

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

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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)

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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**

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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).

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.

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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:

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- 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.

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- I understand that abstracts submitted for the ARHP may not be dually submitted to the ACR and vice versa.
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- 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.

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