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Signal recognition particle (SRP) positive myositis in a patient with cryptogenic organizing pneumonia (COP)

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Abstract We report of a 46-year-old female patient with cryptogen organizing pneumonia preceeding the rare SRP positive necrotising myositis without cardiac involvement and no sign of dysphagia. Myositis showed full regression without oral immune suppression but with extracorporeal treatment, performed as a combined therapy of plasmaexchange and immunoadsorption. After 33-month of treatment, anti-SRP antibodies were not detectable any more.

Keywords Autoimmune myositis · Signal recognition particle · Cryptogen organizing pneumonia · Plasmaexchange · Immunoadsorption

Introduction

A 46-year-old female patient presented with progressive dyspnoea in January 2005. Lung function testing revealed a FEV1 of 1.96 l (61%), a vital capacity of 2.82 l (72%), FEV1/VC of 69%, and a diffusion capacity of 82% calculated to the alveolar volume. The chest X-ray revealed diffuse bilateral opacities of the lower lobes, a high resolution CT scan showed diffuse ground glass opacity of the left

upper and lower lobe as well as the middle lobe. Chemically a normal differential blood count and a normal total IgE (43 kU/l) were found. ASLT was marginally elevated and antinuclear antibodies against signal recognition particle (anti-SRP; titer >1:80) were found. Arterial blood gas analysis at rest was normal whereas at exertion (50 W for 5 min) pO₂ declined to 64 mmHg and pCO₂ was unchanged.

Due to dyspnoea, the patient was investigated in another hospital in December 2004. Bronchoscopy was done and bronchoalveolar lavage revealed a reduced CD4/CD8 ratio, a lymphocytosis of 37% and activated T cells. In the histology alveolitis with lymphocytes, plasma cells, histiocytes, sparse eosinophils but no granulomas were found. The patients mansion repeatedly suffered from water damage and mold growths on the walls of her indoor pool. Indoor air sampling was done, but mold exposure was in the low to moderate range. However, in blood no precipitating IgG antibodies against molds were found. Exogenic allergic alveolitis was suspected and the patient was started on oral steroids with an initial dose of 0.75 mg/kg, which was rapidly reduced.

First treatment: 1/2005 to 4/2005

Our initial steroid dose was 0.75 mg/kg (=Aprednislon 50 mg) and she improved markedly. FEV1 and VC rose to 2.40 and 3.55 l, respectively, however diffusion capacity went down to 79%. The course of lung function parameters is reported in Fig. 1 and Table 1. The HR-CT done in February already indicated regression of opacities and steroid was slowly reduced to a quarter of the initial dose (Aprednislon 12.5 mg). The course of HR-CT scans is shown in Fig. 2.

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Fig. 1 Dynamic of lung function tests (vital capacity VC, FEV1, and diffusion capacity TLCO/VA) Date: from 1/05 until 4/09

In April 2005, the patient clinically deteriorated, computed tomography again showed patchy bilateral opacities and the lung function parameters went down to a FEV1 of 1.70 l, VC of 2.38 l, and a diffusion capacity of 61%. To clarify the diagnosis an open lung biopsy was done in July

2005 [1–3]. Histology revealed centrilobular emphysema, activated alveolar cells, granulated polyps in the terminal bronchioli and alveolar ducts. Fibrin and foam cells were found in the alveoli. Diagnosis: cryptogenic organizing pneumonia [4].

Second treatment: 5/2005 to 4/2006

Steroid dosis was raised to 0.5 mg/kg (Aprednislon 37.5 mg) and the patient improved again. In August 2005, steroid treatment was reduced to 12.5 mg due to massive weight gain, but azathioprine 150 mg was added. In March 2006, steroid was reduced to 10 mg and azathioprine to 100 mg. Two weeks later, the patient was admitted to hospital due to dyspnoea in the course of a cold. Although there was no sign of muscle weakness, an elevated creatine kinase (CK) of 3000 U/l (–145 U/l) was found. As we attributed the elevated CK value to massive cough, no diagnostic steps were taken. Weeks later the patient developed

Table 1 Lung function and blood gas analysis parameters

	1/05	Maximum	Minimum	4/09
Vital capacity in liter and % pred	2.82 (72%)	3.55 (91%)	2.05 (53%)	2.31 (63%)
FEV1 in liter and % pred	1.96 (61%)	2.40 (74%)	1.31 (41%)	1.91 (60%)
Diffusion capacity/alveolar volume	82%	82%	55%	69%
pO ₂ in mm Hg	77	85	66	72
pCO ₂ in mm Hg	31	35	31	35

Fig. 2 HR-CT scan at the start of treatment (**a** 1/05), first improvement (**b** 2/05), first relapse (**c** 4/05), and final CT (**d** 1/09) at stable state (no disease)

Fig. 3 MRT scan at the start of plasmaapheresis showing massive muscle edema (**a** 8/06; CK > 10.000 U/l), improvement (**b** 10/07; CK 3500 U/l), and final MRT (**c** 12/08) at remission (CK in the normal range)

progressive weakness of her upper arms and legs with elevation deficit. She could barely get up from a chair. The neurologist suspected myositis.

Laborchemically, autoantibodies against the signal recognition particle (SRP, titer 1:5120) were found, which could be associated with myositis. Magnetic resonance tomography (MRT) showed muscle edema in several muscles of the pelvis and the upper legs corresponding to active myositis. The radiologic course of MRT is demonstrated in Fig. 3. An open muscle biopsy of the left sartorius muscle was done in May 2005. Histology revealed moderate,

immunologically mediated myositis with necrosis of single muscle fibers with partial regeneration, mild inflammatory cell infiltrates, HLA class-1 upregulation, and deposition of complement complexes in the terminal capillars. Diagnosis: myositis, possibly driven by an anti-SRP autoimmune mechanism.

SRP positive myositis may be associated with difficulties in swallowing and can also affect the heart [5]. The echocardiogram of the heart showed regular ventricular motility and the pulmonary arterial pressure was not elevated. Oesophageal motility was regular.

Fig. 4 Dynamics of creatinine kinase (CK) and myoglobin from the start of plasmaexchange in combination with immunoadsorption ($n = 82$). Date: from 7/06 until 9/08, since 9/08, CK is in the normal range

Third treatment: 5/2006 to 4/2009

Initial treatment consisted of a single dose of 1.000 mg prednisone on five consecutive days and two courses of high dose intravenous immunoglobulin (IVIG) 30.000 mg on five consecutive days (2×150.000 mg total dose, i.e., 2,000 mg/kg body weight), as she refused further long term steroid treatment [6]. However, muscle weakness as well as fatigue increased despite therapy and creatine kinase rose to a total of >10.000 U/l. Finally, the patient was admitted to the general hospital of Vienna (AKH) for immunoadsorption. Treatment was started in August 2006 using Ig-Therasorb®-columns three times a week. Immunoadsorption-procedure was performed as previously described [7, 8].

Due to unsatisfying treatment success, plasmaexchange (3L) was added to immunoadsorption. Within weeks of treatment the patient felt considerably better and her muscle strength improved although creatine kinase was stable at about 10.000 U/l. As myositis developed under steroid and azathioprine, we stopped azathioprine (8/2006), but kept the patient on 5 mg Aprednisone for another 3 months (11/2006). The MRT, done in December 2006 already revealed a significant decrease of muscle edema. Her physical condition improved slowly and she started taking regular walks of increasing distance.

In 2007 and 2008 the patient further improved, plasmaexchange in combination with immunoadsorption was done in intervals of once every 2–3 weeks and creatine kinase went down step by step (Fig. 4). Her physical fitness was fairly good, the daily training program went on, fatigue decreased and lung function parameters were completely stable with a diffusion capacity of 59%. Extracorporeal treatment was and still is tolerated excellently without a single side effect.

In 1/2009, plasmaapheresis is still done every 3 weeks although creatine kinase is almost in the normal range

(157 U/l). The patient is completely stable and lung function even improved. The HR-CT of 1/2009, as already in 10/2007, showed no sign of activity of COP but diffuse fibrosis especially in the middle lobe. Finally, the MRT of 11/2008 was completely normal with no sign of muscle edema. In April 2009, anti SRP antibodies could no longer be detected in the blood.

Discussion

We report of a patient with cryptogen organizing pneumonia and SRP positive necrotising myositis without cardiac involvement and no sign of dysphagia. This case is unusual, as myositis showed full regression without oral immune suppression but with extracorporeal treatment, performed as a combined therapy of plasmaexchange and immunoadsorption, which is said to be of little to no use in treating myositis [5, 9–12]. However, plasmaapheresis is already done for almost 3 years and is still continued without any side effects. According to Hengstman et al. [9, 13], treatment for myositis may be done for years.

The signal recognition particle is a protein targeting system, acting as an adaptor between protein synthesis and the translocation machinery in the membrane. Its function is the co-translocation transport of secretory and membrane proteins. The SRP cycle is a GTP dependent process [14]. In myositis, anti-SRP autoantibodies are found in 5% of cases.

Interstitial lung disease may be found in one quarter of patients suffering from anti-SRP autoimmune myositis, not differing in incidence from polymyositis or dermatomyositis [5, 9, 11, 16]. In our patient, cryptogenic organizing pneumonia occurred about one and a half years prior to the emergence of myositis, although anti-SRP antibodies were present from the very beginning. Although, we can only speculate if SPR-autoantibodies have induced cryptogenic organizing pneumonia, literature reports interstitial lung disease in about one-third of cases preceeding autoimmune myositis [5, 9, 17]. According to Tazelaar [18], polymyositis or dermatomyositis may be accompanied by either usual interstitial pneumonia (UIP), diffuse alveolar damage (DAD) or bronchiolitis obliterans organizing pneumonia (BOOP, now called COP). Diffuse alveolar damage is reported to have the worst prognosis and bronchiolitis obliterans organizing pneumonia (BOOP/COP) has the most favorable prognosis [10, 15, 19]. Anti-SRP antibodies, which are mainly polymyositis specific, may also be found in dermatomyositis or systemic sclerosis [5, 9, 13]. In a recent report, Suzuki et al. [20] suggest to classify SRP positive myositis as a particular form of polymyositis.

SRP positive myositis occurs in adults peaking at about 50 years (5), but it may also be found in children, although

very rarely [20, 21]. The onset of myositis is often accompanied by a viral infection, as was probably the case in our patient. Interestingly, myositis occurred despite immune suppression. It is noteworthy that we reduced the dosis of steroid and azathioprine prior to the emergence of myositis.

The first line treatment for SRP positive myositis is steroid (1 mg/kg), as second line treatment azathioprine or methotrexate is recommended [5]. Intravenous immunoglobulins (IVIG) may also suppress autoimmune disease by binding autoantibodies [6]. However, despite two courses of IVIG (2 g/kg BW) worsening of myositis occurred. Data on plasmapheresis in SRP positive myositis are very rare. Hengstmann et al. [9] reported of two patients in their study, however the treatment outcome was not mentioned.

If SPR-autoantibodies have induced COP, then we are dealing with the more common secondary cryptogenic organizing pneumonia. The first relapse of COP was probably attributed to reducing steroids too quickly, after raising steroids the disease activity was slowly declining. To which extent extracorporeal treatment helped to cure organizing pneumonia will remain unanswered. Lung function parameters were at its worse as myositis emerged, however during the course of myositis the patient never mentioned an impairment of her respiratory muscles.

Although some questionmarks remain, this is a rather unique case of the rare SRP positive myositis showing complete remission to plasmaexchange combined with immunoadsorption without additional immune suppression. Although a case report is no evidence for the efficacy of a treatment, also plasmapheresis may be useful in patients suffering from SRP positive myositis.

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