

Letter to the editor concerning the article: “Potential Cerebrovascular Accident Signal for Risankizumab: A Disproportionality Analysis of the FDA Adverse Event Reporting System (FAERS)”

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March 6, 2023

Hosted file

Letter to editor_Woods_BJCP_RSinvhal et al_submission_rev.docx available at <https://authorea.com/users/592581/articles/628016-letter-to-the-editor-concerning-the-article-potential-cerebrovascular-accident-signal-for-risankizumab-a-disproportionality-analysis-of-the-fda-adverse-event-reporting-system-faers>