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found statistically significant associations between pain VAS and depression score, impacts of pain on daily behaviors and between pain VAS and DAS-28.

Conclusions: Almost 40% of patients had moderate to severe pain in a population of severe RA followed in hospital centers and treated with biotherapies in more than 80% of cases. Therefore, pain is still a major outcome to consider in RA. But, the proportion of patients with moderate to severe pain is less important than that one published by Taylor et al. Pain is associated with the DAS-28 score, but also with the depression score.

References:

[1] Taylor P et al. J Int Med Res 2010; 38: 1213-24.

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SAT0124 HIGHER BMI AND SHORTER DISEASE DURATION ARE ASSOCIATED WITH "FIBROMYALGIC" RHEUMATOID ARTHRITIS - EVALUATION USING POWER DOPPLER ULTRASONOGRAPHY

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Background: Power Doppler ultrasonography (PDUS) can differentiate RA patients who truly have active disease. Patients with persistent joint tenderness may not necessarily have true inflammation; yet composite disease activity indices remain high, leading to treatment escalation. These patients are often referred to having "fibromyalgic RA" where tender joints far exceed that of swollen joints.

Objectives: We evaluated the predictors of "fibromyalgic RA (FMRA)," defined as patient-reported tender joint count (TJC) at least 7 greater than PDUS joint count. Methods: Consecutive RA patients were recruited in a tertiary rheumatology centre in Singapore. Patients self-assessed 28 joints for tenderness, followed by blinded PDUS assessment of the corresponding joints. Patients were grouped into either, (i) PDUS joint count > patient-reported TJC, (ii) PDUS joint count = patient-reported TJC (iii) patient-reported TJC > PDUS joint count, (difference <7) (iv) patient-reported TJC > PDUS joint count, (difference ≥7). Group (iv) is defined as FMRA. Predictors of FMRA were evaluated by multinomial logistic regression

Results: Of the 101 patients, 81% were female, 72% Chinese, median age of 53.8 (IQR 48.1;62.4) and disease duration of 6.2 (1.9;8.8) years. Median BMI was 23.6 (20.8;27.0) and DAS28 was 3.2 (2.5;4.2). FMRA patients (15%) had a high median patient-reported TJC of 13 (10;18) but low median PDUS joint count of 1 (0;4). Compared to the other groups, they had higher median DAS28 of 4.6 (3.3;5.2), shorter mean disease duration of 2.3 (0.4;6.8) years, and higher median BMI of 26.9 (23.7;32.2). No differences in mean ESR or proportion of patients with seropositive disease were observed. Using Group (i) as the reference, patients with FMRA were more likely to have a higher BMI (adjusted OR=1.6, 95% CI 1.2; 2.1,p=0.001 for every unit increase) and shorter disease duration (adjusted OR=0.8, 95% CI 0.7;0.98,p=0.03 for every additional year). Age, gender, ethnicity, disease activity (PDUS score) and seropositivity were not significant predictors of

Conclusions: Higher BMI and shorter disease duration were independent predictors of FMRA. Use of PDUS would be useful in this group, rather than basing treatment decisions on composite disease activity indices alone.

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SAT0125 TENOSYNOVITIS IN RHEUMATOID ARTHRITIS: PREVALENCE AND DETERMINANTS OF TENOSYNOVITIS DETECTED IN THE SONAR-ULTRASOUND EXAMINATION IN THE SCQM COHORT

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Background: Tenosynovitis is one of the common features of rheumatoid arthritis (RA). The diagnosis of tenosynovitis is frequently made clinically but for the detection of tenosynovitis, ultrasound (US) is more sensitive. US and MRI detected tenosynovitis is a predictor for unstable remission and erosive progression in RA

Objectives: The aim of this study was to assess the prevalence and determinants of TS detected in the SONAR (Swiss Sonography Group in Arthritis and Rheumatism) examinations in patients with Rheumatoid Arthritis in the SCQM

Methods: The SONAR ultrasound examination consists of a semi-quantitative score employing both multiplanar gray scale (B-mode) and Doppler-mode (PwD) and a TS composite score (grade 0-3). Pathologic TS was defined as TS grade 2-3. Characteristics of patients with and without TS are shown using standard descriptive methods. In a longitudinal sub-group the change in TS from no TS to pathologic TS or vice versa and DAS28 over time was categorized as "worse" . $(\Delta DAS28 \ge 2.1)$, "same" and "better" ($\Delta DAS28 \le -2.1$) and the correlation between change DAS28 and in tenosynovitis was evaluated.

Results: 941 RA patients with TS score were available. 20% of included patients showed signs of TS. The presence of TS was associated with male gender and higher values of disease activity and physical function disability. Furthermore, 15% of patients in DAS28 Remission had sonographic TS. TS was less frequently observed in patients on biologic therapies (Table 1). The longitiudinal sub-group consisted of 348 patients. The correlation between change in DAS28 and change in TS was poor (polychoric correlation 0.28 [0.14, 0.43]).

Table 1. Patient Characteristics at first Tenosynovitis measurement

Parameter	Tenosynovitis		OR [95% CI]
	Yes (N=184)	No (N=757)	
Male gender, n (%)	59 (32)*	158 (21)*	1.8 [1.2, 2.5]*
Mean age in years	57	56	1.0 [0.99, 1.02]
Mean disease duration in years	5	3	0.98 [0.96, 1.00]
RF positive, n (%)	123 (68)	511 (70)	0.9 [0.7, 1.3]
Anti-CCP positive, n (%)	109 (65)	477 (70)	1.1 [0.6, 2.1]
DAS28 Remission, n (%)	24 (15)*	232 (38)*	0.3 [0.2, 0.5]*
Mean DAS28	3.9*	3.1*	1.6 [1.4, 1.8]*
Mean B-Mode SONAR Score	12.0*	8.0*	1.1 [1.04, 1.08]*
Mean PD-SONAR Score	2.0*	1.0*	1.0 [1.06, 1.12]*
Mean Physician Global Assessment	4.0*	2.0*	1.4 [1.3, 1.5]*
Mean HAQ Score	0.8*	0.5*	1.6 [1.1, 2.2]*
On biologic, n (%)	61 (33)*	399 (53)*	0.4 [0.3, 0.6]*
On synthethic DMARD, n (%)	110 (60)	500 (66)	0.8 [0.6, 1.1]*

^{*}p<0.05.

Conclusions: In the SCQM RA cohort TS is associated with male gender and an overall higher disease activity. One should actively look for TS even in RA patients in DAS28 remission.

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SAT0126 MENTAL HEALTH BENEFITS ASSOCIATED WITH REDUCTION IN DISEASE ACTIVITY AMONG RHEUMATOID ARTHRITIS PATIENTS: A SYSTEMATIC REVIEW OF THE LITERATURE

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Background: The association between physical functioning and rheumatoid arthritis (RA) disease activity is well established. In randomized controlled trials (RCTs) of RA treatment, reductions in disease activity result in improved physical functioning. The mental health benefits associated with reductions in disease activity are less well recognized.

Objectives: Investigate the association between reductions in RA disease activity and measures of mental health status.

Methods: A systematic review of the published literature of RCTs in RA was conducted. The search was limited to those published in the last 10 years that included the Short-Form (SF)-36 Health Survey as an outcome measure.

Results: The search strategy yielded 77 articles of which 38 were excluded for the following reasons: not RCT (23); focus exclusively on early RA (9); psychometric evaluation (4); review study (1); and off topic (1). Of the 39 articles reporting mean changes in scores on the SF-36 mental health (MH) and role emotional (RE) domains by treatment group, 3 articles also presented these changes in association with reductions in RA disease activity measures. 1-3 In relation to American College of Rheumatology (ACR) response criteria, small to moderate (0.2-0.5 standard deviation [SD]) effect size changes in RE and MH domain scores were observed among RA patients whose ACR responses ranged from 20-49%. Moderate to large effect sizes (0.5-0.8 SD) were reported in RE and MH domain change scores among ACR50 responders. Similar changes in RE and MH domain scores were reported by patients with reductions in Simplified Disease Activity Index (SDAI) of ≥10 points.

Conclusions: The benefits of clinically meaningful reductions in disease activity among RA patients not only include improved physical functioning but also meaningful improvements in overall mental health status. The improvement in mental health is positively associated with reduction in disease activity, with only small to moderate improvements associated with less than ACR50 responses. Additional work is needed to investigate the incremental effects of RA therapies on mental health beyond the effects related to improvements in disease activity. References:

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Disclosure of Interest: V. Strand Consultant for: Abbvie, Amgen, AstraZeneca, BiogenIdec, Boehringer Ingelheim, Celltrion, Crescendo, Genentech/Roche, GSK, Janssen, Lilly, Merck, Novartis, Pfizer, Regeneron, Samsung, Sanofi and UCB, R. Rendas-Baum Employee of: Optum, M. Kosinski Employee of: Optum, D. Brooks Shareholder of: GlaxoSmithKline, Employee of: GlaxoSmithKline, R. Ganguly Shareholder of: GlaxoSmithKline, Employee of: GlaxoSmithKline

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