

1 Deadline for submission of abstracts: 24 June 2014 (Noon Eastern Time)

2 <Title character count: 138> [limit: assume 250 characters]

3 <Character count: 2746 (includes 250 for table)> [limit: 2750 characters] (excludes spaces,
4 title, names of authors and affiliations and disclosures; table or figure counts towards
5 character count by ~250 characters, but this should be checked)

6

7 **Comparisons of Quality of Life, Resource Use and Physical Functioning in RA**
8 **Patients Classified as High, Moderate or Low Risk for Rapid Radiographic**
9 **Progression**

10

11 EA Alemao, Bristol-Myers Squibb, Princeton, NJ, United States

12 S Joo, Bristol-Myers Squibb, Hopewell, NJ, United States

13 P Allison, University of Pennsylvania, Philadelphia, PA, United States

14 M Al, Erasmus University, Rotterdam, Netherlands

15 M Rutten-van Molken, Erasmus University, Rotterdam, Netherlands

16 S Banerjee, Bristol-Myers Squibb, Princeton, United States

17 C Iannaccone, Brigham and Women's Hospital, Boston, MA, United States

18 M Frits, Brigham and Women's Hospital, Boston, MA, United States

19 N Shadick, Brigham and Women's Hospital, Boston, MA, United States

20 M Weinblatt, Brigham and Women's Hospital, Boston, MA, United States

21 KP Liao, Brigham and Women's Hospital, Boston, MA, United States

22

23 **Background/Purpose:** We developed and validated a prognostic model to identify subjects
24 with elevated risk of rapid radiographic progression (RRP). The objective of this study was to
25 compare differences in quality of life (QoL), resource use and clinical outcomes at 12 months
26 in patients classified with high, moderate and low baseline risk of RRP by the prognostic
27 model.

28 **Methods:** In a longitudinal cohort of RA patients with clinical and radiographic data in an
29 outpatient setting, we applied the prognostic model to calculate the baseline probability of
30 RRP. Variables to determine the probability of RRP in the prognostic model included
31 seropositivity, body weight, disease duration, DAS28 (CRP) and total Sharp score. Based on
32 the calculated probability of RRP, patients were categorized into low risk (probability 0 to
33 0.25), moderate risk (0.25 to 0.75) and high risk (>0.75) of RRP. The categorization was
34 based on visual inspection of probability plots. QoL outcome measured by EQ5D, healthcare
35 resource use (nursing home visits, home healthcare visits, surgeries, durable medical

36 equipment use, hospitalization and ER visits) and clinical outcome of physical functioning
37 measured by mHAQ at 12 months were compared by baseline RRP risk groups of low,
38 moderate and high using analysis of variance for continuous variables and Chi-square test
39 for categorical variables.

40 **Results:** In the RA cohort, 942 (72.6%) patients had adequate data to calculate RRP. Of
41 these, 414 (43.9%) were classified as low, 477 (50.6%) as medium and 51 (5.4%) as high
42 risk of RRP at baseline. Patients in the low-risk group when compared with those in the
43 moderate- and high-risk groups tended to be younger, have a lower number of swollen or
44 tender joints (mean [SD] 9.4 yrs [11.5], 19.8 [14.2], 33.1 [12.9], respectively), and less likely
45 to be treated with a biologic DMARD. Patients in the low- versus high-risk groups had higher
46 QoL, lower resource use and higher physical functioning at 12 months (Table).

Table: QoL, Resource Use and Physical Functioning at 12 Months in Patients at Low, Moderate and High Baseline Risk of RRP			
Outcomes	Low Risk of RRP	Moderate Risk of RRP	High Risk of RRP
EQ5D, mean (SD)**	0.83 (0.14)	0.79 (0.15)	0.72 (0.19)
ER visits, % of pts*	23.4	25.1	38.2
Nursing home visits, % of pts*	2.4	2.7	14.6
Home healthcare visits, % of pts*	4.8	13.5	36.0
Surgeries, % of pts*	15.4	25.4	38.2
DME use, % of pts*	21.0	33.2	58.4
Hospital visits, % of pts*	13.3	20.4	37.1
mHAQ, mean (SD)**	0.39 (0.42)	0.65 (0.50)	0.72 (0.19)

*p<0.05 based on Chi-square test; **p<0.05 based on analysis of variance

47 **Conclusion:** Patients categorized as having high risk of future RRP at baseline (compared
48 with moderate and low risk of RRP) had worse outcomes at 12 months for QoL, resource
49 utilization and physical functioning. These findings suggest that therapies are needed to
50 improve QoL and resource utilization in these high-risk patients.

**American College of Rheumatology (ACR) Annual Scientific Meeting;
14–19 November 2014; Boston, Massachusetts, United States**

APPENDIX

Key words: Cardiovascular disease, risk management, rheumatoid arthritis

Submission category: Health Services Research, Quality Measures and Quality of Care

Preferred presentation format: No preference

Additional Information

Research Method:

Type of Trial:

Type of Trial Phase:

Track: Clinical practice

Primary research method: Observational

Study sponsor statement: Bristol-Myers Squibb. The study sponsor provided funding for the completion of the study and the development of the abstract.

AUTHOR AGREEMENTS

For information for all authors:

Presenting Author Agreement

The ACR does not condone presentations given by an invited presenter who has not been intimately involved in the development of the data and who cannot meet the criteria for authorship. **Presenting authors will be required to check both statements to be eligible to present.**

- I affirm, I have had full access to all of the data in the study, and take responsibility for the integrity of the data and the accuracy of the data analysis and approved the data for presentation.
- I affirm, I have made significant contributions to the study design, analysis or interpretation of results.

Institutional Review Board Affirmation

An Institutional Review Board (IRB) is charged with protecting the rights and welfare of people involved in research. All Human Subjects Research must receive approval from the IRB. The purpose of the IRB is to protect the rights and welfare of individuals who are participating as subjects in the research.

- I affirm that my research meets received approval from the IRB.
- I affirm that my research did not involve human subjects and therefore no IRB approval was required.

- I accept these terms. By accepting these terms, I agree 1) to allow the College to use my presentation in connection with its education resources, including SessionSelect (a digital copy of my presentation audio and video as presented, and 2) to distribute a PDF copy of my presentation to attendees and users of *SessionSelect*.

Each abstract submission must abide by the following conditions:

**American College of Rheumatology (ACR) Annual Scientific Meeting;
14–19 November 2014; Boston, Massachusetts, United States**

- I affirm that I have read and agree to the ACR Annual Meeting general guidelines and policies for abstract submission outlined in the 2013 Call for Abstracts Brochure.
- I affirm that any work with human or animal subjects reported in the abstract complies with the guiding principles for experimental procedures found in the Declaration of Helsinki of the World Medical Association.
- I understand that case reports are not acceptable and will not be reviewed.
- I understand that if the abstract reports the results of a clinical trial not yet approved by a regulatory agency, the trial phase must be indicated on the submission form.
- I understand that an abstract is ineligible for consideration if it reports work that has been accepted for publication as a manuscript prior to the ACR submission deadline of Tuesday, June 25, 2013.
- I understand that abstracts submitted for the ARHP may not be dually submitted to the ACR and vice versa.
- I understand that, if accepted, the American College of Rheumatology has permission to publish this abstract in printed and/or electronic formats.
- I understand that, if accepted for presentation, the presenting author or co-authors listed on the abstract must present the abstract during an oral and/or poster presentation.

Abstract embargo policy

Accepted abstracts are made available to the public online in advance of the meeting and are published in a special supplement of [Arthritis & Rheumatism](#). Information contained in those abstracts may not be released until the abstracts appear online. Academic institutions, private organizations and companies with products whose value may be influenced by information contained in an abstract may issue a press release to coincide with the availability of an ACR abstract on the ACR website.

However, the ACR continues to require that information that goes beyond that contained in the abstract (e.g., discussion of the abstract done as part a scientific presentation or presentation of additional new information that will be available at the time of the meeting) is under embargo until 4:30 PM Pacific Time on Saturday, November 15, 2014. Violation of this policy may result in the abstract being withdrawn from the meeting and other measures deemed appropriate. Authors are responsible for notifying financial and other sponsors about this policy.

- I understand that this abstract, if accepted, will be under embargo until 4:30 PM Eastern Time on Saturday, November 15.