Disease Activity, Functional Ability and Nutritional Status in Patients with Rheumatoid Arthritis: An Observational Study in Greece

Anastasia G. Markaki, Konstantinos Gkiouras, Christos Papakitsos, Maria G. Grammatikopoulou, Athena Papatsaraki, Roula Ioannou, Amalia Tsagkari, Theodora Papamitsou, Dimitrios P. Bogdanos

Mediterr J Rheumatol 2020;31(4):406-11
Disease Activity, Functional Ability and Nutritional Status in Patients with Rheumatoid Arthritis: An Observational Study in Greece

Anastasia G. Markaki1, Konstantinos Gkiouras2,3, Christos Papakitsos5, Maria G. Grammatikopoulou2, Athena Papatsaraki1, Roula Ioannou1, Amalia Tsagkaris2, Theodora Papamitsou6, Dimitrios P. Bogdanos2

1Department of Nutrition & Dietetics Sciences, Faculty of Health Sciences, Hellenic Mediterranean University, Sitia, Crete, Greece, 2Department of Rheumatology and Clinical Immunology, Faculty of Medicine, School of Health Sciences, University of Thessaly, Larissa, Greece, 3Department of Medicine, School of Health Sciences, Aristotle University of Thessaloniki, Thessaloniki, Greece, 4Department of Nutrition, KAT General Hospital, Athens, Greece, 5Health School, Aegean College, Athens, Greece, 6Department of Histology and Embryology, Department of Medicine, School of Health Sciences, Aristotle University Thessaloniki, Thessaloniki, Greece

ABSTRACT

Aim: The aim of the present pilot study was to assess differences in the nutritional status, Mediterranean diet (MD) adherence, and functional ability among patients with rheumatoid arthritis (RA), according to disease activity. Methods: A total of 48 patients with RA, outpatients of a hospital in Athens, Greece were recruited. Disease activity was evaluated with DAS28, functional status with the Health Assessment Questionnaire (HAQ), MD adherence with the MedDietScore and malnutrition with the patient-generated subjective global assessment (PG-SGA). Results: A relationship was noted between DAS28 and HAQ, indicating a reduced functional status with increased RA activity. Although MD adherence differed between DAS28 categories, no specific differences were noted in the PG-SGA or the MedDietScore in the post-hoc analyses. According to the PG-SGA, no need for nutritional intervention was noted among participants. Conclusions: The origin of the participants might have reduced the differences between MD adherence and DAS28. In parallel, the PG-SGA does not appear sensitive in detecting muscle-related malnutrition among patients with RA.

INTRODUCTION

Rheumatoid arthritis (RA) is a chronic condition, with patients receiving complex and diverse medications, often associated with adverse events.1-3 As a result, research interest on possible lifestyle changes with fewer adverse events has mushroomed recently, with a plethora of diet regimes and oral nutrient supplements (ONS) being investigated.4-7 In particular, studies using specific diets, such as the Mediterranean dietary pattern, have shown positive findings in tampering down pain and reducing the number of swollen joints.6,8 Individual components of the Mediterranean diet (MD) including poly-unsaturated...
fatty acids (PUFA), when provided as ONS, have been shown to reduce inflammation and the progression to pharmacotherapy. In parallel, according to a recent meta-analysis, between 1994 and 2016, was conducted using MEDLINE (via PubMed) to 32% of patients with RA experience rheumatoid cachexia, as a result of joint destruction, subsequent muscle inactivity, the high levels of sarcoactive inflammatory cytokines – including the tumour necrosis factor α (TNF-α) and interleukin (IL) 1β, loss of muscle mass and strength and concomitant increase in fat mass, is very common in patients with rheumatoid arthritis (RA) and due to the systemic inflammatory process experienced in RA. In addition, factors used commonly in RA to tamper down inflammation, including TNF inhibitors and the IL 6 receptor blocker, further aggregate body protein and energy metabolism, inducing body composition alterations. Although important, relevant data from Greece are limited.

The aim of the present pilot study was to assess differences in the nutritional status, MD adherence and functional ability among patients with RA, according to disease activity.

**METHODOLOGY**

Participants and research question

RA participants were recruited from consecutive outpatients visiting the Rheumatology clinic situated at the KAT General hospital in Athens, Greece, during 2014 (from April, to December of the same year). Patient characteristics are presented in Table 1. Eligible patients: 1) were adults (exceeding 18 years of age) on the basis of a definite disease (RA) irrespectively of whether their serological status (rheumatoid factor and anti-cyclic citrullinated peptide antibodies) was known or not to the treating physicians, 2) could communicate efficiently in the Greek language and 3) and provided participation consent. Permission for the study was granted by the Hellenic Mediterranean University and the KAT General Hospital. All patients provided informed consent prior to their participation. The participating institutions (Hellenic Mediterranean University and KAT General Hospital) have obtained an Ethics Committee approval.

Research question and design

The research question of this study in the format of P.E.O. (Population, Exposure and Outcomes) was addressed as follows: In patients suffering from RA (P), do different disease activity levels (E) associate with differences in their nutritional or functional status (O)? The study had a cross-sectional research design, with all measurements of exposure and outcome variables been recorded once for each patient (either measured by the authors of the study, or extracted from the patients’ medical records).

**Anthropometry**

Body weight and height of participants were recorded with a Seca 711 mechanical scale (Seca, Hamburg, Germany) and a Harpenden wall-mounted stadiometer, respectively (Holttain Limited, Crosswell, Crymlyn, Pembs, UK). For each patient, the body mass index (BMI) was calculated as body mass (kg) divided by stature (m²). Weight status was defined according to the World Health Organization BMI cutoffs for normoweight (18.5 < BMI ≤ 24.99 kg/m²), overweight (25 < BMI ≤ 29.99 kg/m²) and obesity (≥ 30 kg/m²).

**Disease activity and functional status**

The Disease Activity Score using 28 joint counts (DAS28) was used based on the erythrocyte sedimentation rate (ESR) of participants was selected. It is based on the swollen joint counts (SJCs) and the tender joint counts (TJCts). Scores <2.6 represent remission, low (≤3.2), moderate (≤5.1) or severe (>5.1) disease activity, respectively.

C-reactive protein (CRP), albumin and haemoglobin levels were recorded for each patient from their medical records. Disease duration was also recorded from medical records and was timed since the diagnosis of each patient (expressed in months).

Functional status/disability of participants was evaluated using a validated translation of the Stanford Health Assessment Questionnaire (HAQ) in the Greek language. The HAQ offers a patient-oriented outcome assessment of eight functional domains based on 20 specific functions, evaluating patient difficulty in performing daily activities during a recall period of one week.

**Malnutrition and diet quality**

Malnutrition was assessed with the Patient-Generated Subjective Global Assessment (PG-SGA). The PG-SGA provides a numerical score rating nourishment of patients. Higher scores are indicative of increased malnutrition risk whereas, a score ≥ 9 indicates an urgent need for nutrition intervention. In particular, the scores allow for triaging nutrition interventions, including symptoms management, the need for nutrition education for both the patient and the family, as well as specific dietary manipulations including (1) the need for enteral or parenteral nutrition, (2) the provision of additional or larger food portions to meet energy requirements, or (3) a need for oral nutrition supplements (ONS) to meet micronutrient needs.

The Mediterranean diet adherence was assessed via the MedDietScore score for each patient as previously proposed. The MedDietScore assesses the frequency of consumption of eleven distinct foods/food groups, including fish, non-refined cereals, fruits, vegetables, olive...
MEDITERRANEAN JOURNAL OF RHEUMATOLOGY

31
4
2020

Table 1. Description of study variables and tests for difference according to sex.

| Characteristic          | Total sample (n=48) | Men (n=8) | Women (n=40) | P-value |
|-------------------------|--------------------|-----------|--------------|---------|
| Age (years)             | 60.29 ± 12.94      | 60.13 ± 12.82 | 60.33 ± 13.13 | 0.969[^6] |
| BMI (kg/m^2)            | 25.67 (23.28–30.13) | 26.06 (23.95–28.96) | 25.31 (22.72–30.28) | 0.611[^7] |
| BMI categories          |                    |           |              |         |
| Normoweight             | 22 (45.83%)        | 3 (37.50%) | 19 (47.50%)  | 0.884[^8] |
| Overweight              | 14 (29.17%)        | 3 (37.50%) | 11 (27.50%)  |         |
| Obese                   | 12 (25.00%)        | 2 (25.00%) | 10 (25.00%)  |         |
| Disease duration (months) | 63.00 (33.00–120.00) | 56.00 (41.50–109.5) | 73.50 (28.50–120.00) | 0.897[^9] |
| MedDietScore            | 30.00 (27.00–31.50) | 29.00 (28.00–30.50) | 29.55±3.84 | 0.377[^10] |
| HAQ                     | 0.54 (0.11–0.95)   | 0.09 (0.01–0.40) | 0.65±0.44 | 0.037[^11] |
| PG-SGA score            | 0.50 (0.50–2.00)   | 0.50 (0.00–1.25) | 0.50 (0.50–2.00) | 0.193[^12] |
| PG-SGA categories       |                    |           |              |         |
| <2                      | 32 (66.67%)        | 6 (75.00%) | 26 (65.00%)  | 1.000[^13] |
| 2 to <4                 | 11 (22.92%)        | 2 (25.00%) | 9 (22.50%)   |         |
| 4 to <9                 | 4 (8.33%)          | 0 (0.00%) | 4 (10.00%)   |         |
| ≥9                      | 1 (2.08%)          | 0 (0.00%) | 1 (2.50%)    |         |
| DAS28 score             | 4.42 (2.74–6.24)   | 4.08 ± 1.74 | 4.55 ± 2.08 | 0.553[^14] |
| CRP (mg/dL)             | 0.80 (0.36–3.57)^[1] | 2.58 ± 2.40^1 | 0.80 (0.38–3.30) | 0.855[^15] |
| Albumin (g/dL)          | 3.90 (3.65–4.00)^[2] | 3.96 ± 0.59^1 | 3.90 (3.65–4.00)^[3] | 0.752[^16] |
| ESR (mm/h)              | 36.50 (15.50–55.00) | 44.25 ± 23.66 | 34.50 (15.00–55.00) | 0.466[^17] |
| Haemoglobin (g/dL)      | 12.39 ± 1.55       | 13.02 ± 2.07 | 12.26 ± 1.43 | 0.211[^18] |

BMI: body mass index; CRP: C-reactive protein; DAS28: Disease Activity Score-28; ESR: erythrocyte sedimentation rate; HAQ: Health Assessment Questionnaire; MedDietScore: Mediterranean diet score; PG-SGA: Patient-Generated Subjective Global Assessment.

Values are means ± standard deviations or medians (with 25th – 75th percentile) for continuous variables or frequencies (with percentages) for categorical variables.

[^6] refers to independent t-tests;[^7] refers to Mann-Whitney tests;[^8] refers to Chi-square tests;[1] 1 missing value;[^2] 5 missing values;[^3] 4 missing values.

Statistical analysis

Data are reported as means along with their standard deviations (SDs) for normally distributed variables, or as medians alongside their corresponding 25th and 75th percentiles when non-normally distributed variables were concerned. Deviations from the normality assumption were assessed with Shapiro-Wilk’s test. Independent t-tests assessed differences in normally distributed variables and the assumption of homoscedasticity was tested with Levene’s test. Categorical variables are presented as frequencies with their corresponding percentages and were assessed with the Chi-square test. Mann-Whitney tests assessed differences in non-normally distributed samples. For comparing nutritional (BMI, MedDietScore, and PG-SGA) and health assessment (HAQ) among the DAS28 categories, the Kruskal-Wallis tests were performed due to lack of normality on ANOVA’s (Analysis of Variance) residuals and/or lack of homogeneity of variances. Post-hoc comparisons were performed with the Bonferroni correction. Correlations among DAS28 as a continuous variable and BMI, MedDietScore, PG-SGA, and HAQ were assessed with Spearman’s rho correlation coefficient. SPSS version 25 (SPSS, Chicago, IL, USA) was used for the analyses and the level of significance was set at 0.05.

oil, alcohol, poultry, red meat, full-fat dairy, potatoes and legumes. The score ranges between 0 to 55, with greater scores being indicative of greater adherence to the Mediterranean diet.
The sample consisted of 48 RA patients with a mean age of 60.3 ± 12.9 years old. Most subjects were normoweight (45.8%) with a median BMI of 25.7 kg/m² and a median MedDietScore of 30 (Table 1). No differences were noted between men and women, except for an increased HAQ score for the latter (p=0.037).

**Associations of DAS28 with nutritional and general health variables**

DAS28 categories differed on HAQ score (p=0.027) (Table 2). Specifically, in post-hoc comparisons, only participants in remission (DAS28 <2.6) compared to participants with moderate DAS28 scores (2.6 to 3.2) had lower HAQ score (p=0.015). Additionally, DAS28 (as a continuous variable) demonstrated a week, positive correlation with HAQ (rho=0.296, p=0.041). Although the Kruskal-Wallis test indicated the presence of at least one difference among the DAS28 categories on MedDietScore (p=0.037) (Table 2), all post-hoc tests remained non-significant (all p>0.05) indicating that no specific DAS28 categories were significantly different from one another. Furthermore, DAS28 (as a continuous variable) did not demonstrate a correlation to the MedDietScore (rho=0.016, p=0.916). No further significant differences or correlations were noted for BMI or PG-SGA.

**Discussion**

The present pilot study indicates the existence of a relationship between disease activity and disability among patients with RA. In parallel, disease activity was associated with various levels of MD adherence. Malnutrition assessment failed to associate with disease activity. Although small in sample size, the current findings are in agreement to previous research indicating a relationship between HAQ and DAS2823-25. Boyd23 suggested that this tight connection is stronger at diagnosis (first visit), with a tendency to weaken as time passes after the diagnosis. On the other hand, age of participants did not appear to induce changes in this association,23 although it has been suggested that in older patients, age-related factors might act synergistically in tampering down functional status further. Previous research has showed that increased disease activity, older age, and female sex consist of HAQ predictors, irrespectively of the DAS28 score.26 In the present analysis, disease duration was long, whereas on the other hand, the majority of participants consisted of women of older age, and these factors might have contributed to weakening the DAS28-HAQ association. Nevertheless, the existence of a connection was noted, indicating that elevated increased RA activity and joint destruction was associated with deteriorated functional capacity. According to Drossaers-Bakker,25 disease activity is in fact the most pivotal determinant of the HAQ score and both should be evaluated at every patient visit to the record course of the disease.

**Table 2. Differences according to the DAS28 categories.**

| Variable          | Remission [DAS28 < 2.6] (n=11) | Low [DAS28 ≤ 3.2] (n=5) | Moderate [DAS28 ≤ 5.1] (n=12) | High [DAS28 >5.1] (n=20) | P-Value |
|-------------------|-------------------------------|------------------------|-----------------------------|-------------------------|---------|
| BMI (kg/m²)       | 28.57 ± 7.04                  | 24.98 (23.88–25.87)    | 24.34 (22.40–28.89)         | 26.40 ± 3.91           | 0.853   |
| MedDietScore      | 27.82 ± 4.21                  | 27.00 (27.00–29.00)    | 31.83 ± 2.72                | 29.3 ± 3.37            | 0.037   |
| HAQ score         | 0.15 (0.04–0.43)               | 0.58 ± 0.64            | 0.86 ± 0.32                 | 0.61 ± 0.49            | 0.027   |
| PG-SGA            | 0.50 (0.50–0.50)               | 1.30 ± 1.20            | 0.50 (0.00–0.75)            | 1.50 (0.50–3.00)       | 0.166   |

BMI: body mass index; DAS28: Disease Activity Score-28; HAQ: Health Assessment Questionnaire; MedDietScore: Mediterranean diet score; PG-SGA: Patient-Generated Subjective Global Assessment.

Values are means ± standard deviations or medians (with 25th – 75th percentile).

**Results**

The sample was consisted of 48 RA patients with a mean age of 60.3 ± 12.9 years old. Most subjects were normoweight (45.8%) with a median BMI of 25.7 kg/m² and a median MedDietScore of 30 (Table 1). No differences were noted between men and women, except for an increased HAQ score for the latter (p=0.037).

DAS28 categories differed on HAQ score (p=0.027) (Table 2). Specifically, in post-hoc comparisons, only participants in remission (DAS28 <2.6) compared to participants with moderate DAS28 scores (2.6 to 3.2) had lower HAQ score (p=0.015). Additionally, DAS28 (as a continuous variable) demonstrated a week, positive correlation with HAQ (rho=0.296, p=0.041). Although the Kruskal-Wallis test indicated the presence of at least one difference among the DAS28 categories on MedDietScore (p=0.037) (Table 2), all post-hoc tests remained non-significant (all p>0.05) indicating that no specific DAS28 categories were significantly different from one another. Furthermore, DAS28 (as a continuous variable) did not demonstrate a correlation to the MedDietScore (rho=0.016, p=0.916). No further significant differences or correlations were noted for BMI or PG-SGA.
MedDietScore. Moreover, given that the sample derived from a Mediterranean country where the MD consists of the main diet prototype adhered, it is possible that most of the participants demonstrated an adequate degree of MD adherence limiting between-group differences. No differences were noted in the malnutrition score in distinct DAS28 categories. Although nutrition appears to be an important component of RA initiation and progression, BMI is not sensitive enough to detect malnutrition with none of the participants herein being underweight according to their BMI. On the other hand, muscle protein degradation has been shown to occur in RA and hand grip strength, skinfold or body composition measurements might have produced more accurate results as compared to the PG-SGA. Rheumatoid cachexia has been suggested to correlate with disease activity and disability, further complicating outcomes and health status of patients. The PG-SGA used herein did not appear sensitive enough to depict malnutrition, as the only anthropometric index used to calculate the PG-SGA is body weight. Furthermore, all participants used in the present study were outpatients and it is likely that the inclusion of in-ward patients might have altered the malnutrition level of the sample. Nevertheless, according to the PG-SGA, no specific triage recommendations were required for the participants. Limitations of the present study include its small sample and cross-sectional design for demonstrating a causal relationship and further limit the generalizability of the findings. In parallel, if the DAS28-CRP was used instead of the DAS28-ESR, results might have differed. Finally, the small sample size and the lack of funding prohibited additional analyses adjusted for potential confounders such as socio-demographic characteristics, occupation, comorbidities and lifestyle parameters. The results indicate that disease activity and functional disability are aligned in RA. Given the design of the study, it is difficult to understand the role of the MD, however, most patients appear to adhere well to the MD prototype, either due to origin, or health consciousness. Further studies are needed to understand the disability and nutritional status of RA patients in Greece, as data are limited.

CONFLICT OF INTEREST
The authors declare no conflict of interest.

REFERENCES
1. Luis M, Freitas J, Costa F, Buttgereit F, Boers M, Da Silva JAP, et al. An updated review of glucocorticoid-related adverse events in patients with rheumatoid arthritis. Expert Opin Drug Saf 2019;18(7):581-90.
2. Dragos D, Glica M, Gamar L, Vlad A, Iosif L, Stoian I, et al. Phyto-medicine in Joint Disorders. Nutrients 2017;9(1):70.
3. de Camargo MC, Barros BCA, Fulone I, Silva MT, Silveira MSDN, de Camargo IA, et al. Adverse Events in Patients With Rheumatoid Arthritis and Psoriatic Arthritis Receiving Long-Term Biological Agents in a Real-Life Setting. Front Pharmacol 2019 Sep 11;10:965.
4. Asteriou E, Gkoutzourelas A, Mavropoulos A, Katsiaris C, Sakkas L, Bogdanos D. Curcumin for the Management of Periodontitis and Early ACPA-Positive Rheumatoid Arthritis: Killing Two Birds with One Stone. Nutrients 2018;10(7):908.
5. Vranou P, Gkoutzourelas A, Aftaniatou D, Zafirou I, Grammatikopoulou MG, Bogdanos DP. Let food be thy medicine: the case of the Mediterranean diet in rheumatoid arthritis. Mediterr J Rheumatol 2020;31(3):325-9.
6. Petersson S, Philippou E, Rodomar C, Nikiforou E. The Mediterranean diet, fish oil supplements and Rheumatoid arthritis outcomes: evidence from clinical trials. Autoimmun Rev 2018;17(11):1105-14.
7. Tedeschi SK, Costenbader KH. Is There a Role for Diet in the Therapy of Rheumatoid Arthritis? Curr Rheumatol Rep 2016;18(5):23.
8. Forsyth C, Kouvari M, D’Cunha NM, Georgoussopoulou EN, Panagiotakos DB, Mellor DD, et al. The effects of the Mediterranean diet on rheumatoid arthritis prevention and treatment: a systematic review of human prospective studies. Rheumatol Int 2018;38(5):737-47.
9. Santo RCE, Fernandes KZ, Lora PS, Filipin LJ, Xavier RM. Prevalence of rheumatoid cachexia in rheumatoid arthritis: a systematic review and meta-analysis. J Cachexia Sarcopenia Muscle 2018;9(5):816-25.
10. Fukuda W, Omoto A, Oku S, Tanaka T, Tsubouchi Y, Kohno M, et al. Contribution of rheumatoid arthritis disease activity and disability to rheumatoid cachexia. Mod Rheumatol 2010;20(5):439-43.
11. Roubenoff R. Rheumatoid cachexia: A complication of rheumatoid arthritis moves into the 21st century. Arthritis Res Ther 2009;11(2):108.
12. Engvall IL, Elkan AC, Tengstrand B, Cederholm T, Brismar K. Fat-storm I. Cachexia in rheumatoid arthritis is associated with inflammatory activity, physical disability, and low bioavailable insulin-like growth factor. Scand J Rheumatol 2006;37(5):321-28.
13. Tournadre A, Pereira B, Dutthel F, Giraud C, Courtieux D, Sapin V, et al. Changes in body composition and metabolic profile during interleukin 6 inhibition in rheumatoid arthritis. J Cachexia Sarcopenia Muscle 2017;8(4):639-46.
14. WHO Expert Consultation. Waist Circumference and Waist-Hip Ratio: Report of a WHO Expert Consultation. Geneva, Switzerland: World Health Organization; 2008.
15. Fuchs HA, Brooks RH, Callahan LF, Fincus T. A simplified twenty-eight–joint quantitative articular index in rheumatoid arthritis. Arthritis Rheum 1989;32(5):531-7.
16. van Gestel AM, Haagsma CJ, van Riel PL. Validation of the Disease Activity Score (DAS) and Health Assessment Questionnaire Disability Index (HAQ), Multidimensional Health Assessment Questionnaire Disability Index (HAQ), Modified Health Assessment Questionnaire Disability Index (MDHAQ), Health Assessment. Arthritis Care Res 2003;4(11):1845-50.
17. Hensor EMA, Emery P, Bingham SJ, Conaghan PG. Discrepancies in categorizing rheumatoid arthritis patients by DAS-28(ESR) and DAS-28(CRP): can they be reduced? Rheumatology 2010;49(8):1521-9.
18. Chatzitheodorou D, Kabitsis C, Papadopoulos NG, Galanopoulou V. Assessing disability in patients with rheumatic diseases: translation, reliability and validity testing of a Greek version of the Stanford Health Assessment Questionnaire (HAQ). Rheumatol Int 2006;28(11):1091-7.
19. Bruce B, Fries JF. The Stanford Health Assessment Questionnaire: dimensions and practical applications. Health Qual Life Outcomes 2003;1(1):20.
20. Masala L, Anderson J, Michaud K. Measures of functional status and quality of life in rheumatoid arthritis: Health Assessment Questionnaire Disability Index (HAQ), Modified Health Assessment Questionnaire (MHAQ), Multidimensional Health Assessment Questionnaire (MDHAQ), Health Assessment. Arthritis Care Res (Hoboken) 2011;63(S1):S4-S13.
21. Ottery FD. Patient-Generated Subjective Global Assessment. In: McCallum PD, Polisena CG, eds. The Clinical Guide to Oncology Nutrition. Chicago, IL: The American Dietetic Association; 2000:11-23.
22. Panagiotakos DB, Pitsavos C, Arvaniti F, Stefanadis C. Adherence to the Mediterranean food pattern predicts the prevalence of hypertension, hypercholesterolemia, diabetes and obesity, among healthy adults; the accuracy of the MedDietScore. Prev Med (Bal-tim) 2007;44(4):335-40.

23. Boyd TA, Bonner A, Thorne C, Boire G, Hitchon C, Harraoui BP, et al. The Relationship Between Function and Disease Activity as Measured by the HAQ and DAS28 Varies Over Time and by Rheumatoid Factor Status in Early Inflammatory Arthritis (EIA). Results From the CATCH Cohort. Open Rheumatol J 2013;7:58-63.

24. Kumar SB, Suneetha P, Mohan A, Kumar PD, Sarma KVS. Comparison of Disease Activity Score in 28 Joints With ESR (DAS28), Clinical Disease Activity Index (CDAI), Health Assessment Questionnaire Disability Index (HAQ-DI) & Routine Assessment of Patient Index Data With 3 Measures (RAPID3) for Assessing Dis. Indian J Med Res 2017;146(Supplement).

25. Dossaers-Bakker KW, De Buck M, Van Zebed M, Zwinderman AH, Breedveld FC, Hazes JMW. Long-term course and outcome of functional capacity in rheumatoid arthritis: The effect of disease activity and radiologic damage over time. Arthritis Rheum 1990;42(9):1854-60.

26. Welting PM, van Gestel AM, Swinkels HL, Kliemeney LA, van Riel PL. The Relationship Between Disease Activity, Joint Destruction, and Functional Capacity Over the Course of Rheumatoid Arthritis. Arthritis Rheum 2001;44(9):2009-17.

27. Cutolo M, Nikiphorou E. Don’t neglect nutrition in rheumatoid arthritis! RMD Open 2018;4(1):e000591.

28. Elkan A-C, Engvall I-L, Tengstrand B, Cederholm T, Hafstrom I. Malnutrition in women with rheumatoid arthritis is not revealed by clinical anthropometrical measurements or nutritional evaluation tools. Eur J Clin Nutr 2008;62(10):1239-47.

29. Collins Jr R, Dunn T, Walthaw J, Harrell P, Alarcon G. Malnutrition in Rheumatoid Arthritis. Clin Rheumatol 1997;6(3):391-8.

30. Fukuda W, Yamazaki T, Akaogi T, Hayashi H, Kusakabe T, Tsubouchi Y, et al. Malnutrition and disease progression in patients with rheumatoid arthritis. Mod Rheumatol 2005;15(2):104-7.

31. Fleischmann RM, Van Der Heijde D, Gardiner PV, Szumski A, Marshall L, Banaris E. DAS28-CRP and DAS28-ESR cut-offs for high disease activity in rheumatoid arthritis are not interchangeable. RMD Open 2017 Jan 30;3(1):e000392.