

## 2017 EULAR recommendations for a core data set to support observational research and clinical care in rheumatoid arthritis

Radner, H; Raza, Karim

DOI:

[10.1136/annrheumdis-2017-212256](https://doi.org/10.1136/annrheumdis-2017-212256)

License:

None: All rights reserved

*Document Version*

Peer reviewed version

*Citation for published version (Harvard):*

Radner, H & Raza, K 2018, '2017 EULAR recommendations for a core data set to support observational research and clinical care in rheumatoid arthritis', *Annals of the Rheumatic Diseases*, vol. 77, pp. 476-479. <https://doi.org/10.1136/annrheumdis-2017-212256>

[Link to publication on Research at Birmingham portal](#)

### **Publisher Rights Statement:**

Final Version of Record published as: Radner, Helga, et al. "2017 EULAR recommendations for a core data set to support observational research and clinical care in rheumatoid arthritis." *Annals of the rheumatic diseases* (2018). Available at: <http://dx.doi.org/10.1136/annrheumdis-2017-212256>

### **General rights**

Unless a licence is specified above, all rights (including copyright and moral rights) in this document are retained by the authors and/or the copyright holders. The express permission of the copyright holder must be obtained for any use of this material other than for purposes permitted by law.

- Users may freely distribute the URL that is used to identify this publication.
- Users may download and/or print one copy of the publication from the University of Birmingham research portal for the purpose of private study or non-commercial research.
- User may use extracts from the document in line with the concept of 'fair dealing' under the Copyright, Designs and Patents Act 1988 (?)
- Users may not further distribute the material nor use it for the purposes of commercial gain.

Where a licence is displayed above, please note the terms and conditions of the licence govern your use of this document.

When citing, please reference the published version.

### **Take down policy**

While the University of Birmingham exercises care and attention in making items available there are rare occasions when an item has been uploaded in error or has been deemed to be commercially or otherwise sensitive.

If you believe that this is the case for this document, please contact [UBIRA@lists.bham.ac.uk](mailto:UBIRA@lists.bham.ac.uk) providing details and we will remove access to the work immediately and investigate.

# THE 2017 EULAR RECOMMENDATIONS FOR A CORE DATASET TO SUPPORT OBSERVATIONAL RESEARCH AND CLINICAL CARE IN RHEUMATOID ARTHRITIS

Radner H<sup>1</sup>, Chatzidionysiou K<sup>2</sup>, Nikiphorou E<sup>3</sup>, Gossec L<sup>4</sup>, Hyrich K<sup>5</sup>, Zabalán C<sup>6</sup>, van Eijk-Hustings Y<sup>7</sup>, Williamson P.R<sup>8</sup>, Balanescu A<sup>9</sup>, Burmester G.R<sup>10</sup>, Carmona L<sup>11</sup>, Dougados M<sup>12</sup>, Finckh A<sup>13</sup>, Haugeberg G<sup>14</sup>, Hetland M.L<sup>15</sup>, Oliver S<sup>16</sup>, Porter D<sup>17</sup>, Raza K<sup>18</sup>, Ryan P<sup>19</sup>, Santos M.J<sup>20</sup>, van der Helm-van Mil A<sup>21</sup>, van Riel P<sup>22</sup>, von Krause G<sup>23</sup>, Zavada J<sup>24</sup>, Dixon WG<sup>25</sup>, Askling J<sup>2</sup>

<sup>1</sup> Department of Internal Medicine III, Division of Rheumatology; Medical University Vienna; Austria

<sup>2</sup> Department of Rheumatology, Karolinska University Hospital, and Department of Medicine Solna, Karolinska Institutet, Stockholm, Sweden

<sup>3</sup> Academic Rheumatology Department, King's College London, London, UK

<sup>4</sup> Sorbonne Universités, UPMC University Paris 06; AP-HP, Hopital Pitie-Salpetriere, Rheumatology Department, Paris, France

<sup>5</sup> NIHR Manchester Musculoskeletal Biomedical Research Unit, Central Manchester NHS Foundation Trust, Manchester Academic Health Science Centre

<sup>6</sup> Patient Partner of Romanian League against Rheumatism, Bucharest, Romania

<sup>7</sup> Department of Patient & Care, Maastricht, The Netherlands

<sup>8</sup> MRC North West Hub for Trials Methodology Research, Department of Biostatistics, University of Liverpool, Waterhouse Building, Brownlow Street, Liverpool, L69 3GL, UK

<sup>9</sup> Research Center of Rheumatic Diseases, "Sf. Maria" Hospital, University of Medicine and Pharmacy "Carol Davila", Bucharest, Romania

<sup>10</sup> Department of Rheumatology and Clinical Immunology, Charité—University Medicine Berlin, Berlin, Germany

<sup>11</sup> Instituto de Salud Musculoesquelética, Madrid, Spain

<sup>12</sup> Medicine Faculty, APHP, Rheumatology B Department, Paris Descartes University, Cochin Hospital, Paris, France

<sup>13</sup> Division of Rheumatology, Department of Internal Medicine Specialties, University Hospitals of Geneva, 26, Ave Beau-Sejour, CH 1211, Geneva, Switzerland.

<sup>14</sup> Department of Rheumatology, Martina Hansens Hospital, Bærum; Norwegian University of Science and Technology, Trondheim, Norway

<sup>15</sup> Copenhagen Center for Arthritis Research, Center for Rheumatology and Spine Diseases, Centre of Head and Orthopaedics, Rigshospitalet, Glostrup, and Department of Clinical Medicine, Faculty of Health and Medical Sciences, University of Copenhagen, Copenhagen, Denmark

<sup>16</sup> North Devon, UK

<sup>17</sup> Rheumatology Department, Gartnavel General Hospital, Glasgow, UK

<sup>18</sup> Institute of Inflammation and Ageing, Arthritis Research UK Rheumatoid Arthritis Pathogenesis Centre of Excellence and MRC Arthritis Research UK Centre for Musculoskeletal Ageing Research, University of Birmingham, Birmingham, UK; NIHR Birmingham Biomedical Research Centre, Birmingham UK; and Sandwell and West Birmingham Hospitals NHS Trust, Birmingham, UK

<sup>19</sup> Department of Biomedical Informatics, Columbia University, New York, NY 10032, United States; Janssen Research and Development, Titusville, NJ 08560, United States

<sup>20</sup> Department of Rheumatology, Hospital Garcia de Orta and Instituto de Medicina Molecular da Faculdade de Medicina, Universidade de Lisboa, Portugal

<sup>21</sup> Department of Rheumatology, Leiden University Medical Center, Leiden; Department of Rheumatology, Erasmus Medical Center, Rotterdam, the Netherlands

<sup>22</sup> Department of Rheumatology, Bernhoven, Uden, The Netherlands

<sup>23</sup> Patient research partner, Paris, France

<sup>24</sup> Institute of Rheumatology, Prague, and Department of Rheumatology, First Faculty of Medicine, Charles University, Na Slupi 4, 12850, Prague, Czech Republic

<sup>25</sup> Arthritis Research UK Centre for Epidemiology, Manchester Academic Health Sciences Centre, The University of Manchester, Manchester, UK

Word Count 1500

Abstract 191

## ABSTRACT

**Background.** Personalised medicine, new discoveries, and studies on rare exposures or outcomes require large samples that are increasingly difficult for any single investigator to obtain. Collaborative work is limited by heterogeneities, both what is being collected, and how it is defined.

**Objective.** To develop a core set for data collection in rheumatoid arthritis (RA) research which (i) allows harmonization of data collection in future observational studies, (ii) acts as a common data model against which existing databases can be mapped, and (iii) serves as a template for standardized data collection in routine clinical practice to support generation of research-quality data.

**Methods.** A multi-step, international multi-stakeholder consensus process was carried out involving voting via online surveys and two face-to-face meetings.

**Results.** A core set of 21 items (“what to collect”) and their instruments (“how to collect”) was agreed: Age, gender, disease duration, diagnosis of RA, BMI, smoking, swollen/tender joints, patient/evaluator global, pain, quality of life, function, composite scores, acute phase reactants, serology, structural damage, treatment and comorbidities.

**Conclusions.** The core set should facilitate collaborative research, allow for comparisons across studies, and harmonise future data from clinical practice via electronic medical record systems.

## INTRODUCTION

Research questions of the current era require ever larger study populations which often exceed the number of patients available in individual studies or registries, hence combined or pooled analyses are often required. Such collaborative work is, however, compromised by heterogeneities in the data collected. (1) More recently, routinely collected data from electronic medical records (EMR) has the potential to support research, but there is no agreed set of data that would be desirable to collect in order to generate research-quality data if used beyond direct clinical care.

To facilitate collaborative research, standardizing items (“what to collect”) and their instruments (“how to collect”) across studies and data collections is critical. Although guidelines on data reporting and outcome measures have been developed, such core sets have so far pertained to randomized control trials and trial extension studies in rheumatoid arthritis (RA) (2, 3), and have not been vetted against the feasibility of data collection in routine clinical care, nor against the needs of observational research. Indeed, any core dataset to be implemented in registers, research cohorts or EMR systems need to strike a balance between clinical feasibility and the potential to generate useful, research-quality data.

For these reasons, a EULAR task force was convened to develop a standardised core set for data collection in RA that should (i) harmonize data collection in future observational studies, (ii) act as a common data model against which existing databases and EMR systems can be mapped, and (iii) ensure that new data collections in routine clinical practice can support the generation of future research-quality data.

## METHODS

The task force comprised a *steering committee* (n=10, including 2 co-conveners (WD, JA), a EULAR methodologist (LG), a patient partner (CZ) and 3 fellows (HR, KC, EN)), a *working group* (additional 15 experts from 11 European countries and the U.S. including patients, clinical researchers and experts in the area of epidemiology, quality of care, and common data modelling, physicians and other health professionals), and a pan-European *expert panel* (n=90).

The task force employed a step-wise process (figure 1) in accordance with the EULAR standard operating procedure (4), underpinned by an inventory of data collection in existing RA clinical cohorts and registers (1):

(a) An hierarchical literature review was carried out by the fellows in PubMed to identify data items and instruments used in existing RA registers and clinical cohorts, and the frequency with which these items and instruments were collected. Results were cross-checked with EULAR recommendations for reporting in RA clinical trials and trial extension studies (2, 3) and with the EULAR outcome measures library. Items and instruments felt to be missing could be added by the steering committee.

(b) In an online survey, the expert panel rated the perceived importance of each of the above items (and instruments) to be included in a core set, using a grading scale from 1 (not important) to 9 (very important). Again, items and instruments could be added.

(c) At the first face-to-face meeting of the working group, electronic voting on individual items and instruments to include in the core set took place. We defined consensus as  $\geq 70\%$  of participants voting either YES or NO to the inclusion of a given item; if no consensus was reached, a second vote took place after a brief discussion which was led by an independent moderator (DP). In this second round, the threshold for consensus was lowered to 60%. The resultant list of items that had reached consensus for inclusion informed a single round of voting on the instruments with which to collect information on the items.

(d) Additional information on items and instruments for which no consensus was reached was collected. Ratification of items included, reasons for any exclusion, and voting on items that had not reached consensus at the face-to-face meeting was performed via an online survey within the working group.

(e) At the second face-to-face meeting of the working group, voting was conducted for outstanding items, then for final approval of the complete core list of items, followed by voting on instruments for all items in the final list using the same method of moderated discussion and voting cut-offs as outlined above.

In all voting processes, the panellists were reminded of the importance of striking a balance between clinical feasibility and the potential to generate research-quality data.

## RESULTS

### *(a) Literature review*

Published articles from 67 different European RA registers and clinical cohorts were included. In total, 39 items and 125 instruments were identified; 8 additional items for consideration were added (supplementary table 1 and 2).

### *(b) Online survey*

90 experts from 28 different European countries including patients (18%), allied health professionals (18%), physicians (55%) and researchers (10%) participated in the survey. 29 of the 47 proposed items were considered important enough for inclusion into a core set (supplementary table 2).

### *(c) First face-to-face meeting*

21/25 (84%) members attended. In the first voting round consensus for inclusion was reached for 16/47 (34%) items, and consensus for exclusion for 16/47 (34%) items. After discussion and a second round of voting on the remaining 15 items, consensus was reached for inclusion of 5 and for exclusion of 7, leaving two items (“health related quality of life”, and “socioeconomic status”) without consensus (supplementary table 2).

### *(d) Online ratification*

23/25 (92%) members participated. The voting results from the face-to-face meeting were confirmed except for socioeconomic status, health-related quality of life, fatigue, and joint surgery (lack of consensus).

### *(e) Second face-to-face meeting*

17/25 (68%) members attended. Voting led to the exclusion of items “fatigue”, “joint surgery” and “socioeconomic status” but inclusion of “health-related quality of life”. Next, the

resultant set of the remaining 21 items was voted on in its entirety, and 100% of the participants agreed (Table 1). Moderated discussion and sequential voting on the instruments with which to collect these 21 items resulted in consensus for all but two instruments (how to collect data for items “glucocorticoids”, and “comorbidity”).

*(f) Second online survey*

A final ad hoc online survey with pre-specified suggestions provided by the steering committee for the outstanding instruments for items ‘glucocorticoids’ and ‘comorbidity’ was performed (supplementary table 3). 21/25 (84%) members participated, and consensus was reached. The final core set thus contained 21 items, each with its preferred instrument (see BOX).

**BOX.** Structure and content of the RA core set developed by the EULAR Task Force

ITEM	RECOMMENDATION OF STANDARDIZED WAY OF ASSESSMENT
1) Age	date of birth
2) Gender	male / female
3) Disease duration	date of diagnosis
4) Diagnosis of RA	Rheumatologist reported diagnosis of RA
5) Body mass index (BMI)	weight and height
6) Smoking	current / previous / never
7) Tender joints	28 joint count
8) Swollen joints	28 joint count
9) Patient global	measured on VAS or NRS scale capturing: 1) global assessment of disease activity; 2) related to arthritis; 3) today Example wording: “Considering all the ways your arthritis has affected you, how do you feel your arthritis is today?” anchors: excellent - very poor
10) Evaluator global	measured on VAS or NRS scale capturing: 1) global assessment of disease activity; 2) related to arthritis; 3) today Example wording: “What is your overall assessment of the patient's RA disease activity today?” anchors: excellent - very poor
11) Pain	measured on VAS or NRS capturing: 1) pain; 2) related to arthritis; 3) last week Example wording: “How much pain did you have due to your arthritis last week?” anchors: no pain – worst imaginable pain
12) Physical function	Health Assessment Questionnaire (HAQ)
13) Health related quality of life	Euro-Qol 5 dimensions (EQ5D)*
14) Composite Scores	Collection of Core Items 7-12 and 15 enables to calculate following composite scores: - CDAI(Clinical disease activity index) - SDAI (Simplified disease activity index) - DAS28 (Disease activity Score 28 joints) - EULAR response criteria - ACR response criteria
15) Acute phase reactants	C-reactive protein (CRP) AND erythrocyte sedimentation rate (ESR)
16) Serology	Rheumatoid factor (RF) AND anti-CCP antibodies (ACPA)
17) Structural Damage	Presence of erosions on X-ray Yes/No
18) DMARD History	name of previous DMARD(s) (including biological, synthetic and targeted synthetic DMARDs)

19) Ongoing / most recent DMARD	name of DMARD
	start and stop date
	reason for stopping
20) Glucocorticoids	current use of oral glucocorticoids: 1) dose taken today ___mg prednisolone equivalent; and 2) continuous (> 3 months) intake YES/NO
21) Comorbidities	binary assessment YES/NO of the six EULAR comorbidity domains (cardiovascular disease, malignancies, infections, gastrointestinal disease, osteoporosis, depression) (5)

\* free licence available after registration for non-commercial parties according to [www.euroqol.org](http://www.euroqol.org)

## DISCUSSION

This EULAR taskforce has defined an RA core set, including both items and instruments to support standardised RA data collection in clinical practice and research. This will enable collaborative research studies, and increase comparability across studies.

Unlike most previous core sets, this set was specifically developed keeping clinical feasibility in mind. Importantly, “core” underscores that the set represents a minimum, acknowledging that individual stakeholders are likely to add items or instruments of particular interest to their own data collection.

Consensus for inclusion was straightforward for the majority of items: 16 of the final 21 items were agreed in the first round of voting. Six additional items were included, and after a ratification survey and discussion, ‘joint surgery’ was excluded in the subsequent process. Discussion for each item is summarized in supplementary table 3. Of the 26 items ultimately excluded from the core dataset 9 were seen as important by the expert group. One item, ‘fatigue’, was excluded from our core dataset yet is present in the OMERACT core outcome set for RA trials.(6) Discussion during the meeting acknowledged that this item was important but depending on many factors unrelated to RA and can be captured by PGA. The group repeatedly revisited the scope of a core set, and agreed that fatigue was a good example of an item that might regularly be collected in addition to the core set for either clinical practice or research.

The task-force was established to include expertise in registers and observational research, clinical practice, use of EMRs, the patient perspective, as well as methodological expertise in the development of core datasets (7) to enable robustness and transparency. To ensure inclusiveness, all national European rheumatology societies were invited to contribute. Dissemination of the core set will be facilitated by publication online (EULAR website) and via national societies.

Following agreement on this core set, it is important that it is adopted by future observational data collections and research studies in RA. For EMR systems to deliver to RA research, the core set also needs to be integrated into EMR systems. Additional clinical benefits of collection of structured data such as informing consultations, viewing longitudinal disease progression, supporting audit, and benchmarking care quality (8) should be recognised and may encourage its clinical adoption.



In summary, through a multi-step, multi-stakeholder, and evidence-based process, this task force has developed an RA core set that may (i) harmonize data collection in future observational studies, (ii) act as a common data model against which existing databases can be mapped, and (iii) serve as a template for standardized data collection in routine clinical practice to support the generation of research-quality data. This effort, which will continue to be updated and revised after evaluation of its implementation, may serve as an example for other conditions beyond RA.

## **ACKNOWLEDGMENT:**

The authors want to thank all contributing experts who participated in the online survey.

## **FUNDING**

This project was funded by a EULAR project grant

## **REFERENCES**

1. Radner H, Dixon W, Hyrich K, et al. Consistency and Utility of Data Items Across European Rheumatoid Arthritis Clinical Cohorts and Registers. *Arthritis Care Res (Hoboken)*. 2015;67(9):1219-29.
2. Aletaha D, Landewe R, Karonitsch T, et al. Reporting disease activity in clinical trials of patients with rheumatoid arthritis: EULAR/ACR collaborative recommendations. *Ann Rheum Dis*. 2008;67(10):1360-4.
3. Buch MH, Silva-Fernandez L, Carmona L, et al. Development of EULAR recommendations for the reporting of clinical trial extension studies in rheumatology. *Ann Rheum Dis*. 2015;74(6):963-9.
4. van der Heijde D, Aletaha D, Carmona L, et al. 2014 Update of the EULAR standardised operating procedures for EULAR-endorsed recommendations. *Ann Rheum Dis*. 2015;74(1):8-13.
5. Baillet A, Gossec L, Carmona L, et al. Points to consider for reporting, screening for and preventing selected comorbidities in chronic inflammatory rheumatic diseases in daily practice: a EULAR initiative. *Ann Rheum Dis*. 2016;75(6):965-73.
6. Kirwan JR, Minnock P, Adebajo A, et al. Patient perspective: fatigue as a recommended patient centered outcome measure in rheumatoid arthritis. *J Rheumatol*. 2007;34(5):1174-7.
7. Kirkham JJ, Gorst S, Altman DG, et al. Core Outcome Set-STAndards for Reporting: The COS-STAR Statement. *PLoS Med*. 2016;13(10):e1002148.
8. Ibfelt EH, Jensen DV, Hetland ML. The Danish nationwide clinical register for patients with rheumatoid arthritis: DANBIO. *Clin Epidemiol*. 2016;8:737-42.
9. Nikiphorou E, Radner H, Chatzidionysiou K, Desthieux C, Zabalán C, van Eijk-Hustings Y, et al. Patient global assessment in measuring disease activity in rheumatoid arthritis: a review of the literature. *Arthritis Res Ther*. 2016;18(1):251.
10. Felson DT, Smolen JS, Wells G, Zhang B, van Tuyl LH, Funovits J, et al. American College of Rheumatology/European League against Rheumatism provisional definition of remission in rheumatoid arthritis for clinical trials. *Ann Rheum Dis*. 2011;70(3):404-13.

11. van der Heijde D, van der Helm-van Mil AH, Aletaha D, Bingham CO, Burmester GR, Dougados M, et al. EULAR definition of erosive disease in light of the 2010 ACR/EULAR rheumatoid arthritis classification criteria. *Ann Rheum Dis.* 2013;72(4):479-81.