FIBRODYSPLASIA OSSIFICANS PROGRESSIVA

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SUMMARY - Fibrodysplasia (myositis) ossificans progressiva (FOP) is a rare autosomal dominant disorder in which there is a progressive ectopic ossification and skeletal malformation, mainly in the connective tissue of muscle. The diagnosis is based on the clinical findings and radiological demonstration of the skeletal malformations. We report a 5 year-old female case with FOP.

KEY-WORDS: fibrodysplasia ossificans progressiva, myositis, ossification.

Fibrodysplasia ossificans progressiva

RESUMO - Fibrodysplasia (miôsite) ossificante progressiva (FOP) é doença rara, de herança autossômica dominante, na qual ocorre ossificação ectópica progressiva e malformação esquelética, principalmente no tecido conectivo dos músculos. O diagnóstico é baseado nos achados clínicos e demonstração radiológica das malformações esqueléticas. Relatamos o caso de uma menina de 5 anos de idade com FOP.

PALAVRAS-CHAVE: fibrodysplasia ossificante progressiva, miôsite, ossificação.

Fibrodysplasia ossificans progressiva (FOP) is a rare autosomal dominant disorder in which there is a progressive ectopic ossification and skeletal malformation mainly in the connective tissue of muscles. The diagnosis is based on the clinical findings and radiological demonstration of the skeletal malformations. The management of the patients must be concentrated on the avoidance of exacerbating factors including trauma of the muscles, biopsies of the lumps, operations to excise ectopic bone, intramuscular injections and dental therapy. The treatment with disphosphonate disodium etidronate (EHDP), an inhibitor of calcification, is given in an attempt to suppress calcification of new lesions. We report a case of FOP.

CASE REPORT

SSM, 5 year-old female child has presented at FMM Hospital de Clínicas complaining of knees and left shoulder movement restriction and pain in legs while walking. Frequently, she's fallen. She does not relate trauma history. Pregnancy was uncomplicated and delivery was normal, at home without medical assistance. She sustained her head at 2 months. Although she crawled at 7 months and walked at 1 year-old, she has not sit until present days. Talking began at the expected age. There have been no significant family illnesses reported. On the physical examination, she has presented flexion deformity of the knees, left shoulder painful movement restriction, difficulty on walking resulting from the pain in legs, absence of muscular weakness, normal muscular tone and normal deep tendon reflexes. Radiological findings have shown ectopic bone in hips, thighs, legs and left shoulder (figure 1).
Figure 1. Patient SSM. Radiographs show ectopic bone in: A) hips and thighs; B) legs; C) left shoulder.

Routine laboratory tests were normal. The plasma calcium and inorganic phosphate were normal. The diagnosis was based on clinical and radiological findings. The EHDP, 20 mg/kg/day, was given to patient.

COMMENTS

FOP, also known as myositis ossificans progressiva, is a rare autosomal dominant disorder in which there is progressive ectopic ossification and characteristics skeletal malformations. The pathological process suggest that the primary lesion is in the connective tissue around the muscle. There is an exaggerated proliferation of connective tissue with subsequent dystrophic calcification and ossification, but in contrast, Smith et al. state that the muscular tissue is intrinsically abnormal before overgrowth of connective tissue.

Different parts of the body are variably affected by ectopic ossification. Early lesions often occur around the neck and over the back, whereas the most crippling ossification around hip may occur in late or early adult life. Ossification is unusual in smaller muscles and in those of the
abdomen. There are cases of circumscribed myositis of temporal muscle and femoral region, and a case associated with infection by Streptococcus in paravertebral area. The diagnosis depends upon the clinical and radiological demonstration of the characteristic skeletal malformation. Routine laboratory tests are usually normal. Actually, the bone image may be helpful in determining the full extent of the pathological process and more accurate than conventional roentgenographies, detecting lesions even before they can be observed with roentgenographies. The radiographs may lead to underestimate the severity of the disease. The biopsy of the lumps should be avoided.

Progression of disability is erratic, with severe limitation of movement in spine and shoulder by the age of 10 years; one or both hips were involved by the age of 20 years and most patients were confined to a chair or to bed with restriction of the joints by the age of 30 years. Progression of the disability did not appear to be influenced by any form of medical treatment.

There is no effective treatment. The management of the patients must be concentrated on the avoidance of exacerbating factors, including trauma of the muscles, biopsies of the lumps, surgeries to excise ectopic bone, intramuscular injections and dental therapy.

Due to our ignorance in this matter, the management of FOP remains empirical; nothing have been outstandingly successful. In our patient, we have used ethane-1-hydroxy-1,1-diphosphonate (EHDP) in an attempt to suppress calcification of the new lesions. The exact mechanism of its action is not known and it is possible that it may have important additional effects on bone cells, either directly or indirectly reducing their activity. Bruni et al. observed a significant improvement in symptoms and recovery of some of the active sites of ossification. Therapeutic progress is likely to be slow while the underlying cause remains unknown, but the advantage of relatively small increase in mobility in these young patients can be considerable.

REFERENCES