

# Measures to the Control of *Trypanosoma cruzi* in Buffaloes

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## ABSTRACT

*Trypanosoma (T.) cruzi*, the causative agent of Chagas disease, poses a significant threat to both human and animal health, particularly in regions where the vector (*triatomine* bugs) are present. Buffaloes are susceptible to *T. cruzi* infection, and their role as reservoir hosts for the parasite raises concerns about transmission to humans and other animals. This article aims to explore various control strategies to combat *T. cruzi* infection in buffaloes. It provides information about preventing transmission, diagnosing infected animals, and enhancing treatment modalities, ultimately helping in the overall control of this neglected tropical disease.

### Introduction:

*Trypanosoma (T.) cruzi* is a parasitic protozoan that causes *Chagas* disease. This disease is a neglected tropical disease that affects millions of people worldwide. The primary vectors of *T. cruzi* are *triatomine* bugs in several mammalian species including buffalo. Controlling *T. cruzi* in buffalo is not important only for preventing the transmission to humans but also for other animals (1, 2).

### Key points for controlling *T. cruzi*

#### 1. Vector control

*Triatomine* bugs are the primary vectors of *T. cruzi*, controlling the populations of *triatomine* is vital to decrease the passing of *T. cruzi* in other animals. Proper policies are helpful to control vector transmission such as insecticide-treated bed nets, covering the wall and other surfaces with residual insecticides, and environmental management methods can help to decrease contact between buffaloes and *triatomines*, decreasing the risk of passing *triatomine* in animals (3-4).

#### 2. Vaccination program

Developing a vaccine plays an important role in controlling *Chagas* disease. Research should need to understand the nature of *T. cruzi* antigens that help in developing a vaccine for Buffaloes and other hosts. This vaccine not only protects buffaloes but also decreases the risk of passing to humans (5).

#### 3. Public awareness and education

Training of buffalo farmers and the locals about the risks of *T. cruzi* infection and the importance of repressive measures is about the disease, its transmission routes, and the role of buffalo in maintaining the parasite's life cycle. This can include proper waste management, the use of insecticide-treated nets for buffalo protection, and reporting any doubtful cases to veterinary authorities (6).

#### 4. Management of infected individuals

If a buffalo is recognized as infected with *T. cruzi*, proper measures should be taken to stop further transmission. Infected animals should be isolated from non-infected buffalo and treated if possible. Additionally, buffaloes have direct contact with infected ones. So close monitoring is helpful to detect new cases early (7).

#### 5. Chemotherapy

Chemotherapeutic agents, such as *benznidazole* and *nifurtimox*, are the most commonly used drugs for the cure of *T. cruzi* infection. A good combination of therapies and careful monitoring of treatment show effective results in decreasing parasite load in infected buffaloes and stopping disease development (7).

### Conclusions

To successfully control *T. cruzi* infection in buffaloes, an integrated approach involving vector control, vaccination research, accurate diagnosis, and appropriate treatment is essential for their control. Implementation of sustainable control strategies, along with ongoing research efforts, will lead to better management of *T. cruzi* infection in buffalo populations and reduce the risk of transmission to humans and other animals.

### References

- [1] Hardison JL, Wrightsman RA, Carpenter PM, Kuziel WA, Lane TE, Manning JE. The CC chemokine receptor 5 is important in the control of parasite replication and acute cardiac inflammation following infection with *Trypanosoma cruzi*. *Infection and immunity*. 2006Jan;74(1):135-43.
- [2] Davila AM, SILVA RA. Animal trypanosomiasis in South America: current status, partnership, and information technology. *Annals of the New York Academy of Sciences*. 2000 Dec;916(1):199-212.
- [3] Jaimes-Dueñez J, Triana-Chávez O, Mejía-Jaramillo AM. Spatial-temporal and phylogeographic characterization of *Trypanosoma* spp. in cattle (*Bos taurus*) and buffaloes (*Bubalus bubalis*) reveals transmission dynamics of these parasites in Colombia. *Veterinary parasitology* 2018Jan15;249:30-42.

- [4] Nuryady MM, Widayanti R, Nurcahyo RW, Fadrijnatha B. Characterization and phylogenetic analysis of multidrug-resistant protein-encoding genes in *Trypanosoma evansi* isolated from buffaloes in Ngawi district, Indonesia. *Veterinary World*. 2019 Oct;12(10):1573.
- [5] Garcia H, Garcia ME, Perez H, Mendoza-Leon A. The detection and PCR-based characterization of the parasites causing trypanosomiasis in water-buffalo herds in Venezuela. *Annals of Tropical Medicine & Parasitology*. 2005 Jun 1;99(4):359-70.
- [6] Villanueva MA, Mingala CN, Tubalinal GA, Gaban PB, Nakajima C, Suzuki Y. Emerging infectious diseases in water buffalo: An economic and public health concern. *IntechOpen*; 2018 Feb 21.
- [7] Jaimes-Dueñez J, Triana-Chávez O, Mejía-Jaramillo AM. Spatial-temporal and phylogeographic characterization of *Trypanosoma* spp. in cattle (*Bos taurus*) and buffaloes (*Bubalus bubalis*) reveals transmission dynamics of these parasites in Colombia. *Veterinary parasitology*. 2018 Jan 15;249:30-42.