

Myositis-Like Syndrome Revealing Hypothyroidism

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Abstract

Introduction: Muscular involvement during hypothyroidism is rare and often infra-clinical. Symptomatic forms simulating authentic inflammatory myopathy remain exceptional and difficult to diagnose. We report a particular observation of primary hypothyroidism presenting initially by a “myositis-like” syndrome.

Case Report: A 66-year-old woman with no medical history was explored for myalgia associated with proximal muscular weakness, which had been evolving for the past month, with an increase in muscle enzymes: CPK at 1570 IU/l and LDH at 607 IU/l. She was sent to us for suspicion of polymyositis. The electromyogram was non-contributory and the muscle MRI was without abnormalities. Anti-nuclear antibodies and anti-soluble antigen antibodies were negative.

The thyroid assessment showed TSH at 15.45 μ IU/ml, fT4 at 10.55 pmol/l, anti-thyroglobulin antibodies positive at 1000 IU/ml, and anti-thyropoxidase positive at 488.5 IU/ml confirming the diagnosis of Hashimoto’s thyroiditis. Under thyroxine, the evolution was favorable with disappearance of muscular complaints and normalization of muscular enzymes.

Conclusion: The “myositis-like” syndrome is an exceptional presentation of hypothyroidism that represents a real diagnostic challenge for the clinician, especially since a true combination of thyroiditis and myositis is possible. It is recommended to screen for thyroid dysfunction in front of presentation suggestive of myositis, especially that this myosite-like syndrome may be the first inaugural manifestation of hypothyroidism.

Keywords: Myosite-like syndrome, hypothyroidism, CPK, myositis, thyroiditis.

INTRODUCTION

Muscular involvement during hypothyroidism is rare and often infra-clinical [1,2]. It may be of the type of myalgia, cramps, muscle weakness, muscle hypertrophy, and increased muscle enzyme, particularly creatine-phosphokinase (CPK) [3], and the most characteristic clinical presentation of the specific muscle involvement of hypothyroidism is the Hoffmann’s syndrome associating muscular stiffness and hypertrophy [4].

Rarely, symptomatic forms of myopathy of hypothyroidism can be very obvious and simulate true primary inflammatory myopathy [1,2,5]. These so-called “myositis-like syndrome” presentations remain exceptional and represent a real challenge for the clinician [1,6].

We report a particular observation of primary hypothyroidism revealed by a myositis-like syndrome.

CASE REPORT

A 66-year-old woman with no medical history was explored for myalgia associated with proximal muscular weakness, which had been evolving for the past month, with an increase in muscle enzymes in the city: CPK at 1570 IU/l and lactate dehydrogenase (LDH) at 607 IU/l. She was sent to us for suspicion of polymyositis. No medication, trauma or recent infections were noted.

The clinical examination noted a proximal muscle weakness at the four limbs, with muscle testing at 4-/4+. No neurological signs or skin lesions were noted. The rest of the somatic examination was without abnormalities. The biological assessment

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confirmed rhabdomyolysis with CPK at 1890 IU/L and LDH at 786 IU/l. The rest of the basic biological tests was without abnormalities (blood cell count, erythrocyte sedimentation rate, C-reactive protein, serum protein electrophoresis, creatinine, ionogram, liver enzymes, lipid parameters, troponin I_c, and urine analysis). The chest X-ray and the electrocardiogram were also without abnormalities. The electromyogram showed nonspecific myogenic changes. The immunological assessment of primary inflammatory myopathies was negative (anti-nuclear antibodies and anti-soluble antigens antibodies). MRI of the quadriceps muscles was also without abnormalities. Thus the diagnosis of primary inflammatory myopathy was eliminated. The thyroid status showed TSH at 15.45 μ IU/ml and fT₄ at 10.55 pmol/l. The thyroid immunological test confirmed the diagnosis of Hashimoto's thyroiditis with positive anti-thyroglobulin antibodies at 1000 IU/ml and anti-thyroperoxidase at 488.5 IU/ml. The evolution was favorable with the disappearance of myalgia and muscle weakness and normalization of muscle enzymes one month after the normalization of TSH under thyroxin: CPK at 86 IU/l and LDH at 240 IU/l.

DISCUSSION

The "myositis-like" syndrome defined as a muscular weakness associated with the elevation of muscle enzymes [2,7] is an exceptional manifestation of hypothyroidism [6]. Indeed, Madariaga MG's 2002 review of the literature, collecting all cases of myositis-like syndrome reported during hypothyroidism over a period of 25 years (1975-2000), found only 32 cases [7]. Similarly, in the large rheumatologic series of Leverenz D, of 192 patients referred for CPK > 1000 IU/l with myositis suspicion, hypothyroidism was at the origin of this syndrome "myositis-like" only in four patients, 2% [8].

This syndrome may be the first sign of hypothyroidism [1-3,9], and may remain the sole manifestation of this endocrinopathy [10].

The clinic can simulate point by point a true polymyositis [11], and more rarely skin signs may be added simulating even dermatomyositis [12,13]. This explains why, often, the hypothyroidism presenting with a myositis-like syndrome remains for a long time misdiagnosed and mismanaged [3].

These myositis-like presentations of hypothyroidism have been reported in both adults [1-3] and

children [8] and all muscles can be affected, but those of the shoulder and pelvic girdles are most commonly involved [3]. More rarely forms of diffuse myopathy with paraspinal muscle involvement can be seen, leading to severe presentations with even a camptocormia [2].

The electromyogram can be normal in 50% of cases; otherwise it can reveal minor and nonspecific myopathic changes [7]. Similarly, muscle biopsy is not contributory to the diagnosis, it shows nonspecific histological lesions such as: type II fiber atrophy, type I fiber hypertrophy, necrosis, central nuclei disposition, inflammatory infiltrate and the presence of core-like structures [7]. Sometimes perivascular lymphocyte infiltration is important, suggesting an authentic primary myositis [12]. The diagnostic challenge is even more difficult to solve, as the combination of hypothyroidism and primary inflammatory myopathy type polymyositis or dermatomyositis is possible [14-16].

Several authors recommend, to systematically screen for hypothyroidism in front of any muscular complaint that is not proven, and in particular in front of a myositis-like syndrome. This screening is recommended even in the absence of signs suggestive of hypothyroidism, since these signs can be masked by those of the primary myositis, especially in rapid and very evolutionary myositis [14]. Thyroid explorations, in front of a myositis-like syndrome, can exceptionally reveal, in addition to hypothyroidism, an associated thyroid cancer [16].

CONCLUSION

Myositis-like syndrome is an exceptional and unusual manifestation during hypothyroidism. It represents a real diagnostic challenge for the clinician, especially as the association of thyroiditis and authentic myositis exists. It is therefore necessary to request thyroid tests in front of any table suggestive of myositis, especially since this presentation may be the inaugural manifestation of hypothyroidism.

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