

Rhinosporidiosis - Cutaneous manifestation^{*}

Rinosporidiose - Manifestação cutânea

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Abstract: Rhinosporidiosis is an infectious mucocutaneous disease caused by *Rhinosporidium seeberi*. It is characterized by sessile or pedunculated polyps which are erythematous, moriform and friable and which mainly affect the ocular and nasal mucosa. The occurrence of skin lesions is occasional and due to dissemination from the adjacent mucosa, direct inoculation or hematogenous dissemination. The authors report the clinical case of an eight-year-old boy with an isolated lesion located in the medial epicanthus of the right eye.

Keywords: Communicable diseases; Parasitic diseases; Rhinosporidiosis; *Rhinosporidium*

Resumo: A rinosporidiose é uma doença infecciosa zoonótica mucocutânea causada pelo *Rhinosporidium seeberi*. Caracteriza-se por massa polipoide, séssil ou pedunculada, eritematosa, moriforme e friável, principalmente, nas mucosas nasais e oculares. A ocorrência na pele é ocasional, por disseminação a partir da mucosa adjacente, inoculação direta ou generalização via hematogênica. Os autores apresentam o caso clínico de um menino de oito anos de idade, com lesão isolada localizada no epicanto medial do olho direito.

Palavras-chave: Doenças parasitárias; Doenças transmissíveis; *Rhinosporidium*; Rinosporidiose

Rhinosporidiosis is an infectious mucocutaneous granulomatous disease caused by *Rhinosporidium seeberi*. It usually affects the nasopharynx, occasionally affecting the conjunctiva and the lacrimal sac, and sporadically, the urethra, genitalia, larynx, paranasal sinuses and the skin (Figures 1 and 2).^{1,3} Cases of rhinosporidiosis have been reported in the Americas, Europe, Africa and Asia, with high prevalence in India and Sri Lanka.^{1,2,4} It is endemic in the western region of northeastern Brazil. It occurs by inoculation of spores present in stagnant water or dust from fields. It is more common in men, especially affecting the nose. Eye infection is more prevalent in women.^{1,2,6,8} The histology and/or cytology performed after fine needle aspiration



FIGURE 1: A nine-year-old mulatto male patient from the state of Maranhao presenting a sessile tumor with an irregular, friable, erythematous, moriform surface with white dots and a diameter of 0.8 cm in the medial epicanthus of the left eye. The lesion had been present for three months. Surgical removal and electrocoagulation were performed

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FIGURE 2: Detail of the lesion

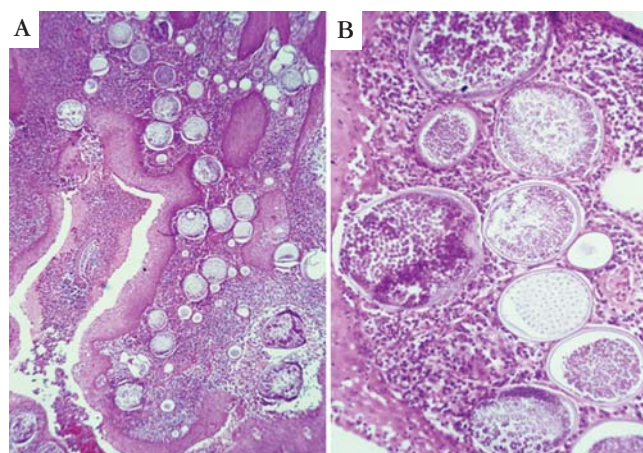


FIGURE 3: A. A histological examination revealed evolutionary forms of sporangia, young, collapsed and empty forms and bigger, mature forms near the surface of the epithelium (HE, 50X); B. The detail shows thousands of spores within the mature forms and inflammatory infiltrate consisting of lymphocytes, histiocytes, plasma cells, some eosinophils and foreign-body type giant cells (HE, 100X)

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(FNAC) serve to confirm the diagnosis (Figures 3 and 4A).³ The first reference was credited to Malbran (1896). Guilherme Seeber (1900) reported the causative agent and classified it as a protozoan from the *Coccidioidaceae* family. A phylogenetic analysis classified it as belonging to the group DRIPs (*Dermocystidium*, rosette agent, *Ichthyophorus*, and *Psorospernim*), an aquatic protistan parasite (Figure 4B).^{2, 4, 9} □

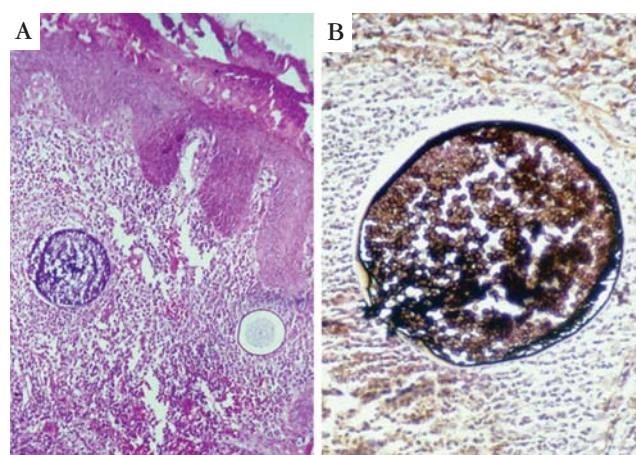


FIGURE 4: A. The stroma shows chronic inflammatory reaction with predominance of plasma cells, lymphocytes and few neutrophils (HE, 50X); B. the sporangia and spores are best visualized with the Grocott-Gomori stain (50X)

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