**CASE STUDY**

**Pulmonary manifestation of systemic mast cell disease**

M. Schmidt*, C. Dercken*, O. Loke*, S. Reimann**, S. Diederich‡, S. Blasius§, S. Heidenreich*

**ABSTRACT:** Systemic mast cell disease is a rare disease of unknown aetiology. Systemic infiltration and proliferation of mast cells in skin, bone marrow, gastrointestinal lymph nodes is the central pathological feature. This study reports a patient with mastocytosis of the skin (urticaria pigmentosa) for 10 yrs. The patient was referred to hospital for dyspnoea. Chest radiograph showed moderate reticular infiltration of both lungs, computerized tomography revealed multiple lymph nodes of the mediastinum and faint nodular lesions of middle and upper areas of lungs. Transbronchial biopsy demonstrated mast cell infiltration of the lung with formation of mast cell granuloma. According to the current literature, systemic mast cell disease with pulmonary involvement is a very rare entity. After a treatment with interferon α-2a over 6 months, the patient's condition and particularly dyspnoea showed improvement in parallel with an amelioration of the lesions as demonstrated by thorax computed tomography.


Mastocytosis is a local or systemic disease characterized by infiltration and expansion of mast cells in various tissues. Urticaria pigmentosa is diagnosed when skin is affected systemically showing multiple brown or livid papules. Whenever mast cell infiltration involves additional organs, the diagnosis of systemic mast cell disease (SMCD) is applied. Involvement of the gastrointestinal tract and bone marrow is most common. The patient presented here showed systemic mastocytosis with a biopsy-proven mast cell infiltration of the bone marrow and lung.

**Case report**

The present 54-yr-old male with a moderate smoking history had chronic pruritus for 10 yrs. Integument revealed exanthematic brown red papules with major involvement of the truncus. Rubbing of the skin evoked urticarial swelling. A cutaneous skin biopsy showed keratinocytes with increased pigmentation and proliferation of melanocytes. In the upper dermis, perivascular infiltrates consisted of lymphocytes, histiocytes and huge numbers of mast cells, all alterations indicative for urticaria pigmentosa. A total of 24 mast cells per high power field (magnification × 400) were counted. Figure 1 shows the truncal skin of the patient with the typical dermal manifestation of mast cell disease. The patient then presented with dyspnoea. The chest radiograph showed signs of interstitial lung disease with a fine reticular infiltration of both lungs (fig. 2A). Computed tomography (CT) of the thorax revealed faint nodular and cystic lesions of the middle and upper fields of the lungs with mediastinal adenopathy (fig. 2B). Blood gas analysis at rest gave a mild hypoxaemia with an arterial oxygentension ($P_{a}O_{2}$) of 8.5 kPa (64 mmHg). Pulmonary function tests showed a modest obstruction with a forced expiratory volume in one second (FEV1) of 3.05 L (77% of predicted value) and a normal vital capacity (VC) of 4.82 L (92% pred). Carbon monoxide transfer factor ($T_{L,CO}$) was markedly diminished (55% pred). Laboratory chemical examinations yielded a peripheral white blood count of 12.250 cells·mm$^{-3}$ with 65% neutrophiles, 25% lymphocytes, 5% monocytes, 4% eosinophilic cells and 1% basophilic cells. Red blood counts and thrombocytes were normal. Standard laboratory values of serum were all in normal range. Serum tryptase was 2.86 μg·mL$^{-1}$ (normal range <1 μg·mL$^{-1}$), histamine in 24 h urine collection was 72 μg·24 h$^{-1}$ (normal <50 μg·24 h$^{-1}$).

Biopsy of the bone marrow was performed and showed typical mast cell granuloma with focal cellular accumulations. Bone marrow presented a mast cell infiltration of 2%, lobulated mast cell nuclei were not found and fat cell content was normal. Because of the clinical symptoms

![Fig. 1. Torso of the presented patient showing generalized papulosis. After rubbing, urticarial skin lesions develop indicative for urticaria pigmentosa.](image-url)
of the patient and the described radiological findings, bronchoscopy and transbronchial fine needle biopsy was carried out. Bronchoscopy revealed signs of chronic bronchitis.

Histology showed infiltration of the lung interstitium by mast cells (basophilic cytoplasm and granula) with abortive mast cell granuloma formation. As shown in figure 3, mast cell granuloma consisted of lung interstitial cells, lymphocytes and mast cells. Mast cells showed signs of maturity as well as of immaturity which were distinguishable by the content of mast cell granula and size and shape of nuclei. A significant fibrosis was not visible.

Gastroscopy showed no pathological condition and biopsies could not demonstrate mast cell infiltration of the gastrointestinal. Cardiac evaluation including echocardiography was normal. For urticaria pigmentosa, therapy with antihistaminic agents was continued, and his pulmonary symptoms inhalative mast cell stabilizers (disodium chromoglycate) were applied. Since dyspnoea did not improve after 6 weeks, therapy with interferon α-2a (Roferon®; Roche, Grenzach, Germany) was started at a dose of 3 million IU. IE Roferon s.c. three times a week. After 6 months of treatment, CT of the thorax showed improvement of nodular and cystic lesions of the lung (fig. 2C). Size and number of mediastinal lymph nodes did not decrease significantly and pulmonary function tests were stable.

**Discussion**

Mastocytosis represents a wide spectrum of disorders characterized by proliferation and accumulation of tissue mast cells. In most patients, mast cell expansion is confined to the skin giving the diagnosis of cutaneous mastocytosis. In systemic mastocytosis different organs can be involved with the bone marrow and the gastrointestinal tract as the most frequently affected sites [1–3].

The lung is typically not involved in systemic mastocytosis although some studies described pulmonary involvement [4, 5]. In a very early study, solitary pulmonary mast cell granuloma or tumours have been described but at the time, pathologists had difficulty in distinguishing...
the histological pattern of mast cell tumours from plasmacytoma because staining procedures and immunohistochemical markers were not well-established [6].

According to a study by Travis et al. [7], where 58 patients with systemic mastocytosis were evaluated, respiratory symptoms occurred in 20% of the SMCD patients. The symptoms were dyspnoea and chronic bronchitis sometimes in association with asthma. Chest radiographic abnormalities were found in 16% of the patients. The abnormalities described were focal or scattered areas of fibrosis, bilateral interstitial fibrosis and multiple pulmonary nodules. In this study, suspected pulmonary involvement by mastocytosis was not confirmed by histology.

Concerning therapy of SMCD, all modes of treatment are aimed to relieve symptoms because curative therapy does not exist. Symptomatic treatment includes H1- and H2-antihistaminic drugs, prostaglandin blockers and mast cell stabilizers such as oral disodium cromoglycate [8–10].

For systemic and malignant mastocytosis different chemotherapeutical schemes exist all of which have been shown to be of limited value. Treatment with prednisolone and interferon α-2a has failed to improve the natural course of the disease but some cases showing a histologically proven response and relief of symptoms have been published [11–13]. Progression of symptoms under conventional therapy was the rationale to treat the presented patient with interferon α-2a. After treatment with this drug for 6 months, pulmonary function tests, the number of enlarged mediastinal lymph nodes and biochemical markers analysed in serum or urine did not improve significantly but symptoms such as dyspnoea and pruritus improved and computed tomography of the lung showed amelioration of interstitial lesions.

References