

MEASURING ADHERENCE TO AND PERSISTENCE WITH ADVANCED THERAPY IN US PATIENTS WITH LUPUS NEPHRITIS IN THE REAL-WORLD SETTING



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BACKGROUND

- Lupus nephritis (LN) is a severe manifestation of systemic lupus erythematosus, marked by kidney inflammation and damage¹⁻³
- While advanced therapies have transformed the management of LN, their real-world clinical benefit relies heavily on sustained patient adherence, which remains under-characterized

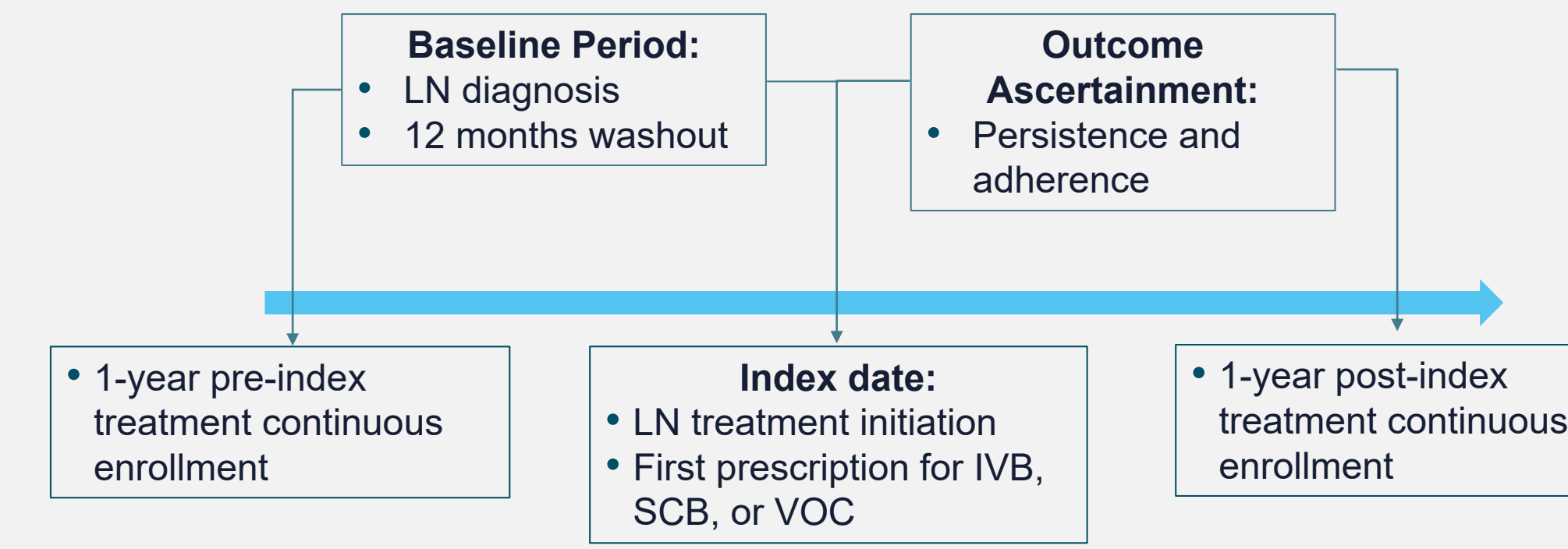
OBJECTIVE

- To measure real-world adherence to, and persistence with, advanced therapies (belimumab [BEL] and voclosporin [VOC]) in patients with LN

METHODS

- The IQVIA PharMetrics Plus closed claims database was used to identify adult US patients diagnosed with LN between January 1, 2020, and September 30, 2025 (study period)
- Adult participants with ≥2 outpatient claims ≥30 days apart or ≥1 inpatient claim for LN between January 1, 2020, and September 30, 2025, with continuous enrollment for a year of pre- and post-index, who were treated with intravenous belimumab (IVB), subcutaneous belimumab (SCB) or VOC, and had a 12-month washout period (no evidence of use of index therapy in the year prior to index date) were included; index date was the date of first index therapy
- Participants with a renal transplant, dialysis, or end-stage renal disease in the 12-month pre-index period were excluded
- The proportion of days covered (PDC) and medication possession ratio (MPR) were calculated over 1 year; measures of medication persistence over 1 year were also reported
 - MPR was defined as the total days' supply, divided by the number of days in the period; refills (including early refills) were summed
 - PDC was defined as the total number of days covered by prescription (i.e., patient had supply on hand), divided by the total number of days in the period. In PDC, early fills were shifted forward, rather than added to existing supply
- All endpoints were stratified by therapy and by payer type

Figure 1. Cohort Definition



Description	N (%)	Change
LN diagnosis* with: • 2 outpatient claims ≥30 days apart or • 1 inpatient claim in the pre-index period between January 1, 2020, and September 30, 2025	31,855 (100.00)	NA
Claim for an advanced therapy* between January 1, 2021, and September 30, 2024, with a 12-month washout for specific treatment	2,610 (8.19)	-29,245
≥18 years old at index date	2,505 (7.86)	-105
Continuous enrollment: 12 months pre- and post-index	736 (2.31)	-1,769
Exclude patients with renal transplant, dialysis, or end-stage renal disease in the 12-month pre-index period	645 (2.02)	-91
No malignancy, HIV/AIDS, HBV, HCV, or TB in the study period	577 (1.81)	-68

CONCLUSIONS

- This real-world study showed that overall adherence to and persistence with advanced therapies (BEL and VOC) in patients with LN were poor
- Only 38–54% of patients treated with IVB, SCB, or VOC achieved the adherence benchmark of PDC ≥80%; adherence was poorer for more frequent SC and oral treatments compared with IV treatment
- Only 50–54% of patients across treatment types persisted with medication at 1 year
- Patients covered by Medicaid demonstrated poorer adherence and persistence compared with those with commercial insurance
- Method of administration appears to be a modulating factor, with better observed adherence and persistence in IV versus SC or oral medications
- Poor adherence and persistence are known drivers of reduced real-world effectiveness compared with efficacy measured in clinical trial populations; our findings suggest an urgent need to improve advanced therapy adherence and persistence in patients with LN

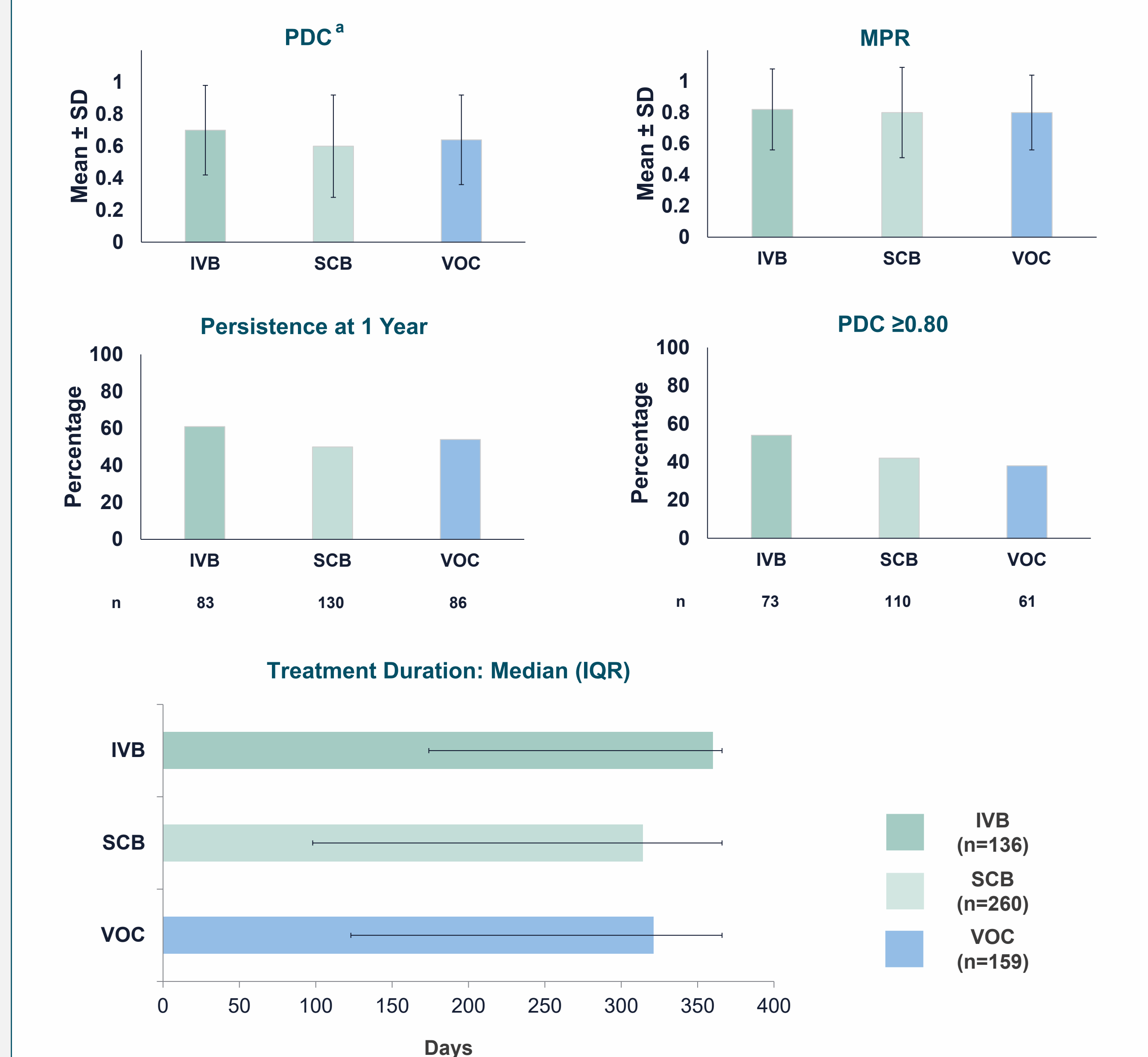
RESULTS

Table 2. Baseline Characteristics

	BEL ^a (N=418)	VOC (N=159)
Age, years		
Mean (SD)	37 (14)	34 (11)
Median (Q1, Q3)	35 (25, 46)	34 (25, 42)
Min, Max	18, 85	18, 70
Age category, years, n (%)		
18–30	159 (38)	63 (40)
31–50	183 (44)	87 (55)
>50	76 (18)	<11 ^b
Sex, n (%)		
Male	44 (11)	24 (15)
Female	374 (89)	135 (85)
Region, n (%)		
South	146 (35)	69 (43)
Midwest	82 (20)	30 (19)
East	13 (3.1)	<11 ^b
West	121 (29)	30 (19)
Northeast	54 (13)	26 (16)
Unknown	<11 ^b	<11 ^b
Payor type, n (%) ^c		
Commercial/self-insured	226 (54)	85 (53)
Medicare Advantage	23 (5.5)	<11 ^b
Medicaid	166 (40)	72 (45)
Insurance product		
PPO	152 (36)	52 (33)
HMO	227 (54)	98 (62)
Other	39 (9.3)	<11 ^b
CCI score		
Mean (SD)	2 (2)	2 (2)
Median (Q1, Q3)	2 (1, 3)	1 (1, 3)
Min, max	1, 9	1, 10

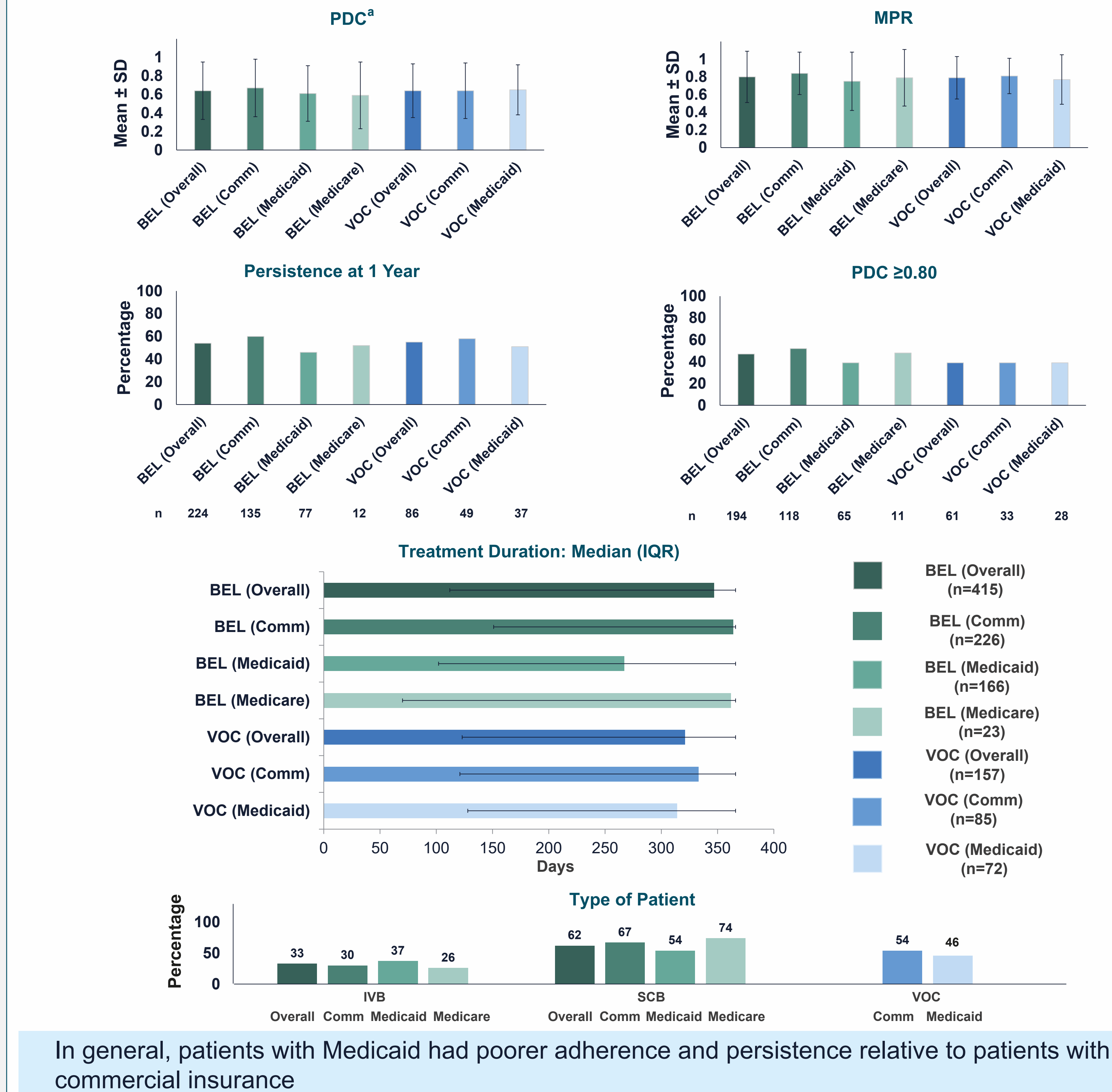
- Of 577 identified patients, 45%, 24%, and 28% were treated with SCB, IVB, and VOC, respectively; the remainder received both SCB and IVB
- Baseline patient characteristics were similar between treatment cohorts
- Median CCI score was higher for BEL vs VOC

Figure 2. Persistence and Adherence



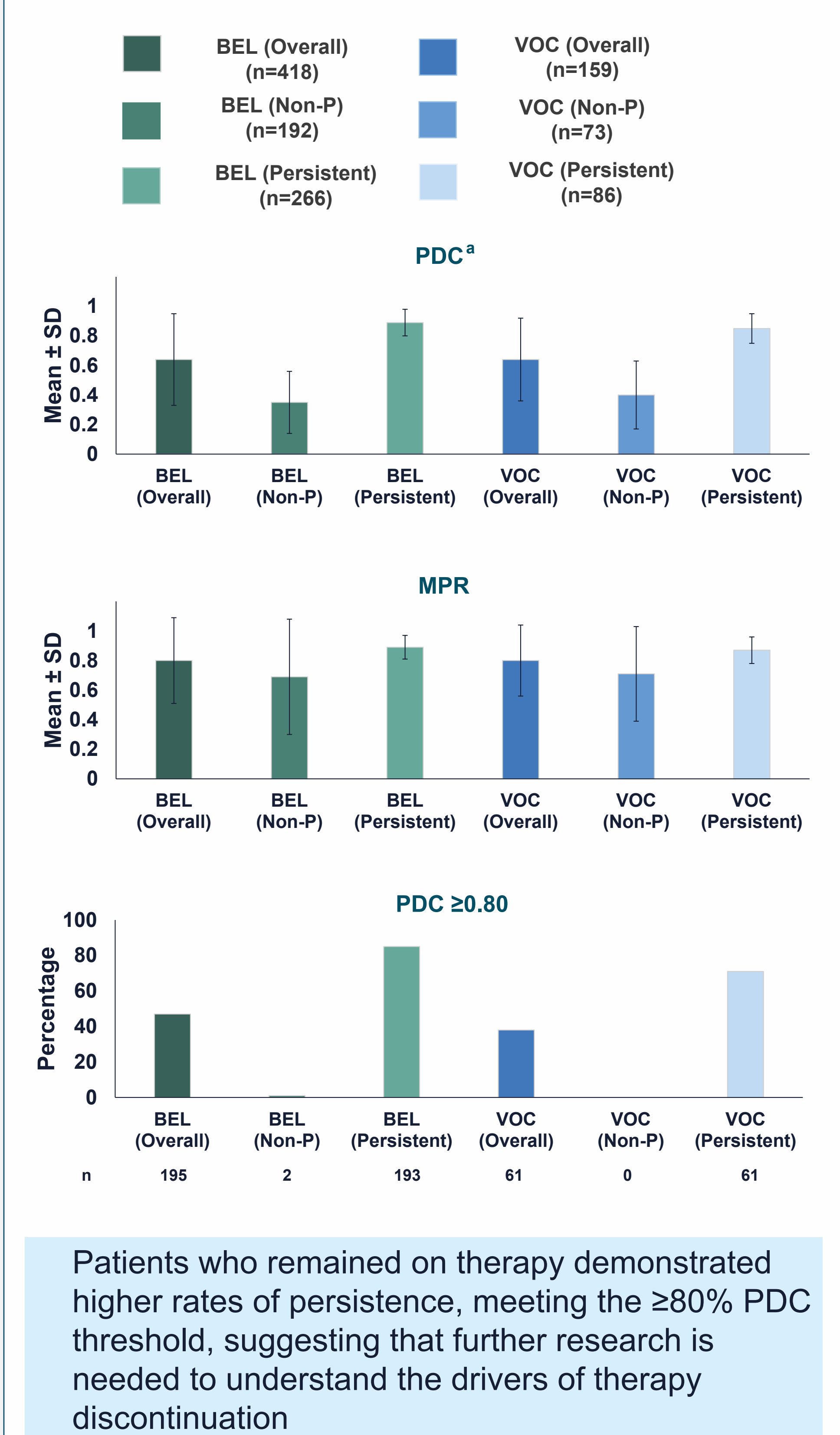
- PDC for the IVB, SCB, and VOC cohorts was 0.70, 0.60, and 0.64, respectively; MPR was 0.82, 0.80, and 0.80, respectively
- The percentage of patients persistent with medication at 1 year was 61%, 50%, and 54% for the IVB, SCB, and VOC cohorts, respectively
- The benchmark of ≥0.80 PDC was achieved by 54% of patients receiving IVB, 42% of patients receiving SCB, and 38% of patients receiving VOC

Figure 3. Results by Insurance Type



In general, patients with Medicaid had poorer adherence and persistence relative to patients with commercial insurance

Figure 4. Results by Persistence



Patients who remained on therapy demonstrated higher rates of persistence, meeting the ≥80% PDC threshold, suggesting that further research is needed to understand the drivers of therapy discontinuation

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ABBREVIATIONS

AIDS, acquired immune deficiency syndrome; BEL, belimumab; CCI, Charlson Comorbidity Index; Comm, commercial insurance (including self-insurance); HBV, hepatitis B virus; HCV, hepatitis C virus; HIV, human immunodeficiency virus; HMO, Health Maintenance Organization; ICD-10, International Classification of Diseases, Tenth Revision; IQR, interquartile range; IV, intravenous; IVB, intravenous belimumab; LN, lupus nephritis; MPR, medication possession ratio; NA, not available; P, persistent; PDC, proportion of days covered; PPO, Preferred Provider Organization; Q, quartile; SC, subcutaneous; SCB, subcutaneous belimumab; SD, standard deviation; TB, tuberculosis; VOC, voclosporin.

DISCLOSURES

All authors are employees of Genentech, Inc. and shareholders of F. Hoffmann-La Roche Ltd.

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