INTRODUCTION

Chromoblastomycosis is a chronic disease caused by dematiaceous fungi found in soil and decaying vegetation, common in tropical and subtropical regions. Infection occurs by traumatic fungal inoculation into the skin, and mycosis is very prevalent in rural men. It is characterized by lesions of polymorphic, papillomatous, nodular, cauliflower, verruciform, hyperkeratotic, cicatricial, plaque or combinations, in the lower limbs, with no tendency to spontaneous cure (Figure 1). Chronic inflammatory infiltrate, microabscesses, dark-brown parasites and granulomatous process are frequent findings in histopathology. Suppurative granuloma rich in fungal cells is almost always observed in verrucous lesions, while tuberculous granulomas with few parasites are mainly detected in cicatricial lesions and well-defined erythematous plaques, suggesting that verrucous lesions have a less competent tecidual inflammatory response. The muriform element is pathognomonic and observed in tissue biopsy specimens or by direct mycological examination from black dotted scrapings of the cutaneous lesion with 10% KOH.

Little is known about histological alterations and variation of the positivity of mycological exams resulting from systemic treatments with antifungals. This study aims to analyze tissue changes during therapy with itraconazole.

METHODS

This is a histological study of cases of chromoblastomycosis from clinical research performed at a university hospital in the Southern region of Brazil. Patients were submitted to at least four biopsies of cutaneous lesions - one biopsy to diagnose the disease and a minimum of three biopsies to follow the tissue modifications during the treatment. Periodic evaluation occurred every four months, and participants were followed for three years from 1988 to 1991. The project to access this historical documentation was approved by the Ethics Committee (protocol 10835/2012).

Skin biopsies, always conducted by the same researcher, consisted of an ellipse of 1cm of tissue on the largest axis, composed of epidermis, dermis and hypodermis, and were intended for mycological and histopathological study. In active lesions, biopsies were performed to spontaneous decrease or disappearance, fibrosis increase and quantitative/morphological changes occurred during treatment.

RESULTS: Fontana-Masson stain was superior to hematoxylin-eosin stain in fungal count in the epidermis (mean difference=0.14; p<0.05). The most distinct mycosis tissue responses were registered in the dermis. Epidermal thinning, granulomatous infiltrate decrease or disappearance, fibrosis increase and quantitative/morphological changes occurred during treatment. Patients could not be located to have their current skin condition examined.

CONCLUSION: Parasitic and tissue changes verified in this study can reflect the parasite-host dynamics under itraconazole action.

Keywords: Chromoblastomycosis; Histology; Itraconazole; Mycology; Pathology
collected in regions of black dotted and/or erythematous borders. In healed lesions during treatment, biopsy was obtained from its border, and desquamative areas with granular appearance were selected. Hematoxylin-eosin (HE) staining was used to verify tissue changes, and HE and Fontana-Masson (FM) staining was used to evaluate fungi throughout the treatment (Chart 1). For the direct mycological examination, 10% KOH solution was used and, for culture, Sabouraud agar medium.

Fungi count occurred on two slides, one stained by HE and the other stained by FM. Counting started always in the upper left corner of the slides, moving the chariot from top to bottom in the epidermis-hypodermic direction until the end of the histological section. Then the chariot was moved from left to right relative to the observer, moving a field at right angle and resuming the count from the bottom up in the hypodermis-epidermis direction. American Optical Spencer microscope was used, with a 40x objective lens and a 10x ocular lens. In the slides with more than one histological section, stained by HE, the section closest to the slide label, to the left of the observer, was used. A reticle was adapted to the 10WF CAT 176 AO ocular lens to delineate areas of histological sections for counting.

In each slide, 100 reticulated areas were counted, registering only the spherical integral parasitic elements found in the odd-numbered reticulated areas. In order to avoid registering a parasite more than once, the following rule was adopted: the parasite that was located on the border lines of the reticulated area, left and above, was counted for this field. The parasite that was situated on the right and below did not belong to this field. For record purposes, each parasitic element found with or without septation, isolated or clustered, was equivalent to one unit. Artifact pigments, fragmented structures and hyphae were excluded from counting.

Criteria used to control the treatment were: clinical (complete healing of all lesions), mycological (absence of parasites at the direct mycological examination and culture) and histological (absence of parasites, absence of microabscesses, atrophy of the epidermis, replacement of the active granulomatous infiltrate in the dermis due to chronic inflammatory infiltrate and dense fibrosis).

In the statistical analysis, Kappa coefficients were estimated and 95% confidence intervals were used to evaluate the concordance between mycological, histological and culture examinations. For the comparison of the HE and FM stainings, Student’s t-test was used for paired samples. Comparison between tumor manifestation and plaque manifestation, in relation to the probability of having a clinical improvement or cure as final result, was performed using Fisher’s exact test. For this same analysis, the relative risk was estimated. Values of p<0.05 indicated statistical significance. Data were analyzed using IBM SPSS Statistics v.20 software.

RESULTS

The sample consisted of 20 individuals, white, adults, farmers, 17 of them were men, with mean age of 58.2 years and standard...
deviation of 11.1 years (42 to 82 years) and 3 women aged 33 to 54 years (mean age of 41.6 and standard deviation of 11.0 years), all receiving treatment with itraconazole 200-400 mg/day. During the study, 135 skin biopsies were performed with a mean of 6.7 biopsies/patient.

Prior to treatment, direct mycological examination was positive in 80% of cases; culture in 85%; and histological examination, in 100% (Figure 2). Culture and microculture allowed the identification of F. pedrosoi in 100% of the patients (Figure 3). During follow-up, there was a good level of concordance between mycological (direct and culture) and histopathological examination (Kappa: 0.64; 95% CI: 0.51 to 0.76); between culture and direct mycological examination (Kappa: 0.67, 95% CI 0.55 to 0.80); and between culture and histopathological examination (Kappa: 0.71, 95% CI: 0.59 to 0.84).

Tissue modifications of the skin during treatment with itraconazole were significant in the epidermis, dermis and hypodermis when the results of the baseline evaluation (month 0) were compared with the results observed in the period from 9 to 12 months after treatment. In the comparison between the baseline and the 4 to 8 months period, the results indicated significant differences for the epidermis and dermis, with a tendency towards significance for fibrosis. Between the period of 4 to 8 months and the period of 9 to 12 months, a significant difference was found for the intensity of fibrosis (Table 1).

Figure 4 illustrates histological aspect prior to treatment. Figure 5 shows parasitic elements in the middle of the dermal inflammatory infiltrate.

Comparing the visualization of the fungi in the FM and HE stainings, it was verified that their identification was facilitated by the FM staining, mainly in the epidermis (mean difference of 0.14 fungus and standard deviation of 0.50; p<0.05) (Figure 6).

In the final evaluation, of the 11 patients with tumor manifestation, 9 (82%) presented improvement or clinical cure and 2 (18%) presented clinical and biological cure. Of the 9 patients with plaque manifestations, 3 (33%) presented improvement or clinical cure and the other 6 (67%) presented clinical and biological cure (p = 0.065). Therefore, it is estimated that, for cases with tumor manifestation, the relative risk of having clinical improvement or cure (and not clinical and biological cure) is 2.5 times, when compared with cases with plaque manifestation.

DISCUSSION

In this study, before the institution of the treatment, the direct mycological examination was positive in 80% of the cases; culture in 85%; and histological examination in 100%. As the mycosis presents centrifugal growth, choosing areas of black dotted region and active lesions borders or, in the absence of these borders, of cicatrizated lesions guided the investigation of fungi, which allowed equivalence between the positivity of the mycological and histological exams. Culture and microculture allowed the identification of F. pedrosoi as an etiological agent of all cases, according to the literature.8,13-16

In chromoblastomycosis, the tissue response to the fungus is non-specific and may be similar to other deep mycoses. Most epidermal tissue alterations on histopathological examination are hyperkeratosis and pseudoepitheliomatous hyperplasia. In der-}

mal and hypodermal tissues, non-specific diffuse inflammatory infiltrates are detected.17 Other findings include suppurative and granulomatous reactions, sclerotic bodies and acanthosis with trans-epidermal fungal elimination.8,18-22 In this study, the intensity and type of histological response were modified according to patient and therapeutic phase. Before treatment, there was a tendency to an exuberant inflammatory condition, epidermis with hyperkeratosis, parakeratosis, leukocyte crusts, Malpighian layer acanthosis, spongiosis, neutrophil and eosinophil exocytosis and microabscesses. Acanthosis was irregular, with typical cells and hypertrophy of the papillae protruding above the cutaneous surface, with enlargement and bifurcation of the interpapillary ridges, constituting pseudoepitheliomatous hyperplasia.

![Figure 2: Positive direct mycological examination: brown spherical parasite elements, with and without septation, observed on tissue fragment with KOH (x1000)](image)

![Figure 3: A) Colony morphology of Fonsecaea pedrosoi in Sabouraud-dextrose-agar medium after 21 days of incubation; B) Microculture of Fonsecaea pedrosoi (X1000)](image)
The most distinct alterations in this study were found in the dermis. Before treatment, hyperplasia, papillomatosis, vascular neoformation and dilations, and edema were noted. The cellular infiltrate combined suppurative and granulomatous foci, being the parasites found amidst the inflammatory response or inside giant cells. The stratum corneum microabscesses probably corresponded to the black dots, observed macroscopically in the skin surface, resulting from transepithelial elimination and from the presence of dermoepidermal fistulas draining outwards. Throughout the treatment, there was progressive atrophy of the epidermis, decrease or disappearance of the granulomatous infiltrate and increase in the degree of fibrosis. Findings such as disappearance of microabscesses and granulomas, disappearance of parasites to mycological exams (direct and culture) and clinical healing of all the lesions allowed the support of the cure criteria, being able to be used to interrupt the treatment and follow up of the cases.23

| Evaluation phase (month) | Epidermis morphology (E1+E2) | Dermal inflammatory infiltrate (D1+D2) | Fibrosis intensity (F1) |
|--------------------------|-----------------------------|----------------------------------------|------------------------|
| 0                        | 5 (25%)                     | 15 (75%)                               | 20 (100%)              |
| 4 to 8                   | 14 (70%)                    | 6 (30%)                                | 15 (75%)               |
| 9 to 12                  | 18 (90%)                    | 2 (10%)                                | 8 (40%)                |

| Compared phases          | P value                     | P value | P value |
|--------------------------|-----------------------------|---------|---------|
| 0 x 4 to 8               | 0.004                       | 0.004   | 0.062   |
| 0 x 9 to 12              | <0.001                      | <0.001  | <0.001  |
| 4 to 8 x 9 to 12         | 0.125                       | 0.250   | 0.016   |

The most distinct alterations in this study were found in the dermis. Before treatment, hyperkeratosis, pseudoepitheliomatous hyperplasia and dermal inflammatory infiltrate (Hematoxylin & eosin, X40)
FM staining was superior to HE staining in detecting fungi with a lighter shade, mainly in the epidermis. Due to their natural pigmentation, these parasites are easily identified on uncolored slides. Special stainings like FM are useful for demonstrating hyphae in keratinized layers. In general, fungi detected in the histological sections had the following aspects: demaceous hyphae and septated spherical parasitic elements in one plane, in two planes (muriform element) or without septation, representing the metamorphosis of the parasite, not being necessary the exclusiveness of the muriform element for the diagnosis of the disease.

In this study, the finding of hyphae in the epidermis was occasional. Parasites presented predominantly dermal location, followed by epidermal, which could be explained by changes in the thickness of their cell wall and decrease of the melanin pigmentation. The location on the surface allows transformation to mycelial form. Fungi located in the dermis undergo the phenomenon of parasitic reduction and generally have a thicker and pigmented wall due to adverse conditions to their parasitism: tissue pressure, local acidification and suppurative and granulomatous inflammatory reaction. Although curious, data on parasite count would lack better investigation, since the amount of fungi recorded was neither static nor definitive, since the histological section represented only one plane of a process that is essentially evolutionary and three-dimensional.

There was a greater tendency for clinical improvement or cure in tumoral forms when compared with plaque manifestation. Favorable clinical evolution during treatment was correlated to negative mycological exams, slower growth of cultures, or non-development of the fungus, and a predominance of histopathology with isolated parasites, reduced in size, with a clearer cell wall, without septation and cytoplasmic organelles, with signs of degeneration. Patients with a less favorable clinical response were positive for mycological exams and presented predominance of parasites with a thickened, well pigmented cell wall with a tendency to cluster, septation and intracytoplasmic organelles, possibly resulting from an adaptive mechanism of sclerotic bodies to escape from the body’s defense. The amount of parasites recorded did not appear to interfere with clinical manifestation prior to treatment. However, the excessive number of parasites and the higher frequency of transepithelial elimination in the most exuberant cases suggest that host factors can determine the quality and quantity of the tissue response.

Limitations of this research were the absence of electron microscopy for a better evaluation of parasitic modifications and immunological techniques for investigation of circulating antigens that could complement the findings of the biopsies with other more specific and sensitive methods. In addition, it was not possible to locate the patients and verify their current skin condition, since most of them came from rural environments in other municipalities and lost hospital follow-up.

CONCLUSION

In this study, there was a greater tendency for clinical improvement or cure of tumoral forms when compared with plaque manifestation. There was concordance between the positivity of mycological and histological exams during treatment. The FM staining facilitated the demonstration of fungi in the epidermis. The most distinct tissue responses of mycosis were recorded in the dermis, with tissue and parasitic changes occurring during treatment with itraconazole, which may reflect parasite-host dynamics.

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Mailing address:
Kátia Sheylla Malta Purim
Serviço de Dermatologia
R. General Carneiro, 181- Centro
80060-900 - Curitiba, PR
Brazil
Email: kspurim@gmail.com

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