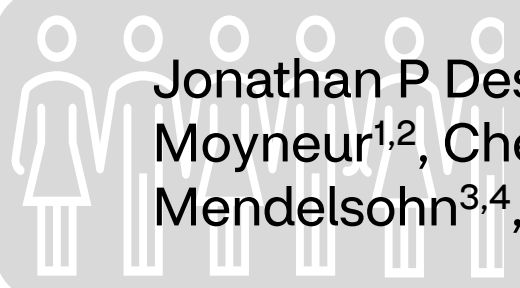


Adherence Differences Between **Self-Administered** and **Provider-Administered** Biologics Among Patients with Rheumatoid Arthritis During the COVID-19 National Emergency



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1) CVS Health, 2) StatLog Inc., 3) Harvard Pilgrim Health Care Institute, 4) Harvard Medical School, 5) Biologics and Biosimilars Collective Intelligence Consortium

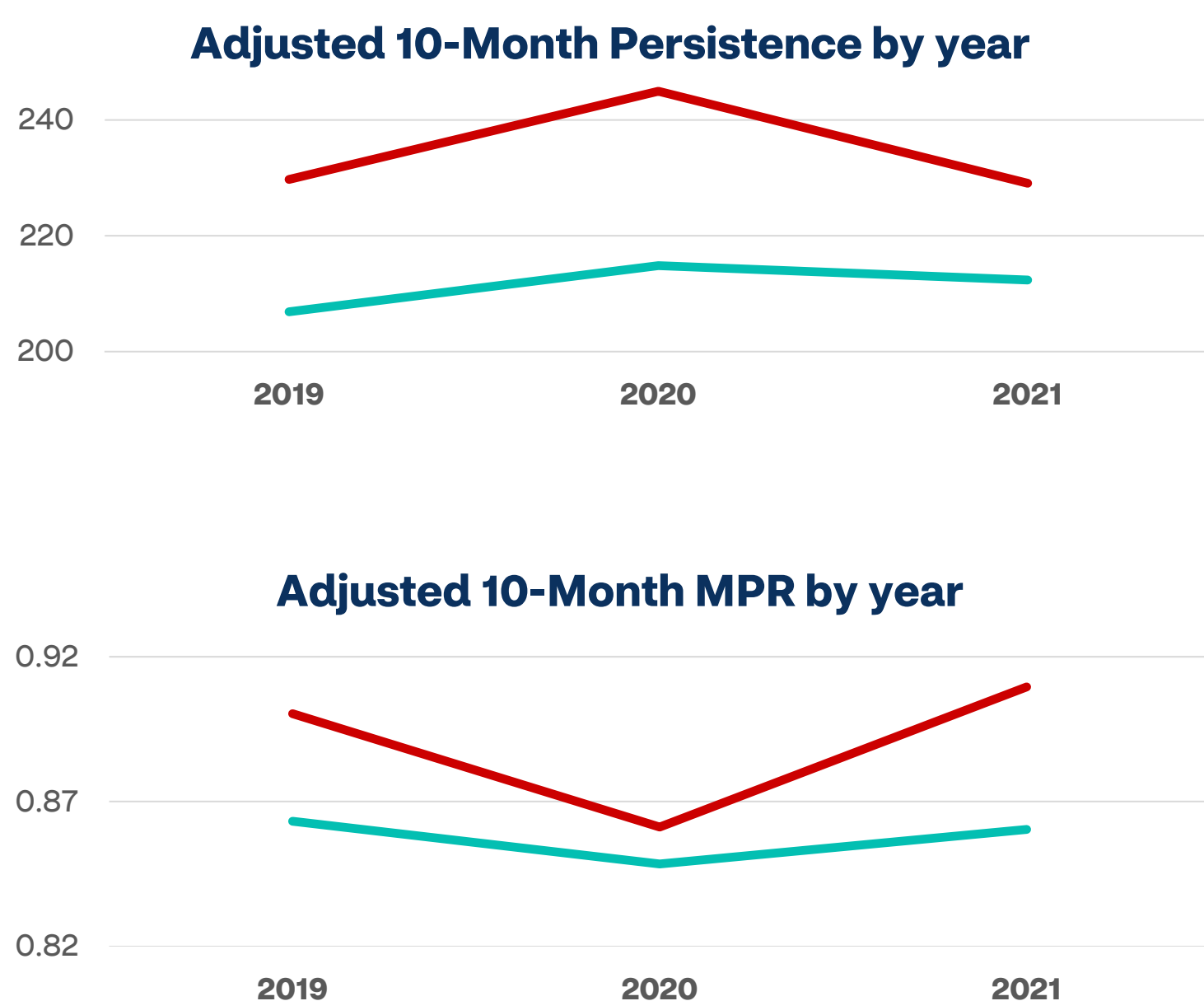
Background

Patients must adhere to the prescribed therapeutic regimen to achieve the best clinical outcomes. COVID-19 presented numerous barriers to Rheumatoid Arthritis (RA) patients taking biologics, particularly those receiving provider-administered (P-A) biologics in a healthcare setting. Previous work has suggested a decrease in P-A biologic adherence after the onset of COVID-19, yet the effects of COVID on RA therapy adherence to P-A and self-administered (S-A) biologics is unknown.

RA Biologics Administration Options		
Self-Administered (S-A)		Provider-Administered (P-A)
✓	Abatacept	✓
✓	Adalimumab	X
✓	Anakinra	✓
✓	Canakinumab	✓
✓	Certolizumab pegol	✓
✓	Etanercept	X
✓	Golimumab	✓
X	Infliximab	✓
X	Rituximab	✓
✓	Sarilumab	X
✓	Tocilizumab	✓

Results

The annual cohorts represented 11,361 patients total and were comparable in biologics, S-A ratio, sex, ethnicity, history of specific comorbidities, and combined comorbidity score.



Objectives

To assess changes in adherence during the COVID-19 emergency among RA patients taking P-A and S-A biologics.

Methods

We conducted a retrospective, repeated, cross-sectional analysis of administrative claims for adults with an RA diagnosis and using a biologic. Infusion/ injection procedures administered by a health care professional and retail pharmacy dispensing were considered P-A and S-A, respectively. We compared fixed-window medication possession ratio (MPR) and persistence among independent cohorts for 2019, 2020, and 2021 using a mixed model adjusting for demographics, biologic, and comorbidities. The analysis window was March 1 until discontinuation or December 31 of the cohort year. Persistence was calculated as days between March 1 and last dispense, plus last days' supply. MPR was calculated as days' supply dispensed divided by days in the analysis window.

In 2019, S-A MPR was 0.90 (95%CL 0.86 - 0.94) and P-A MPR was 0.86 (95%CL 0.83 - 0.90); S-A persistence was 229.7 (95% CL 198.4 - 261.1) and P-A persistence was 206.9 (95%CL 181.0 - 232.7).

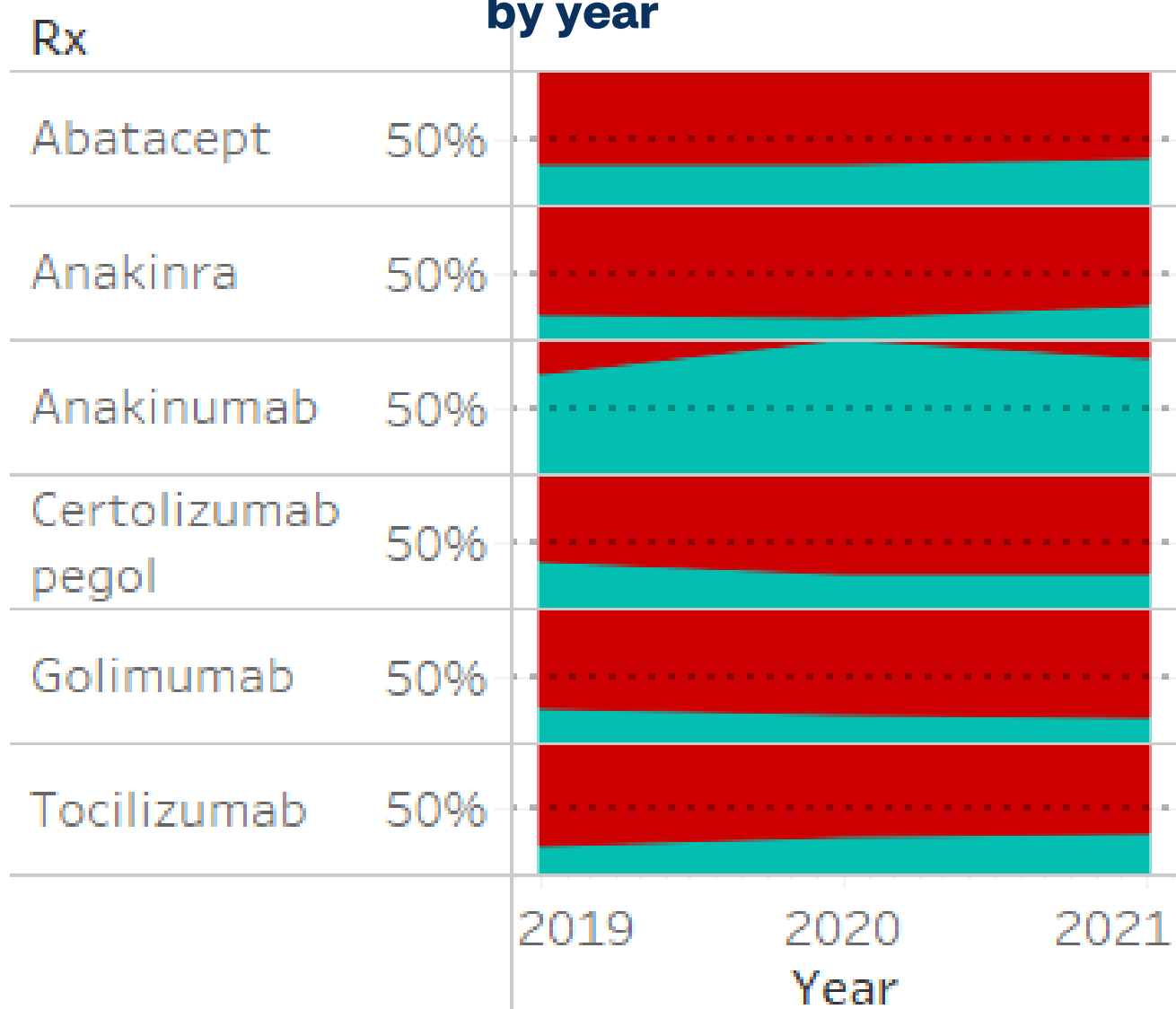
In 2020, **S-A MPR dropped to 0.86 (95%CL 0.82 - 0.90)** and **P-A MPR to 0.85 (95%CL 0.81 - 0.88)**; **S-A persistence increased to 245.0 (95%CL 213.6 - 276.3)** and **P-A persistence to 214.8 (95%CL 189.0 - 240.7)**. Changes in MPR and persistence differed significantly between S-A and P-A. By 2021, both MPR and persistence returned to levels comparable to 2019.

Conclusions

Decreasing MPR alongside increasing persistence suggests that both S-A and P-A patients experienced increased days without therapy during the first year of COVID-19, however differences between administration modalities need to be investigated further.

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% Self-Administered v. Provider-Administered by year



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