Objectives: To clarify how disease activity differs for each generation, especially in menopausal period.

Methods: Using the Japanese large RA cohort database (NinJaNational database of Rheumatic Diseases in Japan) of 2016, we divided 12257 RA females into three groups of age (under 44 years old=1, 45 to 55 years old=2; defined as a menopausal group, over 56 years old=3) and analysed them cross-sectionally. We conducted a one-way ANOVA on disease activity indexes such as Tender joint count (TJC), Swollen joint count (SJc), DAS28, HAQ-DI.

Results: Table 1 shows the number of people per group, the duration of disease, the titer of RF/ACPA, and the proportion of drugs used. The average usage of prednisone and the use of biologics was the most common in group 1. In table 2, TJC was the largest in group 2 (p<0.01). Furthermore, the difference between groups seen in TJC tends to be larger than TJC 28. There was no significant difference in SJc (SJc28) between three groups. Other disease activity indicators (ESR, CRP, DAS 28, HAQ DI) were the largest in group 3 (p<0.01) and the percentage of Boolean remission was also lowest in group 3 (p<0.02).

Abstract AB0255 – Table 1. Characteristics of the three age groups

<table>
<thead>
<tr>
<th>group</th>
<th>1</th>
<th>2</th>
<th>3</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of patients</td>
<td>1071</td>
<td>1699</td>
<td>9487</td>
<td>0.001</td>
</tr>
<tr>
<td>Symptom duration, years</td>
<td>8.2</td>
<td>10.5</td>
<td>15.4</td>
<td>p&lt;0.01</td>
</tr>
<tr>
<td>RF titer, (IU/mL)</td>
<td>233</td>
<td>254</td>
<td>232</td>
<td>p=0.06</td>
</tr>
<tr>
<td>ACPA titer, (IU/mL)</td>
<td>90</td>
<td>102</td>
<td>132</td>
<td>p&lt;0.01</td>
</tr>
<tr>
<td>On NSAIDs use, n(%)</td>
<td>392</td>
<td>570</td>
<td>824</td>
<td>p&lt;0.01</td>
</tr>
<tr>
<td>Symptom duration, years</td>
<td>8.7</td>
<td>10.5</td>
<td>15.4</td>
<td>p&lt;0.01</td>
</tr>
<tr>
<td>RF titer, (IU/mL)</td>
<td>233</td>
<td>254</td>
<td>232</td>
<td>p=0.06</td>
</tr>
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</tr>
</tbody>
</table>

Disclosures of Interest: None declared

Disclosure of Interest: None declared

AB0256

DIFFERENTIAL DIAGNOSIS OF SERONEGATIVE RA: CALCIUM PYROPHOSPHATE DIHYDRATE DEPOSITION DISEASE

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Background: Calcium pyrophosphate deposition (CPPD) disease is caused by calcium pyrophosphate (CPP) crystals and seen mainly in elderly. Clinical presentation can be heterogeneous. The arthropathy of CPPD may mimic RA, particularly if involving common joints seen in RA. Diagnosis of CPPD arthritis is based on CPP crystals seen in synovial fluid (SF) analysis, chondrocalcinosis (CC) seen in radiographs and/or on typical clinical presentation for CPPD. Early diagnosis of CPPD can be challenging and a proportion of CPPD patients may be misdiagnosed. We demonstrate 17 cases with CPPD initially diagnosed and treated as seronegative RA.

AB0257

PHYSICAL ACTIVITY IN EARLY AND LONG-STANDING RA – RELATIONS TO DISEASE ACTIVITY, CARDIOVASCULAR RISK FACTORS AND ATHEROSCLEROSIS

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Background: The excess risk for cardiovascular disease (CVD) in Rheumatoid Arthritis (RA), is partly attributable to traditional cardiovascular risk factors for CVD1,2 and systemic inflammation.3,4 The aim of this cross-sectional study was to objectively measure and to analyse possible associations with disease activity, risk factors for CVD and measures of subclinical atherosclerosis.

Methods: This study included 84 patients with early and 37 with long-standing RA (disease duration, mean [SD] 1.4 [0.4] and 18.3 [2.3] years respectively). Physical activity was measured using a combined accelerometer and heart rate monitor and included total physical activity (counts/min), proportion of moderate to vigorous physical activity (MVPA) and sedentary time. Further assessments were; disease activity (ESR, DAS28), functional ability (HAQ), risk factors for CVD (blood lipids, i.e., triglycerides, high density lipoprotein [HDL], low density protein [LDL], blood glucose, blood pressure, waist circumference, body mass index [BMI]), body fat (Dual-energy X-ray), and early signs of atherosclerosis (pulse wave velocity [PWV], augmentation index [AIx] and carotid intima-media thickness [cIMT]).

Results: Physical activity variables did not differ between patients with early and long-standing RA. Thirty-seven of the patients with early and 43 of the patients with long-standing RA did not reach WHO's recommended levels of MVPA. Univariate linear regression analyses with the two groups combined showed associations between total physical activity and younger age, lower values for HAQ and disease activity (ESR), as well as more beneficial values

REFERENCES:


Disclosure of Interest: None declared

for blood glucose, triglycerides, waist circumference, BMI, body fat, sleeping heart rate (SHR), systolic, diastolic and central blood pressure and pulse pressure, AIX, PWV, and cIMT. More time spent in MVPA was associated with younger age and with favourable values of blood glucose, HDL, LDL, waist circumference, SHR and PWV.

Abstract AB0257 – Table 1. Physical activity variables in patients with early and long-standing RA, presented as median with inter-quartile range (IQR). P-value refers to Mann-Whitney U-test.

<table>
<thead>
<tr>
<th></th>
<th>Early RA</th>
<th>Long-standing RA</th>
</tr>
</thead>
<tbody>
<tr>
<td>(n=84)</td>
<td>(n=37)</td>
<td></td>
</tr>
<tr>
<td>Accelerometry (mean counts/minute)</td>
<td>35.7 (29.9)</td>
<td>38.1 (24.3)</td>
</tr>
<tr>
<td>MVPA (% of wear time)</td>
<td>3.2 (6.6)</td>
<td>2.5 (5.5)</td>
</tr>
<tr>
<td>Sedentary time (% of wear time)</td>
<td>53.9 (11.0)</td>
<td>52.0 (13.3)</td>
</tr>
</tbody>
</table>

MVPA=Moderate to Vigorous Physical Activity=3+1.75 x resting heart rate. Sedentary time=heart rate data with zero accelerometer counts.

Conclusions: Physical activity behaviour was similar in patients with early and long-standing RA. Total physical activity as well as more time spent in moderate to vigorous physical activity were associated with more favourable risk factors for CVD and measures of atherosclerosis. These results stress the importance of promoting physical activity in patients with RA.

REFERENCES:

Disclosure of Interest: None declared

AB0258

IMPACT OF CONTROLLING DISEASE ACTIVITY ON REGAINING NORMAL PHYSICAL FUNCTION, AND ACHIEVING NO OR LIMITED PAIN IN PATIENTS WITH RHEUMATOID ARTHRITIS TREATED WITH BARICITINIB

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Background: Remission or low disease activity (LDA) are the recommended treatment (tx) targets in rheumatoid arthritis (RA).1 It is still unknown whether achieving remission/LDA is associated with normalisation of physical function, and limiting pain.

Objectives: To describe the impact of baricitinib (BARI) tx on regaining normal physical function, and achieving no/limited pain in patients (pts) who achieved remission or LDA, or remained in moderate or high disease activity (MDA, HDA).

Methods: This is a post-hoc analysis of RA-BEAM (NCT01710358) and RA-BEGIN (NCT01711359). Mutually exclusive categories were defined as clinical disease activity index (CDAI) scores of £2.8 (remission), >2.8 to £10 (LDA), >10 to £22 (MDA), and >22 (HDA). Last observation carried forward was used for pain visual analogue scale (0–100 mm VAS) and Health Assessment Questionnaire-Disability Index (HAQ-DI) to impute missing values. Descriptive analyses of the pts achieving normalisation of physical function was defined by a HAQ-DI score of <0.5 (normative value), and limited/no pain by pain VAS of £10 mm at week (wk) 12 and 24 as a function of disease activity.

Results: Overall, 1228 pts in RA-BEAM (448, PBO+MTX; 471, BARI +MTX; 309, ADA +MTX) and 543 pts in RA-BEGIN (190, MTX; 156, BARI; 197, BARI +MTX) were included. In RA-BEAM, among pts in remission at wk 12, % pts achieving limited/no pain was numerically higher in BARI (83%; 33/40) group compared with ADA (73%; 16/22) and PBO (67%; 6/9); at wk 24, these percentages were 81% (61/75), 82% (32/39), and 63% (12/19) for BARI, ADA, and PBO, respectively. Among pts who achieved remission on BARI +MTX tx, normal physical function was reported in 65% (26/40) and 73% (55/75) of pts at wk 12 and 24, respectively (Fig 1). For ADA +MTX treated pts, the proportion was 73% (16/22) at wk 12 and 69% (27/39) at wk 24. In RA-BEGIN, among pts in remission, % pts with limited/no pain at wk 12 was numerically higher for BARI (96%; 21/22) compared with BARI +MTX (82%; 32/39) or MTX (64%; 9/14); limited/no pain at wk 24 was reported in 68% (23/34), 87% (40/46), and 77% (17/22) of pts treated with BARI, BARI +MTX, and MTX, respectively. Among pts in remission, % pts achieving normal HAQ-DI at wk 12 and 24 with BARI monotherapy were 91% (20/22) and 82% (28/34); BARI +MTX, 77% (30/39) and 91% (42/46); and MTX monotherapy, 79% (11/14) and 82% (18/22), respectively.

Abstract AB0258 – Figure 1. Percentage of pts who achieved pain VAS £10 mm, and HAQ-DI <0.5 scores for the different disease activity states at wk 12

Percentage indicates the pts who regained normal physical functions (HAQ-DI <0.5), and low/none pain (pain VAS £10 mm) with different disease activities (CDAI score of £2.8 [remission], >2.8 to £10 [LDA], 10 to £22 [MDA], >22 [HDA]) in different treatment groups.

*In RA-BEAM, MTX was given as a background therapy among all treatment groups.

Conclusions: These data support that controlling the disease activity by achieving remission or LDA increases the chances to regain normal physical function and relieve pain, independent of the tx. The data from RA-BEAM may indicate that achieving limited/no pain at wk 12 of RA may be more likely with BARI vs ADA, when being in remission.

REFERENCE: