

Supplementary Materials

Effectiveness and safety of belimumab in Chinese lupus patients: a multicenter, real-world observational study

Fangfang Sun¹, Zitao Wang², Tong Wu³, Xue Wu⁴, Jie Chen¹, Danting Zhang¹, Chunde Bao¹, Nan Shen¹, Huaxiang Wu², Jing Zhu³, Lijun Wu^{4,5}, Shuang Ye¹

1. Department of Rheumatology, Renji Hospital, School of Medicine, Shanghai Jiaotong University, Shanghai, China.
2. Department of Rheumatology, The Second Affiliated Hospital Zhejiang University School of Medicine, Zhejiang, China.
3. Department of Rheumatology, Sichuan Provincial People's Hospital, University of Electronic Science and Technology of China, Chengdu, China.
4. Department of Rheumatology, People's Hospital of Xinjiang Uygur Autonomous Region, Xinjiang, China
5. Xinjiang Clinical Research Center for Rheumatoid arthritis, Xinjiang, China

Fangfang Sun and Huaxiang Wu contributed equally to this work.

Corresponding authors: Jing Zhu, Lijun Wu.

Table S1. Baseline characteristics of patients in this real-world study from four centers.

	Items	Center 1 (n=52)	Center 2 (n=48)	Center 3 (n=52)	Center 4 (n=72)
Demographic features	Gender (%)	50 (96.2)	44 (91.7)	50 (96.2)	69 (95.8)
	Age (y)	35.54 (11.42)	31.33 (9.44)	33.90 (10.00)	35.03 (11.92)
	Han	52 (100)	48 (100)	52 (100)	59 (81.9)
	Uygur	0 (0)	0 (0)	0 (0)	13 (18.1)
	SLE duration (y)	10.71 (6.34)	8.84 (5.97)	5.71 (4.67)	6.03 (6.54)
	SLE duration≤2 (%)	7 (13.5)	6 (12.5)	15 (28.8)	37 (51.4)
	Follow up (m)	13.29 (5.87)	12.27 (5.33)	10.35 (4.53)	11.22 (4.48)
Laboratory results	Anti-dsDNA+%	35 (67.3)	34 (70.8)	42 (80.8)	58 (80.6)
	Complement 3 (g/L)	0.66 (0.20)	0.67 (0.25)	0.61 (0.19)	0.59 (0.19)
	Complement 4 (g/L)	0.12 (0.08)	0.10 (0.05)	0.12 (0.10)	0.13 (0.09)
	IgG (g/L)	12.63 (4.35)	15.03 (4.71)	12.81 (4.66)	13.18 (6.32)
Organ involvement	Mucocutaneous	8 (15.4)	18 (37.5)	21 (40.4)	13 (18.1)
	Musculoskeletal	4 (7.7)	16 (33.3)	12 (23.1)	15 (20.8)
	Renal	17 (32.7)	9 (18.8)	27 (51.9)	22 (30.6)
	Hematologic	6 (11.5)	5 (10.4)	18 (34.6)	9 (12.5)
	Neurological	2 (3.8)	1 (2.1)	0 (0)	2 (2.8)
Disease evaluation	SLEDAI	6.75 (3.96)	6.44 (4.70)	10.69 (4.54)	7.12 (3.63)
	SDI	0.63 (0.89)	0.31 (0.47)	0.27 (0.53)	0.65 (0.79)
	PGA	1.12 (0.66)	1.36 (0.67)	1.63 (0.63)	1.26 (0.46)
Baseline Therapy	Prednisone (mg/d)	17.46 (11.34)	22.08 (24.61)	30.50 (22.08)	25.21 (17.50)
	HCQ (%)	50 (96.2)	38 (79.2)	45 (86.5)	65 (90.3)

IS (%)	29 (55.8)	32 (66.7)	46 (88.5)	27 (37.5)
MMF (%)	21 (40.4)	14 (29.2)	19 (36.5)	15 (20.8)
CTX (%)	0 (0.0)	0 (0.0)	6 (11.5)	7 (9.7)
CNI (%)	5 (9.6)	17 (35.4)	16 (30.8)	5 (6.9)
Others (%)	3 (5.8)	1 (2.1)	5 (9.6)	0 (0)

Table S2. Baseline characteristics of mild disease subgroup and “reference group” from previous randomized trials before and after PSM.

Before PSM				After PSM		
All patients	Belimumab (n=98)	Reference (n=130)	p	Belimumab (n=57)	Reference (n=57)	p
Gender (%)	90 (91.8)	122 (93.8)	0.607	51 (89.5)	54 (94.7)	0.490
Age (y)	34.23 (9.80)	32.63 (12.41)	0.047	31.81 (9.66)	31.75 (10.45)	0.792
SLE duration (y)	8.98 (7.06)	4.64 (4.77)	<0.001	6.07 (4.74)	5.47 (4.87)	0.431
Anti-dsDNA+%	64 (65.3)	71 (59.2)	0.274	36 (63.2)	36 (63.2)	1.000
Complement 3 (g/L)	0.70 (0.22)	0.79 (0.23)	0.001	0.72 (0.23)	0.75 (0.20)	0.300
Complement 4 (g/L)	0.13 (0.08)	0.14 (0.08)	0.309	0.13 (0.09)	0.13 (0.07)	0.556
SLEDAI	3.91 (1.64)	3.01 (2.63)	<0.001	3.54 (1.69)	3.96 (2.83)	0.971
Prednisone (mg/d)	14.77 (11.70)	12.12 (7.60)	0.412	12.89 (10.26)	13.43 (6.28)	0.125
HCQ (%)	85 (86.7)	114 (88.4)	0.690	48 (84.2)	48 (84.2)	1.000
IS (%)	51 (52.0)	80 (61.5)	0.177	30 (52.6)	36 (63.2)	0.343
Flare	16 (16.3)	47 (36.2)	0.001	10 (17.5)	22 (38.6)	0.021
Major Flare	4 (4.1)	32 (24.6)	<0.001	3 (5.3)	16 (28.1)	0.0026