



Clinical Case

**Chagas' reactivation in a heart transplant patient**

Tânia Mara Varejão Strabelli, PhD  
Director of Infection Control Unit  
Heart Institute of São Paulo School of Medicine

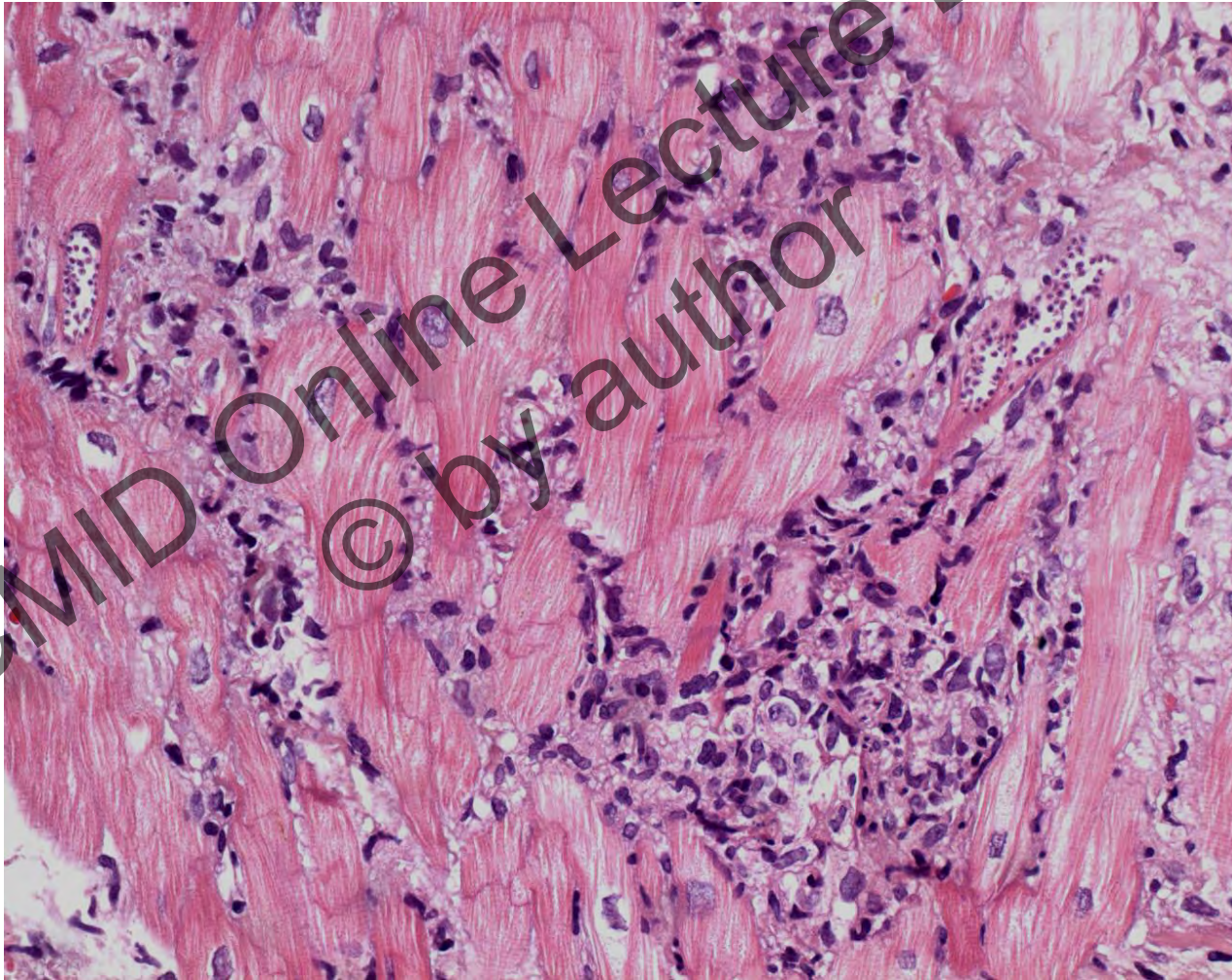
# Clinical case

- A 51-year-old man, diabetes, hypertension
- **Heart transplantation due to Chagas' cardiomyopathy**
  - July, 2013
  - Heart Institute (InCor)
- Immunosuppression
  - Methylprednisolone
  - cyclosporine, azathioprine, prednisone
- Postoperative infection: none
- Prophylaxis: none
- CMV antigenemia: negative
- Endomyocardial biopsy: no rejection
- Discharged on day 23+

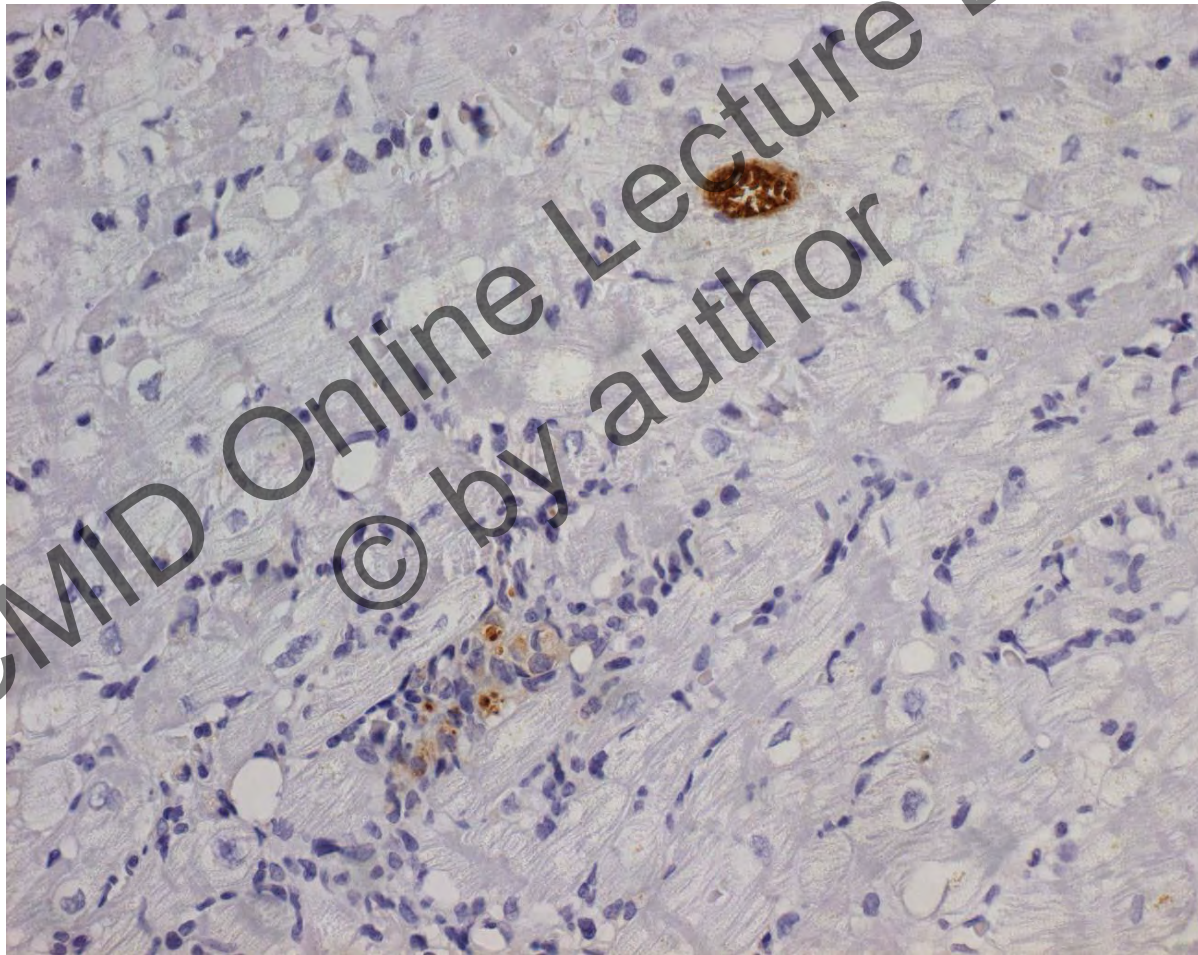
# Clinical case

- Hospital admission on day 60+
- Cellulitis in left leg, no fever
  - Vancomycin – 4 days; teicoplanin – 10 days
- Routine endomyocardial biopsy revealed lymphocytic myocarditis with the presence of *T. cruzi* amastigotes
- No clinical signs or symptoms of congestive heart failure
- Ecocardiography – ejection fraction=65%
- Patient was treated with benznidazole 10mg/kg/day for 60 days

# Endomyocardial biopsy specimen (HE stain)



# Endomyocardial biopsy specimen (immunohistochemistry)



# Clinical case

- Endomyocardial biopsy was repeated after 20 days: chronic mononuclear myocarditis with interstitial fibrosis and no parasites
- Patient was discharged from hospital using cyclosporine (100mg 12/12h), prednisone 20mg/d, azathioprine 120mg/day (reduced dose)
- CMV antigenemia – negative (repeated twice during hospital stay)

# Clinical case

- EB (november 2013) revealed acute rejection (2R AMR 0) and the patient was **readmitted for treatment**
- Methylprednisolone 500mg IV for 3 days
- MMF, tacrolimus, prednisone
- Patient was discharged after a new EB was free of rejection signs.

## Discussion

- This was a case of Chagas' disease reactivation after heart transplantation.
- Frequency at our hospital = 26,5%
- Risk factors associated with reactivation
  - events resulting in greater immunosuppression status



# Discussion

## Risk factors:

- Number of rejection episodes (  $p=0,011$  )
- Neoplasms (  $p=0,009$  )
- Use of mycophenolate mofetil (  $p=0,049$  ) – azathioprine or low dosis MMF

•Campos SV, Strabelli TMV et als. Risk factors for Chagas' disease reactivation after heart transplantation. J Heart Lung Transplant, 2008

•Bacal F, Silva CP et als. Mycophenolate Mofetil Increased Chagas Disease Reactivation in Heart Transplanted Patients: Comparison Between Two Different Protocols. Am J Transplant , 2005.

# Discussion

## Chagas' disease reactivation presentations:

- Cutaneous lesions
- Myocarditis
- Central nervous system lesions are infrequent in heart transplants

The diagnosis of reactivation requires detection of circulating trypomastigotes in the peripheral blood; quantitative polymerase chain reaction (PCR) on serial specimens is a useful technique to show rising parasite load. It provides the earliest and most sensitive indicator of reactivation.

•Bacal F, Silva CP et als. Mychophenolate Mofetil Increased Chagas Disease Reactivation in Heart Transplanted Patients:Comparison Between Two Different Protocols. Am J Transplant , 2005.

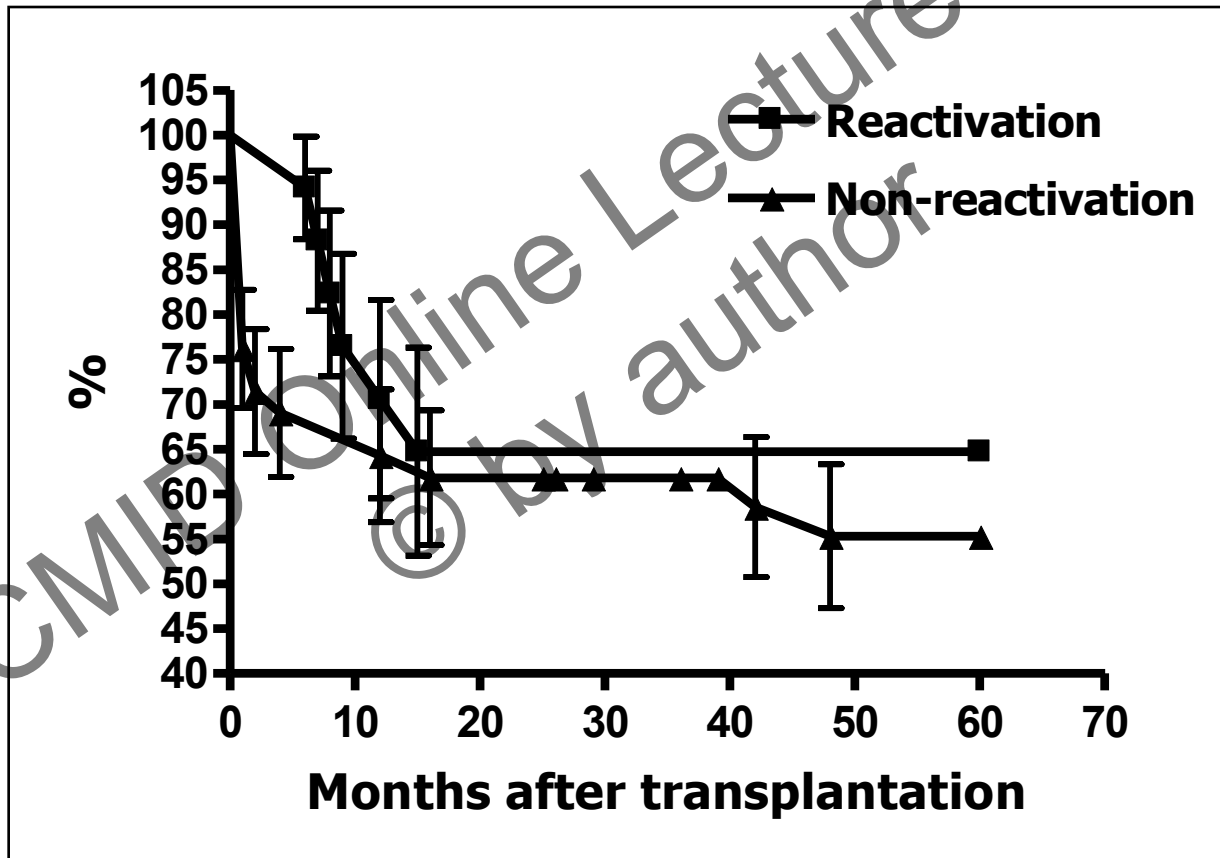
# Discussion

## Monitoring heart transplant patients at regular intervals and after treating rejection episodes

### Treatment options

- Benznidazol
- Allopurinol
- Nifurtimox

# Survival curve for Chagas' disease reactivation group x non-reactivation group (Kaplan-Meier method; $p = 0,46$ Longrank test).



Campos SV, Strabelli TMV et als. Risk factors for Chagas' disease reactivation after heart transplantation. J Heart Lung Transplant, 2008

# Summary

- Chagas' disease is a common cause of heart failure in Brazil
- Chronic disease could reactivate after periods of higher immunosuppression, but treatment is very efficient and safe in heart transplant recipients
- We have to define better methods to monitor disease reactivation and always look for the parasite in tissues.

**THANK YOU FOR ATTENTION!**