FAILURE OF BOTH AZITHROMYCIN AND ANTIMONY TO TREAT CUTANEOUS LEISHMANIASIS IN MANAUS, AM, BRAZIL

Alan César TEIXEIRA(1), Marcilene Gomes PAES(2), Jorge de Oliveira GUERRA(2), Aluízio PRATA(1) & Mario León SILVA-VERGARA(1)

SUMMARY

A non-randomized controlled clinical trial was carried out in order to evaluate both azithromycin and antimony efficacy in cutaneous leishmaniasis in Manaus, AM, Brazil. Forty nine patients from both genders, aged 14 to 70, with cutaneous ulcers for less than three months and a positive imprint for *Leishmania* spp. amastigotes were recruited into two groups. Group I (26 patients) received a daily-single oral dose of 500 mg of azithromycin for 20 days and Group II (23 patients) received a daily-single intramuscular dose of 20 mg/kg of meglumine antimony, also for 20 days. Azithromycin cured three of 24 (12.5%) patients on days 60, 90 and 120 respectively whereas therapeutic failure was considered in 21 of 24 (87.5%) cases. In group II, antimony cured eight of 19 (42.1%) cases as follows: three on day 30, one each on day 60 and day 90, and three on day 120. Therapeutic failure occurred in 11 of 19 (57.9%) individuals. The efficacy of antimony for leishmaniasis was better than azithromycin but analysis for the intention-to-treat response rate did not show statistical difference between them. Although azithromycin was better tolerated, it showed a very low efficacy to treat cutaneous leishmaniasis in Manaus.

KEYWORDS: Azithromycin; Antimony; Leishmaniasis; *Leishmania* spp.

INTRODUCTION

Leishmaniasis has been treated for decades with pentavalent antimony, pentamidine or amphotericin B but patients cure rate presents high variability depending on *Leishmania* spp., clinical picture and geographical location (SARAVIA et al, 1989; ROMERO et al., 2001).

Several studies have evaluated oral medicaments to treat cutaneous leishmaniasis caused by different *Leishmania* spp. without conclusive results (DOGRA & SAXENA, 1996; ALRAJHI et al., 2002). Recently, patients with cutaneous leishmaniasis caused by *L. (Viannia) panamensis* in Colombia received miltefosine and presented cure rate of 91%, whereas, in Guatemala, only 50% of cure was obtained in patients infected by *L. (Viannia) brasiiliensis* (SOTO et al., 2004). The results in patients with visceral leishmaniasis in India also showed good efficacy (93%) with this drug (SUNANDAR et al., 2002).

Azithromycin has been used to treat infections caused mainly by intracellular microorganisms because of its tissular distribution within phagocytes and its long half life (PETERS et al., 1992; WILDFEUER et al., 1996). This drug has shown activity against *Toxoplasma gondii*, *Plasmodium vivax* and *Cryptosporidium* spp., among others (GIACOMETTI et al., 2000; PUKKITTAYAKAMMEE et al., 2003).

In an experiment *in-vitro* and *in-vivo* model, azithromycin showed activity against *Leishmania major* (KROLEWIECKI et al., 2002). In an open trial in Brazil, it cured 16 out of 20 patients with cutaneous ulcers caused by *Leishmania (Viannia) braziliensis*, however, cure time, doses and cycles number presented high variability (PRATA et al., 2003). In the same location; three out of four old individuals with mucosal involvement presented clinical cure after several cycles with this drug (SILVA-VERGARA et al., 2004). Most recently, in an open study, none of 45 Syrian old-world leishmaniasis patients treated with azithromycin were cured (DAOUD & BOUSHI, 2006). The aim of this trial was to compare both azithromycin and antimony efficacy in patients with cutaneous leishmaniasis.

PATIENTS AND METHODS

This trial was carried out in an endemic leishmaniasis area where high active transmission occurs due to deep ecosystem changes introduced in the last decades to enlarge agriculture borders, settlements alongside the roads and non-planned urbanization replacing native jungle. The patients recruited were referred by other clinicians selected among several hundred of local residents that were diagnosed during 2003 at the Fundação de Medicina Tropical do Amazonas, a leishmaniasis state reference center.

Sample size was estimated in 202 cases with Yates correction considering 30% and 50% of cure to antimony and azithromycin.
respectively. Criteria for enrollment were as follows: individuals of both genders, aged 14 to 70, presenting cutaneous ulcers for less than 12 weeks and a positive imprint for *Leishmania* spp. amastigotes. Exclusion criteria: patients with history of treatment for leishmaniasis in the last three months, difficulty to attend follow-up, mucosal or diffuse involvement, fever, previous macrolide or antimony allergy, pregnancy, diabetes mellitus, AIDS, vasculitis, use of corticosteroids or immunosuppressive drugs, renal, hepatic or cardiac diseases and/or engaged in another clinical trial.

The consent to enrollment was obtained prior to allocation. The patients were clinically examined and number, diameter and size of lesions were assessed. A intradermal test using standard antigen was performed in all according to procedure already described (MAYRINK et al., 1979).

Patients were allocated in two groups according to individual choice of treatment. Group I received a daily-single oral dose of 500 mg of azithromycin for 20 days and Group II received a daily-single intramuscular dose of 20 mg/kg of meglumine antimony during the same period under supervision. Clinical follow-up was carried out every 30 days and all complaints, compliance and secondary effects were evaluated by spontaneous report and questionnaire.

Cure criteria were characterized by the whole epithelization of skin ulcers and the absence of erythema and/or inflammatory signs. Therapeutic failure was defined if lesions worsened on days 30, 60, 90 or 120 of follow-up despite an early good response.

Data analysis was performed using statistical for windows, 6.0 (Statsoft, Inc. USA). Shapiro-wilk test for normal distribution and Student’s t test for independent samples, Mann-Whitney for abnormal distributions and $\chi^2$ tests for comparing proportions. The project was approved by the ethical review board from Fundação de Medicina Tropical do Amazonas, Manaus, AM, Brazil.

**RESULTS**

From January to April 2003, forty nine patients with cutaneous leishmaniasis were recruited. Most of them were males under 35 years old. From these, 26 (53%) received azithromycin (Group I) and 23 (47%) received antimony (Group II). The main demographic and clinical characteristics of both groups were similar (Table 1).

Clinical cure was noticed in three of 24 (12.5%) cases of Group I on days 60, 90 and 120 respectively, whereas, 21 of 24 (87.5%) patients presented therapeutic failure as follows: on day 30 of treatment, 19 worsened and two had improved but worsened on day 60, then, they were all switched to antimony.

From Group II, eight of 19 (42.5%) cases were cured as follows: three on day 30, one each on day 60 and day 90, and three on day 120. The remaining 11 of 19 (57.9%) improved on day 30, but seven of them worsened on day 60 and four on day 90 and then received another 20-day cycle of antimony. Two patients of the azithromycin group and four of the antimony group did not attend follow-up (Table 2).

In general, azithromycin was well tolerated, however, diarrhea, abdominal pain, dizziness, headache and nausea were reported by 60%, 28%, 24%, 12% and 12% of them respectively but no patient interrupted the therapy because of these side effects. On the other hand, patients receiving antimony, often complained of arthralgia and myalgia and

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>Group I</th>
<th>Group II</th>
<th>$p$ value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age(years)</td>
<td>N</td>
<td>%</td>
<td>N</td>
</tr>
<tr>
<td>14-34</td>
<td>16</td>
<td>61.5</td>
<td>16</td>
</tr>
<tr>
<td>35-49</td>
<td>10</td>
<td>38.5</td>
<td>10</td>
</tr>
<tr>
<td>Gender</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>21</td>
<td>80.8</td>
<td>21</td>
</tr>
<tr>
<td>Female</td>
<td>5</td>
<td>19.2</td>
<td>5</td>
</tr>
<tr>
<td>Skin lesions(weeks)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt; 4</td>
<td>21</td>
<td>80.8</td>
<td>19</td>
</tr>
<tr>
<td>5-8</td>
<td>4</td>
<td>15.4</td>
<td>4</td>
</tr>
<tr>
<td>Multiple ulcers</td>
<td>17</td>
<td>65.4</td>
<td>13</td>
</tr>
<tr>
<td>Classical ulcer</td>
<td>18</td>
<td>69.3</td>
<td>12</td>
</tr>
<tr>
<td>Supraumbilical lesions</td>
<td>13</td>
<td>50.0</td>
<td>11</td>
</tr>
<tr>
<td>Ulcers with bacterial infection</td>
<td>15</td>
<td>57.7</td>
<td>7</td>
</tr>
<tr>
<td>Ulcer diameter &lt; 2 cm</td>
<td>18</td>
<td>69.0</td>
<td>17</td>
</tr>
<tr>
<td>Positive intradermal test</td>
<td>21</td>
<td>84.0</td>
<td>20</td>
</tr>
</tbody>
</table>

Clinical cure was noticed in three of 24 (12.5%) cases of Group I on days 60, 90 and 120 respectively, whereas, 21 of 24 (87.5%) patients presented therapeutic failure as follows: on day 30 of treatment, 19 worsened and two had improved but worsened on day 60, then, they were all switched to antimony.

From Group II, eight of 19 (42.5%) cases were cured as follows: three on day 30, one each on day 60 and day 90, and three on day 120. The remaining 11 of 19 (57.9%) improved on day 30, but seven of them worsened on day 60 and four on day 90 and then received another 20-day cycle of antimony. Two patients of the azithromycin group and four of the antimony group did not attend follow-up (Table 2).

In general, azithromycin was well tolerated, however, diarrhea, abdominal pain, dizziness, headache and nausea were reported by 60%, 28%, 24%, 12% and 12% of them respectively but no patient interrupted the therapy because of these side effects. On the other hand, patients receiving antimony, often complained of arthralgia and myalgia and

<table>
<thead>
<tr>
<th>Follow-up/day</th>
<th>0</th>
<th>30</th>
<th>60</th>
<th>90</th>
<th>120</th>
</tr>
</thead>
<tbody>
<tr>
<td>I</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>II</td>
<td>26</td>
<td>23</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Improving</td>
<td>-</td>
<td>-</td>
<td>5</td>
<td>16</td>
<td>2</td>
</tr>
<tr>
<td>Clinical cure</td>
<td>-</td>
<td>-</td>
<td>3</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>Worsening</td>
<td>-</td>
<td>-</td>
<td>19*</td>
<td>-</td>
<td>2*</td>
</tr>
<tr>
<td>Lost to follow-up</td>
<td>-</td>
<td>-</td>
<td>2</td>
<td>4</td>
<td>-</td>
</tr>
</tbody>
</table>

Group I: Azithromycin - Group II: Antimony; *: switched to Antimony; **: new treatment with Antimony.
only four withdrew it due to the severity of these symptoms but their lesions cured.

Overall, azithromycin cured three of 24 (12.5%) patients and antimony eight of 19 (42.5%) (χ² 5.22, p = 0.035, OR = 5.33 and IC 95% 0.98-32.3). The analysis for the intention-to-treat response rate did not show statistical difference between the two groups.

**DISCUSSION**

Due to limited evidence about azithromycin activity against *Leishmania* spp., this trial compared it with antimony in 49 patients with cutaneous leishmaniasis from an area where *L. (Viannia) guyanensis* causes 97% of infections (LAINSON & SHAW, 1987; GRIMALDI et al., 1989).

Azithromycin cured only three of 24 (12.5%) cases, similar to the results obtained in a pilot study in the same place (TEIXEIRA et al., 2007). In contrast, an open non-controlled study carried out in an endemic area where *L. (Viannia) braziliensis* prevails, azithromycin cured 16 of 20 (85%) patients with cutaneous ulcers although the doses, treatment and cure time were variable (PRATA et al., 2003).

Spontaneous cure in leishmaniasis often occurs after six months and it was unlikely to have occurred in the three patients cured as the epithelization of their lesions began with azithromycin administration and were wholly cured up to four months, which suggests some drug effect and not spontaneous healing only. Besides, one of the inclusion criteria was ulcer with less than 12 weeks.

It has been suggested that early treatment of patients with leishmaniasis can not represent an advantage at least during the first four weeks when compared with the results observed in patients with long term lesions. Most individuals in this trial presented ulcers with less than four weeks which could have contributed to the poor outcome observed (ROCHA et al., 1999).

The proportion of patients that presented ulcers with evidence of bacterial infection was significantly higher in the azithromycin group (57.7% vs 30.4%) but, no difference related to response was observed between them. That may be explained by the selection bias during allocation. Besides, a former report has shown that bacterial secondary infection did not influence the healing process in cutaneous leishmaniasis (VERA et al., 2001).

The prolonged half life (2-6 days) and high intra-phagocyte concentration of azithromycin together with good tolerance and oral administration, are conditions highly desirable in leishmaniasis treatment. Nevertheless, its action mechanism is still unknown and it might be related to leishmanicide direct effect and/or immunomodulatory action (XU et al., 1996; IANARO et al., 2000).

Several reports have shown rate cure of 60 to 80% using antimy in patients with cutaneous lesions caused by *L. (Viannia) braziliensis* in Latin America. In contrast, in Manaus only 42.5% of the patients were cured with this drug similar to other results already published (OLIVEIRA-NETO et al., 1997; ROMERO et al., 2001). These data support the high variability in cure rate of this disease depending on *Leishmania* spp., the clinical picture and the geographical region (SARAVIA et al., 1989; ROMERO et al., 2001; SOTO & TOLEDO, 2007). Besides, drug resistance that would be a naturally acquired condition in leishmania, has been partially associated with poor response (GROGL et al., 1992).

Due to an impressive decrease of leishmaniasis cases at the time, the study was closed prematurely with a small size with 49 patients, different from the estimated 202 and just before the difference became statistically significant between the treatments. This fact, together with operational difficulties related to the follow-up and species definition, hinder a better evaluation of the results and their correlation with clinical and epidemiological factors, but evidenced a very low efficacy of azithromycin to treat cutaneous leishmaniasis in Manaus.

**RESUMO**

Falha da azitromicina e do antimonial no tratamento da leishmaniose cutânea em Manaus, AM, Brasil

Com o objetivo de avaliar a eficácia da azitromicina no tratamento da leishmaniose cutânea, foi realizado ensaio comparativo, em Manaus. Foram recrutados 49 pacientes de ambos os sexos, com idades entre 14 e 70 anos que apresentassem úlceras cutâneas com menos de três meses de evolução e que tivessem exame direto positivo para amastigotas de leishmanía. Estes pacientes foram alocados em dois grupos assim: Grupo I (26) recebeu uma dose diária de 500 mg de azitromicina pela via oral durante 20 dias e o Grupo II, recebeu uma dose diária de 20 mg/kg de antimonio de meglumina por via intramuscular, durante 20 dias. Do grupo da azitromicina, três (12,5%) de 24 pacientes curaram 60, 90 e 120 dias, respectivamente, enquanto, em 21 (87,5%) de 24 houve falha terapêutica. No grupo do antimonial, oito (42,5%) de 19 pacientes curaram como segue: três no dia 30, um no dia 60, um no dia 90 e três no dia 120. Contudo, em 11 (57,9%) de 19 casos, houve falha terapêutica. A azitromicina foi menos eficaz do que o antimonial, embora, a análise da taxa de resposta por intenção de tratamento não mostrou diferença significativa, entre eles. A azitromicina foi melhor tolerada; porém, mostrou-se pouco eficaz no tratamento da leishmaniose cutânea, em Manaus.

**ACKNOWLEDGEMENTS**

The study was supported by CNPq and FUNASA. We are grateful to the kind assistance of the staff at Fundação de Medicina Tropical do Amazonas, Manaus, AM, and to Angela Azor and Maria Rita de Souza for technical assistance.

**REFERENCES**


Received: 26 September 2007
Accepted: 2 April 2008