Heat therapy for cutaneous leishmaniasis elicits a systemic cytokine response similar to that of antimonial (Glucantime) therapy

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Summary
Controlled heat delivered as radio waves has been used successfully in the treatment of cutaneous leishmaniasis (CL). Here we investigated whether local heat therapy has systemic effects, as measured by the modulation of cytokine production following heat therapy of CL lesions compared with antimonial (Glucantime) treatment. Patients with CL were randomly assigned into this study. Heat (50°C for 30s) was applied once. The control group received Glucantime therapy for 20 d. Cytokine production by peripheral blood mononuclear cells was assayed on days 0, 14 and 28 after onset of treatment. At the end of 28 d, 75% of lesions were healing or healed in the heat therapy group and 90% in the control group (P = 0.1261). There was a decrease in IFN-γ, IL-5 and TNF-α levels comparing day 0 with day 28 in both groups, but no difference between the two therapy groups. In patients with only one of several lesions treated with heat therapy, the untreated lesions also healed. Local heat therapy in CL lesions leads to systemic cytokine responses similar to that induced by systemic Glucantime therapy.

1. Introduction
Heat therapy for cutaneous leishmaniasis (CL) has been reported for half a century. da Silveira and Brenner (1950) described the healing of a patient with CL and one with mucocutaneous leishmaniasis (MCL) 1 month after heat induced by Haemophilus ducreyi injections. In-vitro studies showed the sensitivity of various New and Old World
Leishmania species to heat (Berman and Neva, 1981; Sacks et al., 1983). Subsequently, Neva et al. (1984) showed that diffuse cutaneous leishmaniasis (DCL), but not CL, responded to heated water treatment at 39–41 °C. Junaid (1986) treated CL with infrared at 55 °C for 5 min and also reported the healing of non-treated lesions in the treated patients. Aram and Leibovici (1987) observed healing of CL with ultrasound-induced heat. Navin et al. (1990) carried out a control trial using the original Thermosurgery® instrument in Guatemala. Twenty-two patients received antimonial; 22 received 3-weekly heat treatments of 50 °C for 30 s; and 22 served as placebo controls. Three-quarters of the patients in the study had Leishmania (Viannia) braziliensis infection. Healing was similar in the two therapy groups, which was faster than in the placebo group. Velasco-Castrejon et al. (1997) carried out a feasibility study using the same instrument in Mexico on 201 patients with CL. A single application of 50 °C for 30 s across the anesthetized lesion was used, and resulted in over 90% healing.

CL caused by L. (V.) braziliensis can metastasize, leading to mucocutaneous leishmaniasis (MCL) (Mansden, 1986). Patients with CL and MCL have an inflammatory reaction in the lesions and a strong delayed-type hypersensitivity (DTH) response associated with IFN-γ and TNF-α production (Ribeiro-de-Jesus et al., 1998). By contrast, DCL patients lack the inflammatory response, have no DTH response and diminished Th1 cytokines (Bonifim et al., 1996; Melby et al., 1989). The purpose of this study was to determine whether healing of CL caused by local heat therapy induced by radio waves was associated with a modulation of systemic cytokine responses compared with Glucantime therapy, the standard treatment for CL in Brazil.

2. Materials and methods

2.1. Patients

Between March 1997 and December 2000, the same physician in a health care clinic in Laje attended patients from Laje and neighborhood towns in the Jiquirica Valley, in the southwest of Bahia State, Brazil. At the first visit, the physician determined the clinical status of the patients. When they met the inclusion category described below, tests were done to confirm the diagnosis of CL. A Montenegro skin test was applied on the volar surface of the forearm and the cutaneous reaction measured 48 h later; a blood sample (3—5 ml) was drawn for leishmanial serology and a biopsy of the lesion was done and the specimen was divided, with part put in 10% solution of formaldehyde and part frozen in liquid nitrogen. A 4-mm punch pointed toward the ulcer border was done and a photograph of the lesion was taken. A venous blood sample (20 ml) was collected to determine cytokine responses. Biopsy using a Montenegro skin test and positive serology were randomly assigned to the heat therapy or Glucantime treatment groups. The diagnosis was confirmed by biopsy of the lesion. Physical examination of the patient included visualization of the nasal and oropharyngeal mucosa and checking for lymphadenopathy. The skin lesions were described in detail, including their number (one or two), location and size (expressed in mm, great axis × small axis/2); the duration and clinical evidence of bacterial superinfection was noted, and a photograph of the lesion was taken. A venous blood sample (20 ml) was collected to determine cytokine responses. Biopsy using a 4-mm punch pointed toward the ulcer border was done and the specimen was divided, with part put in 10% solution of formaldehyde and part frozen in liquid nitrogen.

2.2. Design of the study

The study protocol was designed to compare the success of the treatment at the end point of 28 d, as any systemic effects should have appeared during this time. Afterward, all patients in the heat therapy group received standard 20-d Glucantime treatment independently of how well they had responded. The potential risk of future MCL in patients submitted only to heat therapy precluded the proper evaluation of the individual response to heat therapy alone, until we could determine whether this therapy had a systemic effect in addition to the local one. The design of the study took into consideration that it is unethical to treat a group having MCL with L. (V.) braziliensis with a placebo.

At day 0, patients included in the study with a presumptive diagnosis of CL (as determined by a positive skin test and positive serology) were randomly assigned to the heat therapy or Glucantime treatment groups. The diagnosis was confirmed by biopsy of the lesion. Physical examination of the patient included visualization of the nasal and oropharyngeal mucosa and checking for lymphadenopathy. The skin lesions were described in detail, including their number (one or two), location and size (expressed in mm, great axis × small axis/2); the duration and clinical evidence of bacterial superinfection was noted, and a photograph of the lesion was taken. A venous blood sample (20 ml) was collected to determine cytokine responses. Biopsy using a 4-mm punch pointed toward the ulcer border was done and the specimen was divided, with part put in 10% solution of formaldehyde and part frozen in liquid nitrogen.

2.2.1. Heat therapy

This was given in a single session. The lesion was washed with saline, then iodine and anesthetized with 2% lidocaine. The fork-like applicator of the Thermosurgery® instrument, powered by batteries, was placed at the edge of the lesion pointing toward the center and heat at 50 °C was applied for 30 s, then the applicator was moved to an adjacent area until the lesion had been completely covered, taking 4–5 min. The heat is completely localized and produced between the two electrodes of the applicator (an area approxi-mately 3 × 4 mm). The lesion was then covered with a gauze bandage. No additional treatment was administered.

2.2.2. Antimony treatment

Intravenous injections of Glucantime, 20 mg/kg/d, were given during 20 consecutive days for all patients of the antimony group starting at day 0 and to all patients in the heat therapy group after day 28.

On day 14, the lesion was measured and signs of healing or secondary infection noted, and at that time another biopsy...
was taken for immunohistochemistry. The lesions were re-evaluated on day 28. Blood samples were taken for cytokine analysis on days 14 and 28.

2.2.3. Montenegro skin test
At the first visit, a Montenegro skin test was applied to the volar face of the forearm of all patients. Test readings were made 48 h after application, and the size (mm) of reaction was measured. These were considered positive when the diameter was 5 mm in diameter or larger. The antigenic material used was a soluble extract antigen from L. (V.) braziliensis killed promastigotes as previously described (da Costa et al., 1996).

2.2.4. Serology
Patients’ sera were tested for the presence of anti-Leishmania antibodies by ELISA using a crude parasite antigen following the method previously described (Ashford et al., 1993).

2.2.5. Determination of cytokines
Peripheral blood mononuclear cells (PBMC) isolated from a ficoll-hypaque gradient (Pharmacia; Uppsala, Sweden) of heparinized venous blood PBMC were cultured in 24-well plates 5 × 10^6 cells/well in RPMI medium (Life Technologies, Gibco-BRL, Gaithersburg, MD, USA) supplemented with 10% fetal calf serum (Hyclone, Logan, UT, USA), L-glutamine (2 mmol/l) sodium pyruvate (1 mmol/l), HEPES (10 mmol/l) and gentamycin (50 μg/ml) (Sigma, St Louis, MO, USA). Cultures were stimulated with L. (V.) braziliensis antigen prepared by freezing and thawing of axenic promastigotes three times. Cell-free supernatants were collected after 72 h for culture of cytokine analysis. Sandwich ELISA was carried out using antibody pairs from PharMingem (San Diego, CA, USA) and recombinant cytokines from R&D (Minneapolis, MN, USA), following the manufacturers’ instructions. Reaction was developed using 3,3',5,5'-tetramethylbenzidine (TMB peroxidase substrate; Kinkergaard & Perry Laboratories, Gaithersburg, MD, USA) and read at 450 nm.

2.2.6. Immunohistochemistry for parasite detection
All biopsy fragments from skin lesions were fixed by tamponade 10% formalin, for 1—3 d. After dehydration and paraffin embedding, serial sections of 3 μm thickness were made, and immunohistochemistry was used to investigate the presence of amastigote forms of Leishmania in the lesions before and after treatment. A rabbit polyclonal antibody that recognizes different Leishmania species was used to identify amastigote forms. Although this antibody did not identify particular Leishmania species, it is known from previous studies in the region of the Jiquiricá river valley in Bahia that 95% of CL in this area is caused by L. (V.) braziliensis (Jones et al., 1987). Sections were incubated with a biotinylated anti-rabbit IgG followed by streptavidin-peroxidase conjugate, and the reaction was developed using diaminobenzidine (DAB). Parasitic diagnosis from the biopsies was confirmed in 27 of the patients (12 [71%] of the heat therapy group and 15 [79%] of the Glucantime group). Three patients had histopathology suggesting CL, along with positive skin test and/or serology. A number of positive biopsies from 22 patients (14 in the Glucantime group and eight in the heat therapy group) were studied to compare the parasite load before and after treatment. The number of amastigotes seen in situ, before (in biopsy tissue by day 0) and after (in biopsy tissue by day 14) treatment, was determined according to intensity grades: (1) 1 to 10 amastigotes was considered as parasitism of mild intensity; (2) between 10 and 100 amastigotes was considered as parasitism of moderate intensity; and (3) above 100 amastigotes was considered as intense parasitism.

2.3. Statistical analysis
Data analysis was done using Epilinfo 6.04 (CDC, Atlanta, GA, USA) and SPSS (7.0 student version; SPSS Inc., Chicago, IL, USA) statistical software. Nonparametric statistical tests were used to allow for the lack of normality of the distribution of some variables considered in the study. Mann–Whitney and Kruskall–Wallis statistical analysis was used to test for differences between the two groups of treatment; the Wilcoxon signed-rank test was used to verify differences between cytokine levels before and after treatment. A χ² test was applied to categorical variables. Graphics were constructed using Excel 97 (Microsoft, Redmond, WA, USA) and SPSS software.

3. Results
3.1. Demographic and clinical baseline characteristics of patients
The comparison of the demographic and clinical baseline characteristics of the two groups of patients is presented in Table 1. There were no statistical differences between the characteristics of the two groups, except that there were more patients with two lesions in the Glucantime group. Thus, randomization successfully allocated patients with similar characteristics into the two treatment groups. Further, no relation was demonstrated between Montenegro skin test, duration of active lesions and size of ulcers with Leishmania serology when tested by Kruskall–Wallis statistical analysis. Parasitic diagnosis from the biopsies was confirmed in 27 of the patients, 12 (71%) of the heat therapy group and 15 (79%) in the Glucantime group; the remainder had histopathology suggesting CL, along with positive skin test and/or serology.

3.2. Overall clinical response
By day 28, most of the 37 patients were in the process of healing or had completely cured their lesions. One patient, however, who was treated with heat therapy, was excluded from analysis because he had put gunpowder over his ulcer and burned it several days after day 14 evaluation. Process of healing, as determined by decrease in size of the lesion and over 50% re-epithelization of the ulcer, was observed in 75% (12/16) of patients in the heat therapy group and 98% (18/20) in the Glucantime group. Three patients had complete cicatrization of the lesion: one in the heat therapy group and two in the Glucantime group. Three patients in the heat therapy group had no healing response seen by
Table 1  Comparative baseline characteristics of 37 patients treated with heat therapy or Glucantime

<table>
<thead>
<tr>
<th>Patient characteristics</th>
<th>Heat therapy (n=17)</th>
<th>Glucantime (n=20)</th>
<th>Statistical tests</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age</td>
<td>Years (range)</td>
<td></td>
<td></td>
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<tr>
<td></td>
<td>34 ± 14.6 (18–65)</td>
<td>16 ± 17.2 (18–67)</td>
<td><em>P</em>= 0.7078 (ANOVA)</td>
</tr>
<tr>
<td>Gender</td>
<td>Male, n (%)</td>
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<tr>
<td></td>
<td>9 (53)</td>
<td>15 (75)</td>
<td><em>P</em>= 0.2913 *ν² = 1.11 (Yates’ corrected)</td>
</tr>
<tr>
<td>Skin color</td>
<td>Female, n (%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>8 (47)</td>
<td>5 (25)</td>
<td><em>P</em>= 0.9633 *ν² = 0.00 (Yates’ corrected)</td>
</tr>
<tr>
<td>Total no. lesions</td>
<td>Brown/black, n (%)</td>
<td></td>
<td></td>
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<tr>
<td></td>
<td>11 (65)</td>
<td>12 (60)</td>
<td><em>P</em>= 0.0481 *ν² = 3.04 (Yates’ corrected)</td>
</tr>
<tr>
<td></td>
<td>One, n (%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>16 (94)</td>
<td>13 (65)</td>
<td></td>
</tr>
<tr>
<td>Ulcer location</td>
<td>Two, n (%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>1 (6)</td>
<td>7 (35)</td>
<td><em>P</em>= 0.4276 *ν² = 2.78</td>
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<td></td>
<td>Face, n (%)</td>
<td></td>
<td></td>
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<tr>
<td></td>
<td>0</td>
<td>3 (15)</td>
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<tr>
<td></td>
<td>Throat, n (%)</td>
<td></td>
<td></td>
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<tr>
<td></td>
<td>2 (12)</td>
<td>2 (10)</td>
<td></td>
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<tr>
<td></td>
<td>Upper limbs, n (%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>3 (18)</td>
<td>3 (15)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Lower limbs, n (%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>12 (70)</td>
<td>12 (60)</td>
<td><em>P</em>= 0.784 (ANOVA)</td>
</tr>
<tr>
<td>Age of lesion (days)</td>
<td>Median (range)</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>45 (15–180)</td>
<td>50 (20–240)</td>
<td><em>P</em>= 0.7799 (Mann–Whitney)</td>
</tr>
<tr>
<td>Secondary infection *a</td>
<td>Present, n (%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>4 (24)</td>
<td>9 (45)</td>
<td><em>P</em>= 0.3087 *ν² = 1.04 (Yates’ corrected)</td>
</tr>
<tr>
<td>Satellite ganglia *a</td>
<td>Absent, n (%)</td>
<td></td>
<td></td>
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<tr>
<td></td>
<td>10 (58)</td>
<td>15 (75)</td>
<td><em>P</em>= 0.4869 *ν² = 0.48 (Yates’ corrected)</td>
</tr>
<tr>
<td>Montenegro test</td>
<td>Absent, n (%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>4 (24)</td>
<td>9 (45)</td>
<td></td>
</tr>
<tr>
<td>Leishmanial serology *a</td>
<td>Present, n (%)</td>
<td></td>
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</tr>
<tr>
<td></td>
<td>14 (82)</td>
<td>32 (86)</td>
<td><em>P</em>= 0.3644 *ν² = 7.65</td>
</tr>
<tr>
<td>Amastigotes in biopsy *a</td>
<td>Present, n (%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>12 (71)</td>
<td>15 (79)</td>
<td><em>P</em>= 0.7060 (Fisher’s exact test: 2-tailed)</td>
</tr>
<tr>
<td>Montenegro test</td>
<td>Absent, n (%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>5 (29)</td>
<td>4 (21)</td>
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</table>

* Day 0.

Day 28 evaluation. There was no significant statistical difference on the overall rate of response between the two groups (χ² = 4.14, *P*= 0.1261). The size of ulcers was tested for association with the healing process by the two treatments, using χ². There was no statistical evidence that size of ulcers influenced cicatrization.

No significant adverse effect was seen or reported by patients who submitted to heat therapy, except for secondary bacterial infection after treatment. A total of eight patients presented secondary bacterial infection, seven of them after heat therapy and one after Glucantime treatment (Yates’ corrected χ² = 3.86, *P*= 0.0351, Fisher’s exact test).

Of special interest, only one lesion of the patient with two lesions in the heat therapy group was heat-treated. Nevertheless, the untreated lesion on the opposite leg healed as rapidly as the treated lesion (Figure 1).

3.3. Parasite burden determination in the lesions

Comparative evolution of parasite burden between day 0 and day 14 in the treatment groups was analyzed using the Wilcoxon signed-rank test. In the Glucantime group, 64% of previously positive biopsies for Leishmania became negative on day 14 (*P*= 0.05). In the heat therapy group, 25% of previously positive biopsies became negative on day 14 (*P*= 0.131), and a reduction in intensity of parasitism was seen in two patients of this group. Both groups had similar intensity of parasitism in day 0 biopsies, as analyzed by Kruskal–Wallis test (*P*= 0.8947).

3.4. Systemic cytokine response

The results of cytokine production by PBMC (TNF-α, IFN-γ, IL-5 and IL-10) were analyzed. A decrease in the levels of TNF-α, IFN-γ and IL-5 comparing day 0 with day 28 was found in both groups: in the heat therapy group (*P*= 0.016, *P*= 0.025 and *P*= 0.013, respectively) and in the Glucantime group (*P*= 0.091, *P*= 0.046 and *P*= 0.075, respectively) using the Wilcoxon signed-rank test (Figure 2). No statistically significant difference was found in IL-10 levels in either treatment group when day 0 was compared to day 28. When comparing the cytokine levels of the two treatment groups, no statistically significant differences were found, as summarized in Table 2 (the lack of difference between the decreased IFN-γ levels of both groups was due to the variations in each group).
4. Discussion

The results of this study show that a single local application of heat therapy to CL lesions (radio waves induced by a ThermoSurgery instrument) will alter the production of systemic cytokines by PBMC similar to that seen by the systemic injection of Glucantime for 20 d. Whether the alteration of the cytokine response is a direct result of the heat therapy and Glucantime, or a result of the healing process stimulated by the treatments, is not known at present. Further evidence of the systemic effect of heat therapy, however, is given below. In addition, this study confirms that heat therapy for CL has beneficial effects similar to those of Glucantime in this limited 28-day evaluation, confirming previous studies (Navin et al., 1990; Velasco-Castrejon et al., 1997).

The finding that *Leishmania* species, especially those causing CL, are temperature-sensitive organisms (Berman and Neva, 1981; Sacks et al., 1983) has led to the use of heat for the treatment of cutaneous forms of leishmaniasis. Heat has been produced by the injection of bacteria (da Silveira and Brener, 1950) or applied directly to lesions using hot water (Neva et al., 1984), infrared (Junaid, 1986), ultrasound (Aram and Leibovici, 1987) and radio waves (Navin et al., 1990; Velasco-Castrejon et al., 1997). The delivery and control of the temperature, however, was previously
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Figure 2  Cytokine response of peripheral blood mononuclear cells (PBMC) of patients treated with Glucantime or heat therapy. Levels of IFN-γ, TNF-α, IL-5 and IL-10 in supernatants of PBMC stimulated with leishmanial antigen before (day 0) and 28 d after heat therapy or Glucantime treatment.

This was mostly solved by the first Thermo-Surgery instruments (Navin et al., 1990; Velasco-Castrejon et al., 1997), and especially in the latest model used in this study, which has an improved temperature control and sensor (±0.2 °C).

The inflammatory response and control of infection in CL are mainly Th1-mediated. There is a polarity between the extreme inflammation of MCL with high IFN-γ and TNF-α, the moderate inflammation in CL, with less elevation of these cytokines and the absence of inflammation of DCL.
with low or lacking IFN-γ (Bacellar et al., 2002; Bomfim et al., 1996; Castes et al., 1988; Follador et al., 2002). In our study, the production of IFN-γ and TNF-α decreased following the therapies. Previous reports on the production of IFN-γ and TNF-α in response to antimonial therapy has been variable, going down (Ribeiro-de-Jesus et al., 1998), not significantly altered (Coutinho et al., 1998) or raised, although lowered when chemotherapy was combined with immunotherapy (Toledo et al., 2001). In our study, the Th2 associated cytokines were either diminished (IL-5) or variable (IL-10) after 28 d of therapy. IL-10 is known to partially regulate Th1 responses, diminishing IFN-γ production. Because IL-10 levels are low in MCL patients, in which IFN-γ is the highest (Bacellar et al., 2002), the maintenance of IL-10 after treatment of CL may be important in the prevention of MCL.

In addition to the controlled trial described above, seven CL patients resistant to Glucantime therapy were treated with heat. These patients were not healing after being injected with as many as 80 to 100 or more ampuoles of Glucantime. In every case, a dramatic healing response occurred to the heat therapy. Further, several CL patients who had to be excluded from the controlled trial because Glucantime was contraindicated (two pregnant women and two patients having no contact with the leishmanial antigen released and incoming inflammatory cells at the burn site induce an immune response similar to a vaccine. Alternatively, the burn itself may cause the release of inflammatory molecules, which lead to the distal healing. A controlled study, designed to confirm that not all lesions on a patient need to be treated should be carried out, especially with patients having multiple CL lesions.

For the last 60 years, pentavalent antimony has been used as the drug of choice for the treatment of leishmaniasis (Mardens, 1985). It is well known that these compounds are toxic and that in some circumstances, as in localized single cutaneous lesions, the treatment can be more harmful than the disease (Convid et al., 1987). Many physicians no longer use pentavalent antimony because of its toxicity. Several alternatives have been tried, but so far none is considered convenient or efficacious enough to replace antimonial therapy. Heat therapy would appear to be a good alternative in CL, as it is effective and usually requires only one application. In addition, the Thermosurgery instrument can be taken into the field (it is a small equipment with an internal rechargeable battery). We now put antibiotic ointment on the lesion after the heat therapy to reduce the risk of bacterial infection.

In our trial, four patients had not responded to heat therapy at day 28. We were unable to prolong the evaluation of heat therapy because everyone in that group received Glucantime after 28 d as part of the study design approved by the ethical committee. Now that a systemic effect of local heat therapy has been demonstrated, a large controlled trial involving heat therapy and Glucantime, with a 10-year follow-up, has been initiated by others in Ceará, Brazil to determine whether there is a difference in the incidence of MCL.

Conflicts of interest statement
The authors have no conflicts of interest concerning the work reported in this paper.

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