



## **WHOLE BODY MAGNETIC RESONANCE IMAGING IN JUVENILE DERMATOMYOSITIS. A LONGITUDINAL STUDY.**

CARLA HELENA CAPPELLO (UNICAMP, CAMPINAS, SP, Brasil), SERGIO DERTKIGIL (UNICAMP, CAMPINAS, SP, Brasil), EDUARDO BRONZATTO (UNICAMP, CAMPINAS, SP, Brasil), ROBERTO MARINI (UNICAMP, CAMPINAS, SP, Brasil), SIMONE APPENZELLER (UNICAMP, CAMPINAS, SP, Brasil)

### **BACKGROUND**

Juvenile dermatomyositis (JDM) is a rare multisystem connective tissues disorder of unknown aetiology.

Assessment of disease activity is a challenge in clinical practice.

### **MATERIALS AND METHODS**

We included consecutive JDM patients followed in the rheumatology unit. All patients were submitted to clinical

and laboratory evaluation. WB-MRI images were obtained using a 1.5 T MRI scanner and short T inversion recovery sequences (STIR). Muscle, peripheral inflammation and subcutaneous inflammations signal abnormalities

were scored in 42 muscular groups. Muscle inflammation was classified as: 0 = absent; 1 = Mild to moderate /

involvement less than 50% of muscle extension and 2 = Accentuated / greater than 50%. Peripheral and subcutaneous inflammations were classified as: 0 = absent; 1 = present; and on proximal and distal extremities.

WB-MRI and clinical assessments were performed concurrently and results compared. Evaluation was repeated

after 12 months. Statistics was performed according to the nature of the variable.

### **RESULTS**

WB-MRI revealed muscle inflammation in 6 (31.6%) at study entry. We observed grade 2 muscle inflammation of

the right and left scapular girdle (1/19 patients), right and left pelvic girdle (2/19 patients) and right and left thigh

(1/19 patients). Grade 1 inflammation was observed in peripheral right and left arm (2/19 patients), peripheral

right and left thigh (1/19 patients). Grade 1 subcutaneous inflammation was observed in right and left thigh (1/19

patients) and left leg (1/19 patients). Additionally we observed sacroiliitis (1/19 patients), spinal cord infarction

(21%) and osteonecrosis (5.2%). All patients were treated with standardized treatment. After 12 months 13/19

(68.4%) patients repeated the WB-MRI. Five (38.4%) patients had new/worsening of muscle and subcutaneous

inflammation, one (7.7%) patient had tibial medullary infarction. Correlations between WB-MRI muscle score and

disease activity measures were excellent (Manual Muscle Test:  $r=-0.88$ , Childhood Myositis Assessment Scale:

$r=-0.81$ ). Patients with subcutaneous inflammation developed clinically evident subcutaneous calcifications

during follow-up.

## **CONCLUSION**

WB-MRI provides additional information to clinical evaluation and represents a promising tool to determine the

grade of muscle inflammation to additional peripheral and subcutaneous tissue inflammation