

LETTER

Further evaluation of a claims-based algorithm to determine the effectiveness of biologics for rheumatoid arthritis using commercial claims data

Jeffrey R Curtis*¹, Benjamin Chastek², Laura Becker², David J Harrison³, David Collier³, Huifeng Yun⁴ and George J Joseph³

See related research by Curtis *et al.*, <http://arthritis-research.com/content/13/5/R155>, and related editorial by Kim and Solomon, <http://arthritis-research.com/content/13/5/129>

Background

As more biologics are approved, there is increasing interest in comparative effectiveness research (CER). Health insurance claims databases contain information about outpatient visits, hospital discharges, procedures, and outpatient pharmacy dispensing but seldom contain clinical outcomes. In a previous issue of *Arthritis Research & Therapy*, we presented an algorithm that assessed the clinical effectiveness of rheumatoid arthritis (RA) biologics which used Veterans Affairs (VA) claims data and which was validated against the DAS28-ESR (Disease Activity Score 28 using erythrocyte sedimentation rate) [1]. The algorithm had a sensitivity of 72% (95% confidence interval (CI) = 67% to 77%) and a specificity of 91% (95% CI = 89% to 93%). In an editorial in the same issue, Kim and Solomon [2] commented the following: 'a claims-based effectiveness algorithm with acceptable performance characteristics across different data settings will be a powerful and desired tool for CER of RA. Such an algorithm will enable large-scale, population-based studies comparing the effectiveness of different DMARD [disease-modifying antirheumatic drug] regimens. Such studies will facilitate head-to-head comparisons, supplementing typical randomized controlled trials and prospective registries that usually include disease activity. Whether the algorithm will have a similar performance in other claims databases therefore needs to be further examined.' We performed an independent analysis to evaluate the algorithm's positive predictive value (PPV) in a commercial claims data source compared with a clinical gold standard.

Methods

Data came from a previous comparative effectiveness study linking outpatient medical records from multiple US institutions and physician practices to commercial claims data from OptumInsight (Eden Prairie, MN, USA) [3] that evaluated the effectiveness of etanercept (ETN), adalimumab (ADA), and infliximab (INF) in biologic naïve adult RA patients persistent on their initial biologic for at least 1 year from 2006 to 2008. Two teams of two rheumatologists reviewed each medical record and categorized clinical change around 1 year as 'much better', 'better', 'no change', 'worse', or 'much worse'. For this study, the biologic was considered effective if the patient was rated as 'better' or 'much better'. Sensitivity, specificity, and negative predictive value could not be determined, because patients switching biologic agents were excluded from the original study. The PPV compared the classification from the algorithm to the rheumatologist rating. Different compliance thresholds with the biologic medications used by the algorithm were evaluated as sensitivity analyses.

Result

The majority (76%) of the 429 patients in the study were female, and the mean age was 51 years. The PPVs were 86.6% in the primary analysis and 86.5% in sensitivity analyses, similar to that of the original algorithm using VA data. PPV did not differ significantly by biologic ($P > 0.2$): INF (PPV = 95%), ETN (PPV = 86%), and ADA (PPV = 85%).

Conclusions

This previously published administrative claims-based effectiveness algorithm had a high PPV across commercial claims data and VA data. This algorithm may be useful in evaluating the effectiveness of biologic agents by administrative claims data in future studies.

*Correspondence: jcurtis@uab.edu

¹Division of Clinical Immunology and Rheumatology, University of Alabama at Birmingham, 510 20th Street South, FOT 802D, Birmingham, AL 35294, USA
Full list of author information is available at the end of the article

Abbreviations

ADA, adalimumab; CI, confidence interval; ETN, etanercept; INF, infliximab; PPV, positive predictive value; RA, rheumatoid arthritis; VA, Veterans Affairs.

Competing interests

The authors declare that they have no competing interests.

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Author details

¹Division of Clinical Immunology and Rheumatology, University of Alabama at Birmingham, 510 20th Street South, FOT 802D, Birmingham, AL 35294, USA. ²OptumInsight, 12125 Technology Drive, Eden Prairie, MN 55344, USA. ³Amgen Inc., One Amgen Center Drive, Thousand Oaks, CA 91320, USA.

⁴Department of Epidemiology, School of Public Health, University of Alabama at Birmingham, 1700 University Boulevard, Birmingham, AL 35294-0013, USA.

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