

**Title:** Analyzing inpatient vaccine reactions amongst systemic lupus erythematosus patients in the COVID-19 era: a national analysis

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**Introduction:** Patients with systemic lupus erythematosus (SLE) are prone to severe infection given the aberrant immune responses inherent to their disease and immunosuppressive medications used to manage their disease.<sup>1</sup> During the coronavirus of 2019 (COVID-19) and subsequent vaccine rollout, SLE patient hospitalizations including those due to COVID-19 vaccine reactions were areas of concern. While the VACOLUP trial of 700 SLE patients showed disease flares in 3% of patients after COVID-19 vaccine,<sup>2</sup> subsequent studies reported flares in one third of patients after vaccine.<sup>3</sup> We aimed to characterize hospitalization trends on a national scale amongst SLE patients following COVID-19 vaccine rollout.

**Methods:** The 2016-2020 Nationwide Inpatient Sample was queried for SLE hospitalizations using International Classification of Diseases, Tenth Revision (ICD-10) code “M32”. SLE hospitalizations with co-morbid vaccine reactions (RX+) were identified using ICD-10 codes “T50.x”. Chi-square and t-tests were used to compare clinical and demographic features and rates of high-risk underlying medical conditions, as defined by the CDC, between RX+ and RX-SLE inpatients.

**Results:** Among 1,082,960 total SLE hospitalizations between 2016-2020, 15,625 (1.4%) were RX+. RX+ patient hospitalizations increased from 1.2-1.3% in 2016-2018 to 1.6% in 2019 and 2020 ( $p<0.001$ ). RX+ patients were on average 3 years older (55.0 vs. 52.1 years,  $p<0.0001$ ), with a higher proportion of males (12.4% vs. 11.7%,  $p=0.01$ ). RX+ patients were more likely to identify as Black (31.6% vs. 30.9%,  $p=0.02$ ), be in the highest income quartile (19.7% vs. 17.6%,  $p<0.0001$ ), and use either Medicare (54.0% vs. 49.0%) or self-pay insurance (3.6% vs. 3.0%) ( $p<0.0001$ ). RX+ patients had a 5% greater proportion of non-routine discharges, with higher proportions transferred to other facilities (40.5% vs. 35.3%,  $p<0.0001$ ). SLE RX+ patients exhibited significantly different rates of high-risk underlying medical conditions. RX+ patients had a greater burden of heart disease (48.5% vs. 34.1%,  $p<0.0001$ ) and chronic kidney disease (36.3% vs. 29.7%,  $p<0.0001$ ). RX+ patients were more likely to be elderly, have chronic liver or lung disease, HIV, and obesity ( $p<0.01$ ).

**Conclusion:** Increases in vaccine-related SLE hospitalizations were observed during the COVID-19 pandemic. Hospitalized SLE patients with vaccine reactions had a higher proportion of co-morbidities, most notably heart and kidney disease. Limitations included possible error in coding of vaccine reaction by recording physician, lack of data regarding specific vaccine formulation triggering reaction, and medication information.

## References

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