Conclusions: In a Swedish population with RA, ERS-RA performed well in identifying patients with a very low and very high CV 10 year CV risk. In clinical routine practice, ERS-RA could be used to identify low and high risk individuals, who might be considered for additional CV risk factor evaluation and subsequent intervention.

REFERENCES:

Disclosure of Interest: None declared

THU0145 CHANGES IN BONE METABOLISM AND TRABECULAR BONE SCORE IN PATIENTS WITH EARLY RHEUMATOID ARTHRITIS
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Background: Rheumatoid arthritis (RA) is a systemic inflammatory disease which can lead to bone and joint damage including local bone erosion and general osteoporosis. Dual X-ray Absorptiometry (DXA) is the established standard for measuring Bone Mineral Density (BMD), but it does not provide any informations about the bone microarchitecture, which is an essential parameter to define bone strength. The Trabecular Bone Score (TBS) is a new structural parameter that can be obtained by DXA scanning and it is related to bone microarchitecture and provides data on bone quality irrespective of bone density.

Objectives: The aim of this study is to evaluate the changes of BMD, TBS and bone remodelling parameters in subjects with recent-onset rheumatoid arthritis, treated or not with high doses of glucocorticoid, compared to age and sex matched healthy controls

Methods: The study included 42 subjects (31F, 11M), fulfilling the 2010 EULAR/ACR diagnostic criteria for RA and recent onset of joint symptoms (<6 months of synovitis) (Early RA), which were treated according to the current EULAR guidelines. As control group, 25 sex and age matched healthy subjects (21F, 4 M) were recruited. Post-menopausal women were excluded from the study. Lumbar spine and femoral BMD and TBS were evaluated at recruitment and after 12 months. The following parameters of bone remodelling and regulatory cytokines were measured at recruitment and every 3 months for 12 months: type I N-terminal pro-peptide (P1NP), osteocalcin, alkaline phosphatase, sclerostin, dickkopf-1 (DKK1), osteogrostererin (OPG), Receptor Activator of Nuclear Factor Kappa-B ligand (RANKL). The clinical and demographic characteristics, including disease activity index and quality of life, were also evaluated

Results: No difference in BMD (spine and hip) and TBS values were detected between Early RA and control group at recruitment time (0.893 g/cm² vs 0.972 g/cm²; 0.790 g/cm² vs 0.770 g/cm²; 1598 vs 1521 respectively). After 12 months, the BMD at spine and hip and TBS values were significantly lower in patients with RA compared to healthy controls (0,601 g/cm² vs 1,011 g/cm²; 0,560 g/cm² vs 0,981 g/cm²; 1335 vs 1488 respectively). After 12 months, patients treated with high-dose of corticosteroids showed lower mean TBS values compared to patients untreated or treated with low-dose of corticosteroids (1,210 vs 1,430), whereas BMD values were similar. No differences were observed in osteocalcin and ALP between Early RA patients and healthy subjects at any time. Compared to healthy subjects, Early RA patients showed a significantly higher RANKL/OPG ratio and DKK1 serum levels, beginning from 6 month of observation, that correlated with disease activity (DAS28)

Conclusions: These preliminary data confirm that even in the early stages of disease, RA exert a negative effect on bone metabolism, whose pathogenesis is very likely involved by dysfunctions of bone homeostasis. This leads to a reduction of BMD and to changes of parameters of bone quality (TBS), that are more pronounced in patients treated with higher doses of corticosteroids compared to patients treated with lower corticosteroid doses

REFERENCE:

Disclosure of Interest: None declared

THU0146 GLYCEMIC PROFILE AND INSULIN RESISTANCE IN PATIENTS WITH EARLY RHEUMATOID ARTHRITIS
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Background: In established rheumatoid arthritis (RA) there is a high incidence of patients with increased insulin resistance, which can favour the development of Diabetes Mellitus (DM) and the appearance of cardiovascular complications. These aspects, however, have not been studied in depth in patients with early rheumatoid arthritis.

Objectives: To describe the glycemic profile and insulin resistance (IR) in patients with early diagnosis of RA who had not received any background treatment or steroids.

Methods: Observational study in which patients were included 18 years old, diagnosed with RA according to criteria ACR 1987 and/or ACR-EULAR 2010 from the service of Arthritis of Recent Beginning of the Hospital Universitario Central de Asturias, between December 2015 and December 2017. In the basal visit, values of insulin, glucose, fasting glycosylated haemoglobin (HbA1c), body mass index (BMI) and abdominal perimeter were collected. In addition, RI and beta cell dysfunction were estimated with HOMA-IR and HOMA-B respectively.

Results: 66 patients were collected, of which 4 were excluded because they were known to be DM and 17 because they had not completed the analytical studies. Of the 45 that were finally studied, 80% (36 patients) are women, the mean age of 54.1±12.9 years, the progression of the disease to diagnosis is 22.8±15.3 weeks, 46.7% (21 patients) are FR and AGPA positive, and the mean disease activity measured by SDAI is 28.0±12.2.

Eighteen patients (45%) and 71.1% (31.5 patients) had HOMA-IR, with both 17 patients (37.7%) showing a statistically significant association of SDAI>11 (p=0.001), BMI >24 (p=0.03) and an increased abdominal perimeter (p=0.013).

Based on the WHO’s diagnostic criteria for DM, 2 patients (4.4%) were diagnosed with DM (all based on a previously unknown HbA1c>6.5%) and 21 (46.7%) as pre-diabetic (14 with glucose between 100–125,16 with HbA1c>5.7 and 9 patients with both alterations). In this subgroup the mean of HOMA-IR and HOMA-B was higher than the mean of the complete sample, 3.4 and 139 respectively.

These patients compared with the rest of the sample had a statistically significant association with BMI >24 (p=0.07), SDAI >11 (p=0.01).

No correlations were found between these metabolic alterations and the sex, age or positivity for RF.

Conclusions: In patients with AR of less than one year of evolution and who have not received previous treatment with FAME or steroids, a high percentage of RI, pancreatic B cell dysfunction and alterations of the glycemic profile are detected. These alterations are significantly correlated with the presence of overweight and obesity, as well as with the high degree of activity of the disease. Follow-up of these patients is necessary to determine the effect of therapy for rheumatoid arthritis on these metabolic alterations.

Disclosure of Interest: None declared

THU0147 TREATMENT RESPONSE IN ANTIDEPRESSANTS-TREATED RA PATIENTS WITH DEPRESSIVE AND ANXIETY DISORDERS RECEIVING DMARDS AND BIOLOGICS ON A FIVE-YEAR FOLLOW-UP
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Background: Anxiety and depressive disorders (ADD) significantly affect disease activity and prognosis, treatment compliance and response in rheumatoid...