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the Brazilian Society of Tropical Medicine XLVIII Congress of and

XXVIII Brazilian Annual Meeting of Applied Research on Chagas Disease, XVI Brazilian Annual Meeting of Applied Research on Leishmaniasis and III Latin American Congress on Travel Medicine

Certificate

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This is to certify that Saraiva, RM; Junqueira, ACV; Costa, FAC; Boia, MN.

Congress of the Brazilian Society of Tropical Medicine, held in Rio de Janeiro from September 23 to 27, has attended the XVIII International Congress for Tropical Medicine and Malaria and XLVIII

Poster Presentation: Long-term evaluation of etiologic treatment with benznidazole in patier with indeterminate chronic Chagas' disease 2012, as

Rio de Janeiro, September 27, 2012.

Professor Pierre Ambroise-Thomas resident of the IFTM

President of the XVIII ICTMM

President of the Scientific Committee of the XVIII ICTMM

Professor Carlos Henrique Nery Costa President of the SBMT

also positive for *T. cruzi* in PCR technique. **Conclusion**: Dogs of the surveyed area are roosting *Leishmania* spp. and *T. cruzi*, being an important reservoir for these protozoosis, enabling the domestic cycle of these parasites. Once canine infection precede or have a significant relationship to human nfection, it should be emphasized the possibility of speading the disease among domestic animals in the evaluated regions, as well as starting a new focus for human disease. **E-mail:** larallyn@hotmail.com

Chagas 065- Long-term evaluation of etiologic treatment with benznidazole in patients with indeterminate chronic Chagas' disease

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ntroduction: The aim of this study was to assess the effect of benzonidazol treatment on the electrocardiographic, serological and parasitological evolution of patients with indeterminate form of Chagas disease. Material and Methods: We retrospectively analyzed a group of 62 patients that were reated with benzonidazol and compared their clinical findings with the findings of an untreated group n=62) of age-matched patients with indeterminate form of Chagas disease. Results: The frequency of electrocardiographic alterations in the treated group was followed for 118.3 ± 61.7 months (7333.7 patients-months) and in the untreated group was followed for 144.51 ± 49 months (8959.9 patientsnonths). The rate of disease progression from indeterminate to the cardiac form of Chagas disease pased on electrocardiographic findings was 12.9% (8/62) among treated patients and 16.1% (10/62) among untreated (p = 0.4). The incidence densities of this outcome was 1.09/1000 patients-months and 1.11/1000 patients-months in treated and untreated groups, respectively (relative risk = 0.98). There was no correlation between the progression to cardiac form and age, sex or place of birth. The serological iters of benzonidazol treated and untreated patients were followed for 88.7 \pm 49.6 months and 140.9 \pm 17.8 months, respectively. The titers were converted to a linear scale, where 1 represented a 1:40 dilution and 6 a 1:1280 dilution. The serological titer reduced significantly after benzonidazol treatment (4.07 perfore treatment vs. 2.85 post-treatment, p < 0.001). Such reduction was not observed among untreated persons (4.22 at the beginning of the follow-up vs. 4.11 at the end, p = 0.503). Despite the observed eduction of the serological titers after benzonidazol treatment, the post-treatment titers tended to ncrease throughout the follow-up period, mainly after 100-150 months. All treated patients were submitted to xenodiagnosis, before treatment, and 32 proved positive. After treatment, xenodiagnosis emained positive in only one patient. Main Conclusions: We concluded that: i) among studied patients here was no relationship between etiologic treatment and electrocardiographic progression, ii) treatment with benznidazole is associated with a significant reduction in serological titers, and iii) parasitaemia seems to be suppressed soon after specific treatment. E-mail: alejandro.hasslocher@ipec.fiocruz.br

Chagas 066- Benznidazole evaluation in the treatment of the experimental Chagas' disease

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ntroduction: Chagas' disease is caused by the flagellate parasite *Trypanosoma cruzi*. Although parasite ransmission by its natural vector has mostly been controlled, Chagas' disease still deserves attention rom researchers since many people remain infected and new infection routes have been described. New irugs must be tested and different diagnostic methods must be used to determinate the parasitological cure. The total parasite elimination is crucial, since its presence can trigger a new acute phase, mainly in he immunosuppressed patient. The present study proposes to evaluate the Benznidazole effectiveness and the efficacy of the diagnostic methodologies in the experimental Chagas' disease treatment. **Material and methods:** Swiss mice were infected intraperitoneally with 10⁴ *T. cruzi* trypomastigotes forms and in he early acute phase the treatment was initiated with Benznidazole at 100mg/kg body weight for 90 consecutive days, by oral route. Every other day, blood samples were collected from the tail vein for



Long-term evaluation of etiologic treatment with Benznidazole in patients with indeterminate chronic Chagas' disease.



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INTRODUCTION

The aim of this study was to assess the effect of benzonidazol treatment on the electrocardiographic, serological and parasitological evolution of patients with indeterminate form of Chagas' disease.

MATERIAL and METHODS

We retrospectively analyzed a group of 62 patients that were treated with benzonidazol and compared their clinical findings with the findings of an untreated group (n=62), both with indeterminate form of Chagas disease.

RESULTS

Variáveis	Evolução com surgimento de alterações eletrocardiográf icas (n = 18)	Manutenção na forma indeterminad a (n = 106)	p- valor
Idade	$35,2 \pm 9,6$	34,8 ± 6,7	0,82
Sexo			
Masculino, n (%)	12 (14,5)	64 (85,5)	0.61
Feminino, n (%)	6 (12,5)	42 (87,5)	0,61
Sorologia inicial, média ± desvio padrão*	4,22 ± 1,11	4,09 ± 1,19	0,67
Naturalidade			
Minas Gerais, Bahia ou Goiás, n (%)	8 / 65 (12,3)	57 / 65 (87,7)	
Pará ou Nordeste exceto Bahia, n (%)	9 / 37 (24,3)	28 / 37 (75,7)	
Rio de Janeiro, São Paulo, n (%)	0 /12 (0)	12 / 12 (100)	0,15
Rio Grande do Sul, Paraná, Mato Grosso do Sul ou Bolívia, n (%)	1 / 10 (10)	9 / 10 (90)	
Tratamento		54 / 62	
Sim	8 / 62 (12,9)	(87,1)	0,40
Não	10 / 62 (16,1)	52 / 62 (83,9)	,

Comparison of patients who developed cardiac form with those who remained with the indeterminate form of Chagas' disease,

electrocardiographic frequency of The alterations in the treated group was followed for 118.3 ± 61.7 months (7333.7 patientsmonths) and in the untreated group was followed for 144.51 ± 49 months (8959.9 patients-months). The rate of disease progression from indeterminate to the cardiac disease form of Chagas based electrocardiographic findings was 12.9% (8/62) among treated patients and 16.1% (10/62) among untreated (p = 0.4). The incidence densities of this outcome was 1.09/1000 patients-months and 1.11/1000 patientsmonths in treated and untreated groups, respectively (relative risk = 0.98). There was no correlation between the progression to cardiac form and age, sex or place of birth. The serological titers of benzonidazol treated and untreated patients were followed for 88.7 ± 49.6 months and 140.9 ± 47.8 months, respectively. The titers were converted to a linear scale, where 1 represented a 1:40 dilution and 6 a 1:1280 dilution. The serological titer reduced significantly after benzonidazol treatment (4.07 before treatment vs. 2.85 posttreatment, p < 0.001). Such reduction was not observed among untreated persons (4.22 at the beginning of the follow-up vs. 4.11 at the end, p = 0.503). Despite the observed reduction of the serological titers after benzonidazol treatment, the post-treatment titers tended to increase throughout the follow-up period, mainly after 100-150 months. All treated patients were submitted to xenodiagnosis, before treatment, and 32 proved positive. After treatment, xenodiagnosis remained positive in only one patient.

Características	Pacient	Pacientes	p-
Cur deter istreas	es	não tratado.	
	tratado	(n = 62)	r
	s com	(11)	
	benzoni		
	dazol		
	(n =		
1-15	62)		
Sorologia inicial*	4,00 ±	$4,23 \pm 1,16$	0,735
	1,87		,
Idade	$33,21 \pm$	$36,51 \pm$	0,009
	8,4	5,01	
Sexo (n [%] sexo	41	35 (56,5)	0,269
masculino)	(66,1)		
		11 1 1	
Naturalidade			
Minas Gerais, Bahia ou	31 (50)	34 (54,8)	
Goiás, n (%)			0,683
Pará ou Nordeste	18 (29)	19 (30,6)	
exceto Bahia, n (%)			
Rio de Janeiro, São	8 (12,9)	4 (6,5)	
Paulo, n (%)			
Rio Grande do Sul,	5 (8,1)	5 (8,1)	
Paraná, Mato Grosso	/		
do Sul ou Bolívia, n			
(%)			
Tempo de			
acompanhamento			
(evolução			
eletrocardiográfica)			
Meses (média ± desvio	$118,3 \pm$	4 4 4 - 40	
1.10000 (1110010 - 000.10	· '	$144,5 \pm 49$	0,001
padrão)	61,7	$144,5 \pm 49$	0,001
,	· '	$144,5 \pm 49$ 8959,9	0,001
padrão) Pacientes-meses	61,7	,	0,001
padrão) Pacientes-meses Tempo de	61,7	,	0,001
padrão) Pacientes-meses Tempo de acompanhamento	61,7	,	0,001
padrão) Pacientes-meses Tempo de acompanhamento (evolução sorológica)	61,7 7333,7	8959,9	0,001
padrão) Pacientes-meses Tempo de acompanhamento (evolução sorológica) Meses (média ± desvio	61,7 7333,7 88,7 ±	8959,9 140,9 ±	<
padrão) Pacientes-meses Tempo de acompanhamento (evolução sorológica)	61,7 7333,7	8959,9	
padrão) Pacientes-meses Tempo de acompanhamento (evolução sorológica) Meses (média ± desvio padrão)	61,7 7333,7 88,7 ±	8959,9 140,9 ±	<
padrão) Pacientes-meses Tempo de acompanhamento (evolução sorológica) Meses (média ± desvio padrão) Tempo de	61,7 7333,7 88,7 ±	8959,9 140,9 ±	<
padrão) Pacientes-meses Tempo de acompanhamento (evolução sorológica) Meses (média ± desvio padrão) Tempo de acompanhamento	61,7 7333,7 88,7 ±	8959,9 140,9 ±	<
padrão) Pacientes-meses Tempo de acompanhamento (evolução sorológica) Meses (média ± desvio padrão) Tempo de acompanhamento (evolução	61,7 7333,7 88,7 ±	8959,9 140,9 ±	<
padrão) Pacientes-meses Tempo de acompanhamento (evolução sorológica) Meses (média ± desvio padrão) Tempo de acompanhamento (evolução parasitológica,	61,7 7333,7 88,7 ±	8959,9 140,9 ±	<
padrão) Pacientes-meses Tempo de acompanhamento (evolução sorológica) Meses (média ± desvio padrão) Tempo de acompanhamento (evolução	61,7 7333,7 88,7 ±	8959,9 140,9 ±	<

Sample characteristics of 124 patients treated and not treated with benznidazole.

5,05 realizado**

CONCLUSIONS

We concluded that: i) among studied patients there was no relationship between etiologic treatment and electrocardiographic progression, ii) treatment with benznidazole is associated with a significant reduction in serological titers, and iii) parasitemia seems to be suppressed soon after specific treatment.

^{*} After conversion of serological evidence of dilution values in a linear scale in which the value 1 is a dilution of 1:80 and the value 6 is a dilution of 1:1280.