10): e3245. DOI: <https://doi.org/10.1371/journal.pntd.0003245>
64. **Hofflin JM, Sadler RH, Araujo FG, Page WE, Remington JS**. Laboratory-acquired CD. *Transactions of the Royal Society of Tropical Medicine and Hygiene*. 1987; 81(3): 437–440. DOI: [https://doi.org/10.1016/0035-9203\(87\)90162-3](https://doi.org/10.1016/0035-9203(87)90162-3)
65. **Salvatella R**. Una visión de la enfermedad de Chagas desde su propia historia. *Organización Panamericana de la Salud y Fundación Mundo Sano*. 2006: 19–22.
66. **Pereira G, Louzada-Neto F, Barbosa V, Ferreira-Silva MM, de Moraes-Souza H**. Performance of six diagnostic tests to screen for CD in blood banks and prevalence of *Trypanosoma cruzi* infection among donors with inconclusive serology screening based on the analysis of epidemiological variables. *Revista brasileira de hematologia e hemoterapia*. 2012; 34(4): 292–297. DOI: <https://doi.org/10.5581/1516-8484.20120074>
67. **Pinazo M, Miranda B, Rodríguez-Villar C, Altclas J, Brunet Serra M, García-Otero E, de Almeida E, de la Mata García M, Gascon J, García Rodríguez M, Manito N, Moreno Camacho A, Oppenheimer F, Puente S, Riarte A, Salas Coronas J, Salavert Lletí M, Sanz G, Torrico H, Tor-rús Tendero D, Ussetti P, Shikanai-Yasuda M**. Recommendations for management of CD in organ and hematopoietic tissue transplantation programs in nonendemic areas. *Transplant Rev*. 2011; 25(3): 91–101. DOI: <https://doi.org/10.1016/j.trre.2010.12.002>
68. **Alonso-Padilla J, Cortés-Serra N, Pinazo M, Bottazzi M**, et al. Strategies to enhance access to diagnosis and treatment for CD patients in Latin America. *Expert Review of Anti-infective Therapy*. 2019: 145–157. DOI: <https://doi.org/10.1080/14787210.2019.1577731>
69. **Cucunubá Z, Manne-Goehler J, Diaz D, Nouvellet P, Bernal O, Marchiol A**, et al. How universal is coverage and access to diagnosis and treatment for CD in Colombia? A health systems analysis. *Soc Sci Med*. 2017; 175: 187–198. DOI: <https://doi.org/10.1016/j.socscimed.2017.01.002>
70. **Verani J, Seitz A, Gilman R, LaFuente C, Galdos-Cardenas G, Kawai V**, et al. Geographic variation in the sensitivity of recombinant antigen-based rapid tests for chronic *Trypanosoma cruzi* infection. *Am J Trop Med Hyg*. 2009; 80(3): 410–415. DOI: <https://doi.org/10.4269/ajtmh.2009.80.410>
71. **Manne J, Snively C, Ramsey J, Salgado M, Barnighausen T, Reich M**. Barriers to treatment access for CD in Mexico. *PLoS Negl Trop Dis*. 2013; 7(10): e2488. DOI: <https://doi.org/10.1371/journal.pntd.0002488>
72. **Manne-Goehler J, Reich M, Wirtz V**. Access to Care for CD in the United States: A Health Systems Analysis. *Am J Trop Med Hyg*. 2015; 93(1): 108–113. DOI: <https://doi.org/10.4269/ajtmh.14-0826>
73. **Stimpert K, Montgomery S**. Physician Awareness of CD. *Emerg Infect Dis*. 2010; 16(5): 871–872. DOI: <https://doi.org/10.3201/eid1605.091440>
74. **Pérez-Molina JA, Molina I**. Chagas Disease. *The Lancet*. 2018; 391(10115): 82–94. DOI: [https://doi.org/10.1016/S0140-6736\(17\)31612-4](https://doi.org/10.1016/S0140-6736(17)31612-4)
75. **De Rissio A, Riarte A, García M, Esteva M, Quaglino M, Ruiz A**. Congenital *Trypanosoma cruzi* infection. Efficacy of its monitoring in an urban reference health center in a non-endemic area of Argentina. *Am J Trop Med Hyg*. 2010; 82(5): 838–845. DOI: <https://doi.org/10.4269/ajtmh.2010.08-0383>
76. **Bern C, Verastegui M, Gilman R**, et al. Congenital *Trypanosoma cruzi* transmission in Santa Cruz, Bolivia. *Clin Infect Dis*. 2009; 49(11): 1667–1674. DOI: <https://doi.org/10.1086/648070>

77. **Mora M, Sanchez Negrette O, Marco D**, et al. Early diagnosis of congenital *Trypanosoma cruzi* infection using PCR, hemoculture, and capillary concentration, as compared with delayed serology. *J Parasitol.* 2005; 91(6): 1468–1473. DOI: <https://doi.org/10.1645/GE-549R.1>
78. **Rivero R, Bisio M, Velazquez E**, et al. Rapid detection of *T. Cruzi* by colorimetric loop-mediated isothermal amplification (LAMP): A potential novel tool for the detection of congenital Chagas infection. *Diagn Microbiol Infect Dis.* 2017; 89: 26–28. DOI: <https://doi.org/10.1016/j.diagmicrobio.2017.06.012>
79. **Cura C, Ramírez J, Rodríguez M, Lopez-Albizu C, Irazu L, Scollo K, Sosa-Estani S.** Comparative Study and Analytical Verification of PCR Methods for the Diagnosis of Congenital CD. *J Mol Diagn.* 2017; 19(5): 673–681. DOI: <https://doi.org/10.1016/j.jmoldx.2017.05.010>
80. **Besuschio S** et al. Analytical sensitivity and specificity of a loop-mediated isothermal amplification (LAMP) kit prototype for detection of *Trypanosoma cruzi* DNA in human blood samples. *PLoS Negl Trop Dis.* 2017; 11(7). DOI: <https://doi.org/10.1371/journal.pntd.0005779>
81. **Schijman AG.** Molecular diagnosis of *Trypanosoma cruzi*. *Acta Tropica.* 2018; 184: 59–66. DOI: <https://doi.org/10.1016/j.actatropica.2018.02.019>
82. **Riarte A, Luna C, Sabatiello R, Sinagra A, Schiavelli R, De Rissio A, Maiolo E, García MM, Jacob N, Pattin M, Lauricella M, Segura EL, Vázquez M.** Chagas' Disease in Patients with Kidney Transplants: 7 Years of Experience, 1989–1996. *Clinical Infectious Diseases.* 1999; 29(3): 561–567. DOI: <https://doi.org/10.1086/598634>
83. **Altchek J, Moscatelli G, Moroni S**, et al. Adverse Events After the Use of Benznidazole in Infants and Children With CD. *Pediatrics.* 2011; 127. DOI: <https://doi.org/10.1542/peds.2010-1172>
84. **Pan American Health Organization.** Guidelines for the diagnosis and treatment of Chagas disease. Washington, DC: PAHO; 2019.
85. **Cerisola JA.** Chemotherapy of Chagas' infection in man. *Scientific Publication PAHO.* 1977; 347: 35–47.
86. **Russomando G, De Tomassone M, De Guillen I**, et al. Treatment of congenital chagas' disease diagnosed and followed up by the polymerase chain reaction. *American Journal of Tropical Medicine and Hygiene.* 1998; 59(3): 487–491. DOI: <https://doi.org/10.4269/ajtmh.1998.59.487>
87. **Schijman AG, Altchek J, Burgos JM**, et al. Aetiological treatment of congenital Chagas' disease diagnosed and monitored by the polymerase chain reaction. *Journal of Antimicrobial Chemotherapy.* 2003; 52(3): 441–449. DOI: <https://doi.org/10.1093/jac/dkg338>
88. **Moscatelli G, Moroni S, García Bournissen F, González N, Ballering G, Schijman A, Corral R, Bisio M, Freilij H, Altchek J.** Longitudinal follow up of serological response in children treated for CD. *PLoS Negl Trop Dis.* 2019; 13(8). DOI: <https://doi.org/10.1371/journal.pntd.0007668>
89. **Alonso-Vega C, Billot C, Torrico F.** Achievements and challenges upon the implementation of a program for national control of congenital Chagas in Bolivia: results 2004–2009. *PLoS Negl Trop Dis.* 2013; 7(7). DOI: <https://doi.org/10.1371/journal.pntd.0002304>
90. **Moya PR, Paolasso RD, Blanco S.** Treatment of CD with nifurtimox during the first months of life. *Medicina.* 1985; 45(5): 553–558.
91. **Blanco SB, Segura EL, Cura EN**, et al. Congenital transmission of *Trypanosoma cruzi*: An operational outline for detecting and treating infected infants in northwestern Argentina. *Tropical Medicine and International Health.* 2000; 5(4): 293–301. DOI: <https://doi.org/10.1046/j.1365-3156.2000.00548.x>
92. **Fragata-Filho A, França F, Fragata C**, et al. Evaluation of parasiticide treatment with benznidazole in the electrocardiographic, clinical, and serological evolution of CD. *PLoS Negl Trop Dis.* 2016; 10(3). DOI: <https://doi.org/10.1371/journal.pntd.0004508>
93. **Morillo C, Marin-Neto J, Avezum A, Sosa-Estani S, Rassi A Jr., Rosas F**, et al. Randomized Trial of Benznidazole for Chronic Chagas' Cardiomyopathy. *N Engl J Med.* 2015; 373(4): 1295–1306. DOI: <https://doi.org/10.1056/NEJMoa1507574>
94. **Sperandio da Silva G, Mediano M, Hasslocher-Moreno A, Holanda M, Silvestre de Sousa A, Sanguis L**, et al. Benznidazole treatment safety: The Medecins Sans Frontieres experience in a large cohort of Bolivian patients with Chagas' disease. *J Antimicrob Chem.* 2017; 72(9): 2596–2601. DOI: <https://doi.org/10.1093/jac/dkx180>
95. **García-Bournissen F, Moroni S, Marson M, Moscatelli G, Mastrantonio G, Bisio M, Cornou L, Ballering G, Altchek J.** Limited infant exposure to benznidazole through breast milk during maternal treatment for CD. *Arch Dis Child.* 2015; 100(1): 90–94. DOI: <https://doi.org/10.1136/archdischild-2014-306358>

96. **Moroni S, Marson M, Moscatelli G, Mastrantonio G, Bisio M, Gonzalez N, Ballering G, Altcheh J, García-Bournissen F.** Negligible exposure to nifurtimox through breast milk during maternal treatment for CD. *PLoS Negl Trop Dis.* 2019; 13(8). DOI: <https://doi.org/10.1371/journal.pntd.0007647>
97. **Sguassero Y, Cuesta C, Roberts K, Hicks E, Comandé D, Ciapponi A, Sosa-Estani S.** Course of Chronic *Trypanosoma cruzi* Infection after Treatment Based on Parasitological and Serological Tests: A Systematic Review of Follow-Up Studies. *PLoS One.* 2015; 10(10): e0139363. DOI: <https://doi.org/10.1371/journal.pone.0139363>
98. **Sguassero Y, Roberts K, Harvey G, Comandé D, Ciapponi A, Cuesta C, Aguiar C, Castro A, Danesi E, de Andrade A, de Lana M, Escribà J, Fabbro D, Fernandes C, Flores-Chávez M, Hasslocher-Moreno A, Jackson Y, Lacunza C, Machado-de-Assis G, Maldonado M,** et al. Course of serological tests in treated subjects with chronic *Trypanosoma cruzi* infection: A systematic review and meta-analysis of individual participant data. *Int J Infect Dis.* 2018; 73: 93–101. DOI: <https://doi.org/10.1016/j.ijid.2018.05.019>
99. **Sosa-Estani S, Segura E.** Etiological treatment in patients infected by *Trypanosoma cruzi*: Experiences in Argentina. *Current opinion in infectious diseases.* 2006; 19(6): 583–587. DOI: <https://doi.org/10.1097/01.qco.0000247592.21295.a5>
100. **Müller Kratz J, Garcia Bournissen F, Forsyth CJ, Sosa-Estani S.** Clinical and pharmacological profile of benznidazole for treatment of CD. *Expert Review of Clinical Pharmacology.* 2018; 11(10): 943–957. DOI: <https://doi.org/10.1080/17512433.2018.1509704>
101. **Sosa Estani S, Segura E, Ruiz A, Velazquez E, Porcel B, Yampotis C.** Efficacy of chemotherapy with benznidazole in children in the indeterminate phase of CD. *The American Journal of Tropical Medicine and Hygiene.* 1998; 59(4): 526–529. DOI: <https://doi.org/10.4269/ajtmh.1998.59.526>
102. **Viotti R, Vigliano C, Lococo B,** et al. Long-term cardiac outcomes of treating chronic CD with benznidazole versus no treatment: A nonrandomized trial. *Annals of Internal Medicine.* 2006; 144(10): 724–734. DOI: <https://doi.org/10.7326/0003-4819-144-10-200605160-00006>
103. **Fabbro D, Streiger M, Arias E, Bizai M, del Barco M, Amicone N.** Trypanocide treatment among adults with chronic CD living in Santa Fe city (Argentina), over a mean follow-up of 21 years: Parasitological, serological and clinical evolution. *Revista da Sociedade Brasileira de Medicina Tropical.* 2007; 40(1): 1–10. DOI: <https://doi.org/10.1590/S0037-86822007000100001>
104. **Cardoso C, Ribeiro A, Oliveira C, Oliveira L, Ferreira A, Bierrenbach A, Silva J, Colosimo E, Ferreira J, Lee T, Busch M, Reingold A, Sabino E.** Beneficial effects of benznidazole in CD: NIH SaMi-Trop cohort study. *PLoS Negl Trop Dis.* 2018; 12(11). DOI: <https://doi.org/10.1371/journal.pntd.0006814>
105. **Morillo C, Waskin H, Sosa-Estani S,** et al. Benznidazole and Posaconazole in Eliminating Parasites in Asymptomatic *T. Cruzi* Carriers: The STOP-CHAGAS Trial. *Journal of the American College of Cardiology.* 2017; 69(8): 939–947. DOI: <https://doi.org/10.1016/j.jacc.2016.12.023>
106. **Torrice F, Gascon J, Ortiz L,** et al. Treatment of adult chronic indeterminate CD with benznidazole and three E1224 dosing regimens: A proof-of-concept, randomised, placebo-controlled trial. *The Lancet Infectious Diseases.* 2018; 18(4): 419–430. DOI: [https://doi.org/10.1016/S1473-3099\(17\)30538-8](https://doi.org/10.1016/S1473-3099(17)30538-8)
107. **Molina I, Gómez I, Prat J, Salvador F,** et al. Randomized Trial of Posaconazole and Benznidazole for Chronic Chagas' Disease. *New England Journal of Medicine.* 2014; 370(20): 1899–1908. DOI: <https://doi.org/10.1056/NEJMoa1313122>
108. **DNDi.** Chagas BENDITA Study Briefing Document. 2019.
109. **Forsyth C, Hernandez S, Flores C, Roman M, Nieto J, Marquez G,** et al. "It's Like a Phantom Disease": Patient Perspectives on Access to Treatment for CD in the United States. *The American Journal of Tropical Medicine and Hygiene.* 2018; 98(3): 735–741. DOI: <https://doi.org/10.4269/ajtmh.17-0691>
110. **Andrade J,** et al. Latin American guidelines for the diagnosis and treatment of Chagas' heart disease: Executive summary. *Bras Cardiol.* 2011; 96(43).
111. **Filigheddu M, Górgolas M, Ramos J.** Orally-transmitted CD. *Med Clin (Barc).* 2017; 148(3): 125–131. DOI: <https://doi.org/10.1016/j.medcli.2016.10.038>
112. **Botoni FA,** et al. A Randomized Trial of Carvedilol after Renin-Angiotensin System Inhibition in Chronic Chagas Cardiomyopathy. *American Heart Journal.* 2007; 153(4). DOI: <https://doi.org/10.1016/j.ahj.2006.12.017>
113. **Novartis Pharmaceuticals.** Efficacy and Safety of Sacubitril/Valsartan Compared With Enalapril on Morbidity, Mortality, and NT-proBNP Change in Patients With Chagas Cardiomyopathy (PARACHUTE-HF). *ClinicalTrials.gov.* 2019.

114. **Fernandes F.** Colchicine for Patients With Chagas' Disease (B1 Stage) (COACH). University of Sao Paulo General Hospital; 2018.
115. **Cardoso R, Macedo F, Garcia D, Benjo A, Aguilar D,** et al. Chagas cardiomyopathy is associated with higher incidence of stroke: A meta-analysis of observational studies. *J Card Fail.* 2014; 20(12): 931–938. DOI: <https://doi.org/10.1016/j.cardfail.2014.09.003>
116. **de Sousa A, Xavier S, de Freitas G, Hasslocher-Moreno A.** Prevention strategies of cardioembolic ischemic stroke in Chagas' disease. *Arq Bras Cardiol.* 2008; 91(5): 306–310. DOI: <https://doi.org/10.1590/S0066-782X2008001700004>
117. **Villar J, Rodriguez D.** A Trial Testing Amiodarone in Chagas Cardiomyopathy (ATTACH). Identification No. NCT03193749, 2017. [Online]. Available: <https://clinicaltrials.gov/ct2/show/NCT03193749>.
118. **Martinelli M.** Amiodarone Against ICD Therapy in Chagas Cardiomyopathy for Primary Prevention of Death (CHAGASICS). Identification No. NCT01722942. 2014. [Online]. Available: <https://clinicaltrials.gov/ct2/show/NCT01722942>.
119. **Kransdorf E, Zakowski P, Kobashigawa J.** CD in solid organ and heart transplantation. *Curr Opin Infect Dis.* 2014; 27(5): 418–424. DOI: <https://doi.org/10.1097/QCO.0000000000000088>
120. **Clark E, Sherbuk J, Okamoto E, Jois M, Galdos-Cardenas G,** et al. Hyperendemic CD and the Unmet Need for Pacemakers in the Bolivian Chaco. *PLOS Neglected Tropical Diseases.* 2014; 8(6): e2801. DOI: <https://doi.org/10.1371/journal.pntd.0002801>
121. **Jackson Y, Alirol E, Getaz L, Wolff H, Combescure C, Chappuis F.** Tolerance and safety of nifurtimox in patients with chronic CD. *Clin Infect Dis.* 2010; 51(10): e69–e75. DOI: <https://doi.org/10.1086/656917>
122. **Hidron A, Gilman R, Justiniano J, Blackstock A, LaFuente C,** et al. Chagas Cardiomyopathy in the Context of the Chronic Disease Transition. *PLOS Neglected Tropical Diseases.* 2010; 4(5): e688. DOI: <https://doi.org/10.1371/journal.pntd.0000688>
123. **de Almeida AE,** et al. Co-infection *Trypanosoma cruzi*/HIV: Systematic review (1980–2010) *Rev Soc Bras Med Trop.* 2011; 44(6): 762–770. DOI: <https://doi.org/10.1590/S0037-86822011000600021>
124. **Ribeiro I, Sevcsik A, Alves F, Diap G, Don R, Harhay M, Chang S, Pecoul B.** New, improved treatments for CD: From the R&D pipeline to the patients. *PLoS Negl Trop Dis.* 2009; 3(7): e484. DOI: <https://doi.org/10.1371/journal.pntd.0000484>
125. **Kong A, Warren M, Edwards D, Karrar K, Iyer JK.** Are pharmaceutical companies making progress when it comes to global health? *Access to Medicine Foundation.* May 2019.
126. **Pratt B, Loff B.** Linking Research to Global Health Equity: The Contribution of Product Development Partnerships to Access to Medicines and Research Capacity Building. *American Journal of Public Health.* 2013; 103(11): 1968–978. DOI: <https://doi.org/10.2105/AJPH.2013.301341>
127. **Sánchez-Valdéz F, Padilla A.** In Situ Detection of Dormant *Trypanosoma cruzi* Amastigotes Using Bioluminescent-Fluorescent Reporters. *Methods Mol Biol.* 2019; 1955: 179–186. DOI: https://doi.org/10.1007/978-1-4939-9148-8_13
128. **Beaumier C, Gillespie P, Strych U,** et al. Status of vaccine research and development of vaccines for CD. *Vaccine.* 2016; 34: 2996–3000. DOI: <https://doi.org/10.1016/j.vaccine.2016.03.074>
129. **Jones K, Versteeg L, Damania A,** et al. Vaccine-linked chemotherapy improves benznidazole efficacy for acute CD. *Infect Immun.* 2018; 86. DOI: <https://doi.org/10.1128/IAI.00876-17>
130. **Pinazo M-J, Pinto J, Ortiz L,** et al. A strategy for scaling up access to comprehensive care in adults with Chagas disease in endemic countries: The Bolivian Chagas Platform. *PLOS Neglected Tropical Diseases.* 2017. DOI: <https://doi.org/10.1371/journal.pntd.0005770>
131. **Batista C, Forsyth C, Herazo R, Certo M, Marchiol A.** A four-step process for building sustainable access to diagnosis and treatment of Chagas disease. *Rev Panam Salud Publica.* 2019; 43: e74. DOI: <https://doi.org/10.26633/RPSP.2019.74>
132. **Pennington P, Juárez J, Arrivillaga M, De Urioste-Stone S, Doktor K, Bryan J, Cerdón-Rosales C.** Towards Chagas disease elimination: Neonatal screening for congenital transmission in rural communities. *PLoS Negl Trop Dis.* 2017; 11(9): e0005783. DOI: <https://doi.org/10.1371/journal.pntd.0005783>
133. **Engam D, Engam B, Engam K.** Chagas Heart Disease Pathogenesis: One Mechanism or Many? *Current Molecular Medicine.* 2008; 8(6): 510–518. DOI: <https://doi.org/10.2174/156652408785748004>

134. **Tanowitz HB**, et al. Developments in the Management of Chagas Cardiomyopathy. *Expert Review of Cardiovascular Therapy*. 2015; 13(12): 1393–1409. DOI: <https://doi.org/10.1586/14779072.2015.1103648>
135. **Chevillard C**, et al. Disease Tolerance and Pathogen Resistance Genes May Underlie *Trypanosoma cruzi* Persistence and Differential Progression to Chagas Disease Cardiomyopathy. *Frontiers*. 2018. DOI: <https://doi.org/10.3389/fimmu.2018.02791>
136. **Pinazo M, Thomas M, Bua J, Perrone A, Schijman A, Viotti R, Ramsey J, Ribeiro I, Sosa-Estani S, Lopez M, Gascon J**. Biological markers for evaluating therapeutic efficacy in Chagas disease: A systematic review. *Expert Rev Anti Infect Ther*. 2014; 12(4): 479–496. DOI: <https://doi.org/10.1586/14787210.2014.899150>
137. **Albareda M, Laucella S**. Modulation of *Trypanosoma cruzi*-Specific T-Cell Responses after Chemotherapy for Chronic Chagas Disease. *Memórias Do Instituto Oswaldo Cruz*. 2015; 110(3): 414–421. DOI: <https://doi.org/10.1590/0074-02760140386>
138. **Albareda M**, et al. Distinct Treatment Outcomes of Antiparasitic Therapy in *Trypanosoma cruzi*-Infected Children Is Associated With Early Changes in Cytokines, Chemokines, and T-Cell Phenotypes. *Frontiers in Immunology*. 2018; 9. DOI: <https://doi.org/10.3389/fimmu.2018.01958>
139. **Sánchez-Valdéz F, Padilla A, Wang W, Orr D, Tarleton R**. Spontaneous dormancy protects *Trypanosoma cruzi* during extended drug exposure. *Elife*. 2018; 7. DOI: <https://doi.org/10.7554/eLife.34039>
140. **Mendoza N, de la Torre Avilé L**. Chagas Workshop. *Pasa la Voz project*. 2016.
141. **Sanmartino M, Saavedra A, Prat J, Barba M, Albajar-Viñas P**. Que no tengan miedo de nosotros: el Chagas según los propios protagonistas. *Interface – Comunicação, Saúde, Educação*. 2015; 19: 1063–1075. DOI: <https://doi.org/10.1590/1807-57622014.1170>
142. **Forsyth C**. I Cannot Be Worried: Living with CD in Tropical Bolivia. *PLoS Negl Trop Dis*. 2017; 11(1). DOI: <https://doi.org/10.1371/journal.pntd.0005251>
143. **Sanchez D, Traina M, Hernandez S, Smer A, Khamag H, Meymandi S**. CD awareness among Latin American immigrants living in Los Angeles, California. *The American Journal of Tropical Medicine and Hygiene*. 2014; 91(5): 915–919. DOI: <https://doi.org/10.4269/ajtmh.14-0305>
144. **Ozaki Y, Guariento M, de Almeida E**. Quality of life and depressive symptoms in CD patients. *Qual Life Res*. 2011; 20(1): 133–138. DOI: <https://doi.org/10.1007/s11136-010-9726-1>
145. **Ventura-Garcia L, Roura M, Pell C, Posada E, Gascón J, Aldasoro E**, et al. Socio-Cultural Aspects of CD: A Systematic Review of Qualitative Research. *PLOS Neglected Tropical Diseases*. 2013; 7(9): e2410. DOI: <https://doi.org/10.1371/journal.pntd.0002410>
146. **Oliveira W, Jr**. All-around care for patients with CD: a challenge for the XXI century. *Memorias do Instituto Oswaldo Cruz*. 2009; 104(1): 181–186. DOI: <https://doi.org/10.1590/S0074-02762009000900024>
147. 2nd Brazilian Consensus on CD. In *Epidemiol*. Brasília; 2015.
148. Manual de procedimiento para la atención de pacientes con enfermedad de Chagas. *Plan Nacional de Control y Prevención de Enfermedad de Chagas Ministerio de Salud de Chile*; 2017.
149. **Ministerio de Salud de la Nación**. Chagas: Atención del paciente infectado con *Trypanosoma cruzi*. BAJA: Dirección de Epidemiología, Buenos Aires; 2018.
150. Guia para el Diagnóstico, Atención y Manejo Clínico de la Enfermedad de Chagas en Venezuela. *República Bolivariana de Venezuela Ministerio del Poder Popular para la Salud*; 2014.
151. **Ministério da Saúde, Secretaria de Ciência, Tecnologia e Insumos Estratégicos**. Protocolo Clínico e Diretrizes Terapêuticas em Doença de Chagas. Brazilian Ministry of Health, 2018.
152. **Pérez- Molina J, Rodríguez-Guardado S, Soriano A, Pinazo M**, et al. Guidelines on the treatment of chronic coinfection by *Trypanosoma cruzi* and HIV outside endemic areas. *HIV Clin Trials*. 2011; 12(6): 287–298. DOI: <https://doi.org/10.1310/hct1206-287>
153. **Pinazo M, Cañas E, Elizalde J, Garcia M**, et al. Diagnosis, management and treatment of chronic Chagas' gastrointestinal disease in areas where *Trypanosoma cruzi* infection is not endemic. *Gastroenterol Hepatol*. 2010; 33(3): 191–200. DOI: <https://doi.org/10.1016/j.gastrohep.2009.07.009>
154. **Gascón J, Albajar P, Cañas E, Flores M**, et al. Diagnosis, management and treatment of chronic Chagas' heart disease in areas where *Trypanosoma cruzi* infection is not endemic. *Rev EspCardiol*. 2007; 60(3): 285–293. DOI: [https://doi.org/10.1016/S1885-5857\(07\)60153-4](https://doi.org/10.1016/S1885-5857(07)60153-4)
155. **Abbott, AstraZeneca, Bayer, Becton Dickinson, Bill & Melinda Gates Foundation**, et al. London Declaration on Neglected Tropical Diseases. London; 2012.
156. **World Health Organization**. Accelerating work to overcome the global impact of Neglected Tropical Diseases. A Roadmap for Elimination. *WHO*; 2012.

157. **Mathers C, Ezzati M, Lopez A.** Measuring the burden of neglected tropical diseases: The global burden of disease framework. *PLoS Negl Trop Dis.* 2007; 1: e114. DOI: <https://doi.org/10.1371/journal.pntd.0000114>
158. **TDR, the Special Programme for Research and Training in Tropical Diseases.** MosquitiaMed: Shortening Distances Through Telemedicine. *Social Innovation in Health Initiative*; 2019. [Online].
159. **Marcolino M, Palhares D, Ferreira L, Ribeiro A.** Electrocardiogram and CD: A large population database of primary care patients. *Global Heart.* 2015; 10(3): 167–172. DOI: <https://doi.org/10.1016/j.gh.heart.2015.07.001>
160. **Nascimento BR, Beaton AZ, Nunes MCP, et al.** PROVAR+ (Programa de Rastreamento da Valvopatia Reumática and Other Cardiovascular Diseases) investigators. Integration of echocardiographic screening by non-physicians with remote reading in primary care. *Heart.* 2019; 105(4): 283–90. DOI: <https://doi.org/10.1136/heartjnl-2018-313593>
161. **Cormick G, Ciganda A, Cafferata M, Ripple M, et al.** Text message interventions for follow up of infants born to mothers positive for CD in Tucumán, Argentina: A feasibility study. *BMC Res Notes.* 2015; 8. DOI: <https://doi.org/10.1186/s13104-015-1498-9>
162. **Jiang F, Jiang Y, Zhi H, Dong Y, Li H, Ma S, et al.** Artificial intelligence in healthcare: past, present and future. *Stroke and Vascular Neurology.* 2017; 2(4): 230–243. DOI: <https://doi.org/10.1136/svn-2017-000101>

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