Amulirafusp alfa (IMM0306)
- First-in-Class CD47×CD20 Bispecific Antibody for SLE
- Preliminary Phase I Results

ImmuneOnco (01541.HK)

BIO International Convention

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Amulirafusp Alfa (IMM0306) - A Novel CD47xCD20 Bispecific Antibody with Best-in-disease Potential in SLE

Dual targeting of CD47 and CD20

- Engineered IgG1 Fc
- Strong ADCC/ADCP
- Safe to RBC in vitro

First-in-class for autoimmune diseases

- Rapid, efficient and sustained B-cell depletion
- Immune reconstitution with lower risk of infection

Best-in-disease potential

- 83.3% response in SLEDAI-2K at 1.2 mg/kg [1]
- No CRS
- Improvement in multiple measurements

Multiple indications in development

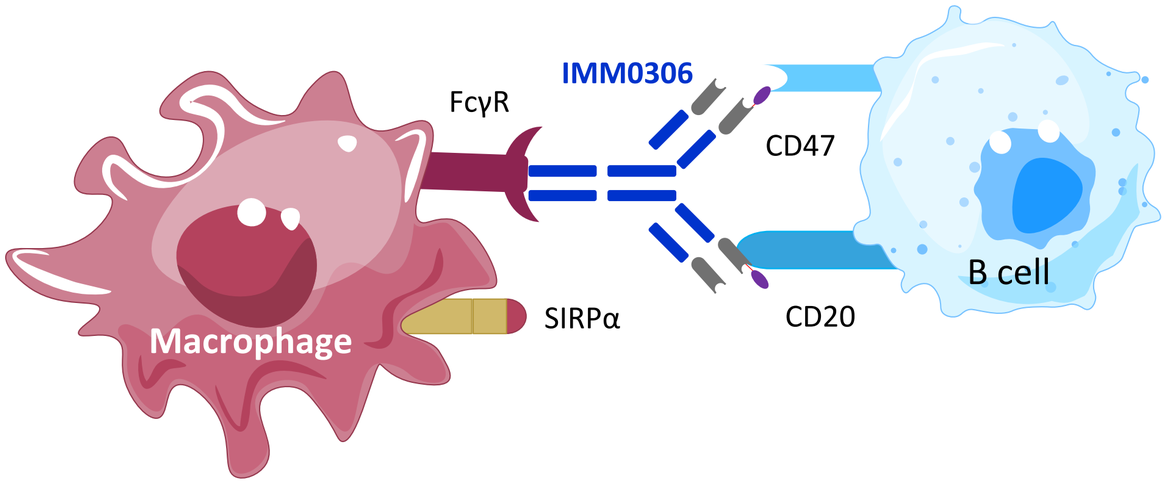
- Phase II in follicular lymphoma ongoing: CRR 64.7%
- Approved IND
  - China: SLE, LN, NMOSD, NHL
  - USA: NHL

1. Defined as the percentage of patients (SLEDAI-2K ≥8) achieving ≥4-point reduction from baseline.

ADCP: Antibody-dependent cellular phagocytosis; ADCC: Antibody dependent cell-mediated cytotoxicity. RBC: red blood cell;

CRR: complete response rate; SLE: systemic lupus erythematosus; NMOSD: neuromyelitis optica spectrum disorder; LN: Lupus nephritis; NHL: Non-Hodgkin lymphoma

Mechanism of Action - Amulirafusp alfa (IMM0306)

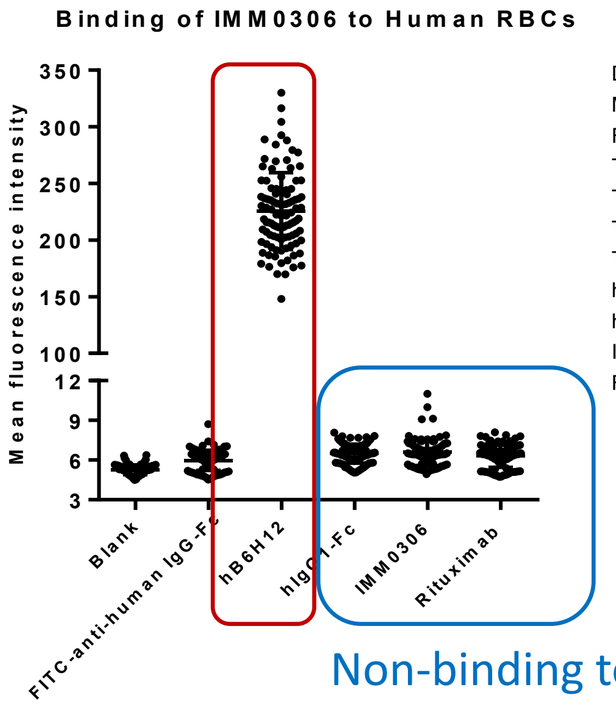


- IMM0306 is a **fusion protein of CD20 mAb with the CD47 binding domain of SIRPα** on both heavy chains.
- IMM0306 possesses:
  - **Stronger ADCC/ADCP activity compared to rituximab**
  - **No in vitro binding to human RBC**
  - **Higher affinity to CD20 better avoid normal cell killing**

ADCP: Antibody-dependent cellular phagocytosis; ADCC: Antibody dependent cell-mediated cytotoxicity.

Amulirafusp alfa (IMM0306) is In Vitro Safe to Red Blood Cells (RBC)

In vitro binding assay to RBC



Donor Characteristics (n=100)
Male (n=62)
Female (n=38)
Type A blood group (n=29)
Type B blood group (n=31)
Type AB blood group (n=10)
Type O blood group (n=30)
hB6H12:500nM
hIgG1-Fc:5000nM
IMM0306:5000nM
Rituximab:5000nM

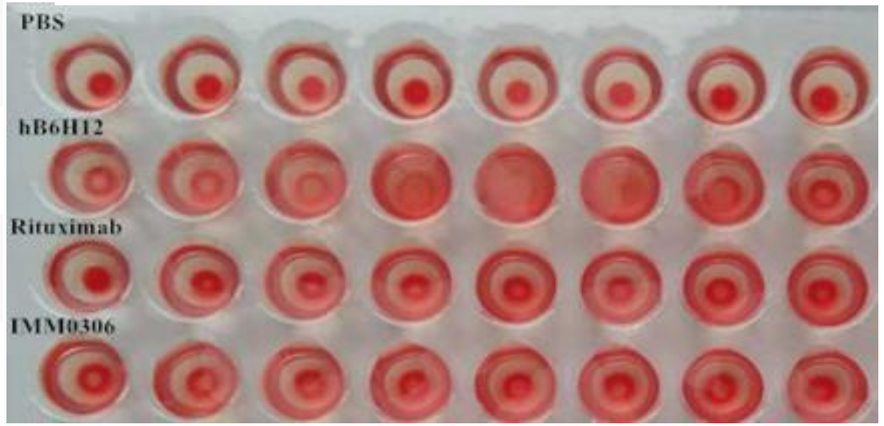
Non-binding to human RBC

Hemagglutination assay

	Hemagglutination
PBS (negative control)	No
hB6H12 (positive control)	Yes
Rituximab	No
IMM0306	**No**

Concentration (nM)

4000 1333 444 148 49.4 16.5 5.49 1.83



IMM0306 is Efficacious as Monotherapy, in Combo with Lenalidomide and to anti-CD20-treated Lymphoma Patients

	Phase I	Phase II	Patients with prior anti-CD20 treatment (obinutuzumab)
Treatment	Monotherapy[1]	Combined with Lenalidomide[2]	Combined with Lenalidomide[2]
	Follicular lymphoma (n = 17)	Follicular lymphoma (n = 34)	n = 10
CR	**4 (23.5%)**	**22 (64.7%)**	**5 (50%)**
PR	**3 (17.6%)**	**8 (23.5%)**	**3 (30%)**
SD	4 (23.5%)	2 (5.9%)	1 (10%)
PD	6 (35.3%)	2 (5.9%)	0
ORR	**7 (41.2%)**	**30 (88.2%)**	**8 (80%)**
DCR	11 (64.7%)	32 (94.1%)	9 (90%)

1. IMM0306 monotherapy data is as of April 18, 2024, among 17 efficacy evaluable patients with r/r FL who received doses 0.8-2.0 mg/kg. 2. Cut off date as June 9, 2025

Significant Unmet Needs Among Systemic Lupus Erythematosus (SLE) Patients

3.4 million

Global SLE population[1]

10th leading cause of death

in females 15-24 yr, USA[2]

400 k/yr

Newly diagnosed SLE patients[1]

Top 20 leading cause of death

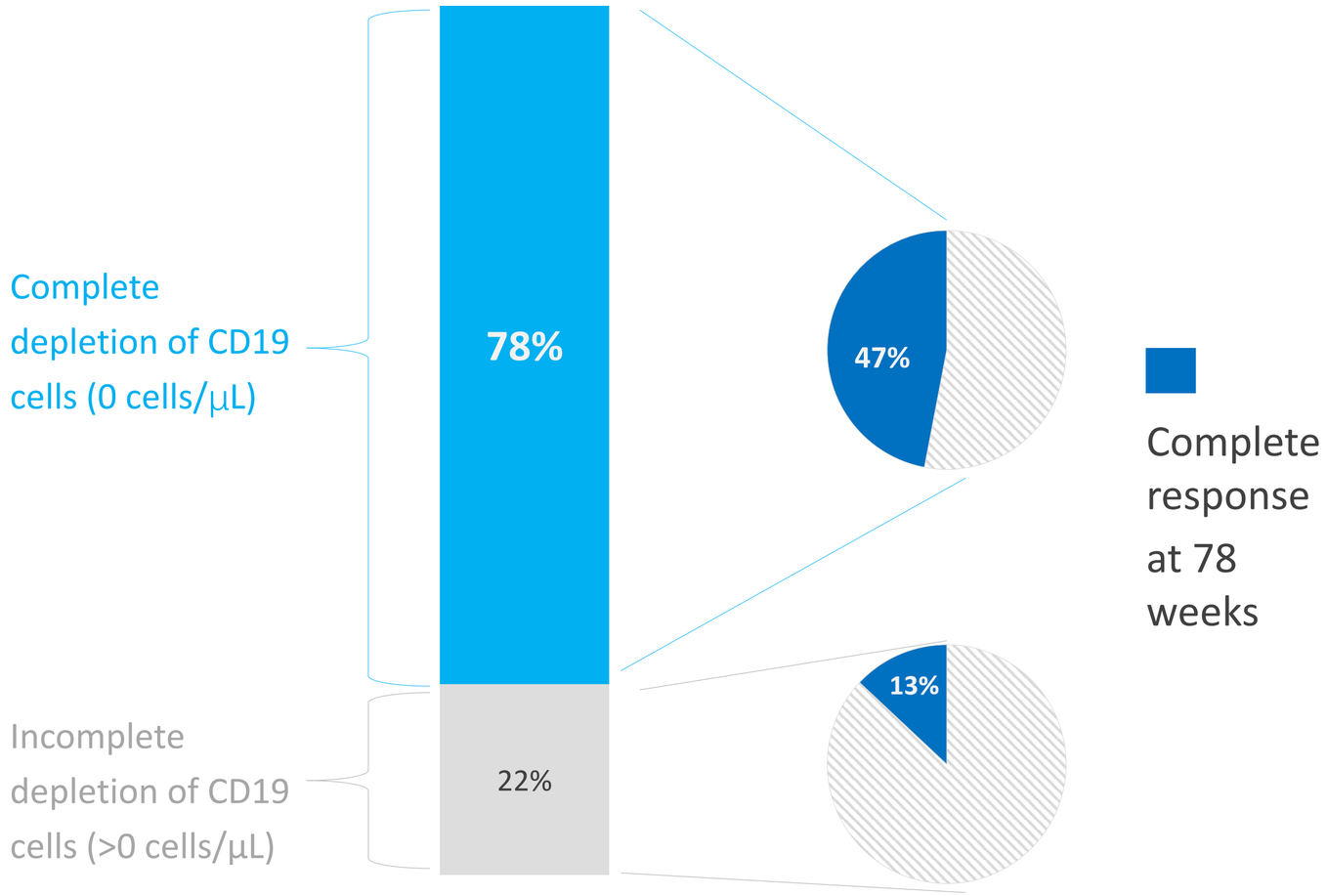
in females 5-64 yr, USA[2]

1. Tian et al. Ann Rheum Dis. 2023 Mar;82(3):351-356. 2. Yen et al. Arthritis Rheumatol. 2018 Aug;70(8):1251-1255.

Enhancing B-Cell Depletion for Greater Efficacy

"Although the B cell depletion agent rituximab failed to reach its primary end points in randomized controlled trials in systemic lupus erythematosus (SLE), favorable clinical experience has led to its frequent off-label use in patients with SLE."[1]

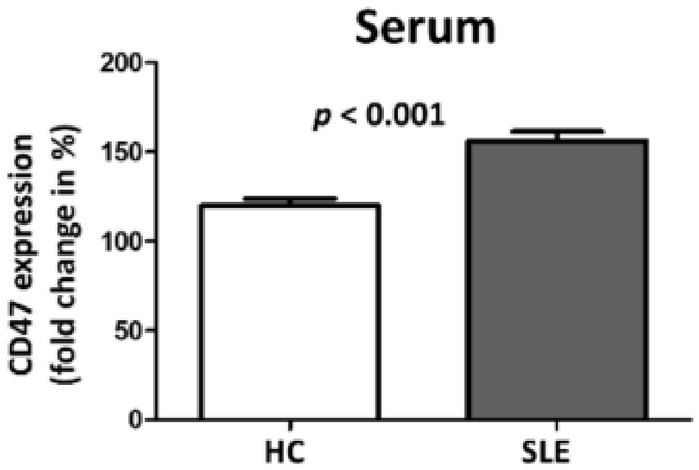
Percentage of patients in LUNAR study - anti-CD20 rituximab in lupus nephritis[2]



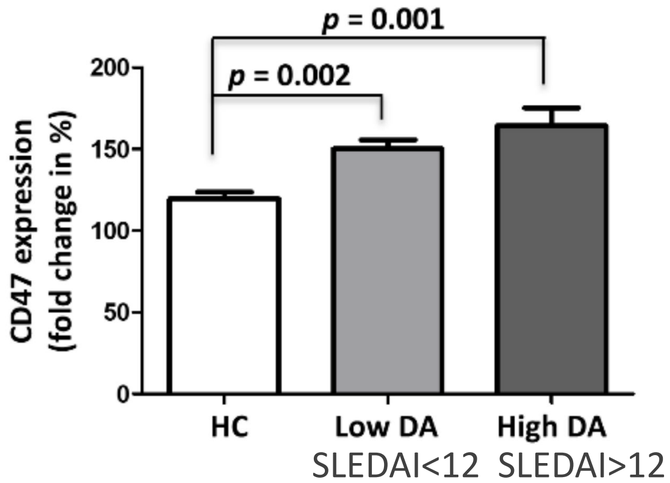
1. Stockfelt et al. Nat Rev Rheumatol. 2025 Feb;21(2):111-126. 2. Mendez et al. Clin J Am Soc Nephrol. 2018 Aug 8;13(10):1502–1509.

CD47 Expression Links to SLE Disease Activity and IFN-α Upregulation

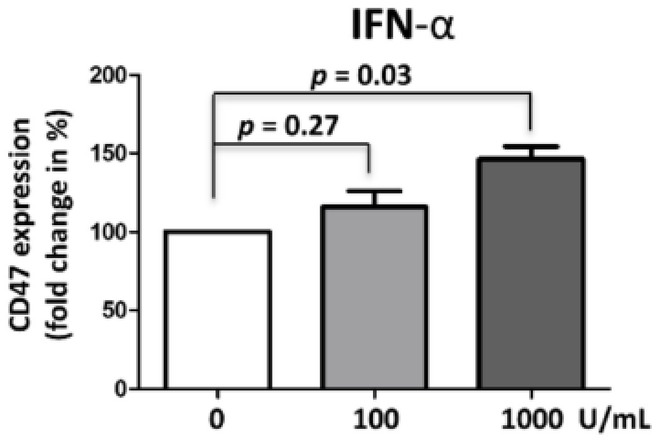
Upregulation of CD47 by SLE serum



Subgroup analysis



Expression of CD47 at presence of IFN-α



- Elevated CD47 expression makes it a promising therapeutic target for SLE.

Park et al. Cells. 2021 May 10;10(5):1151. HC: Healthy control serum; DA: disease activity.

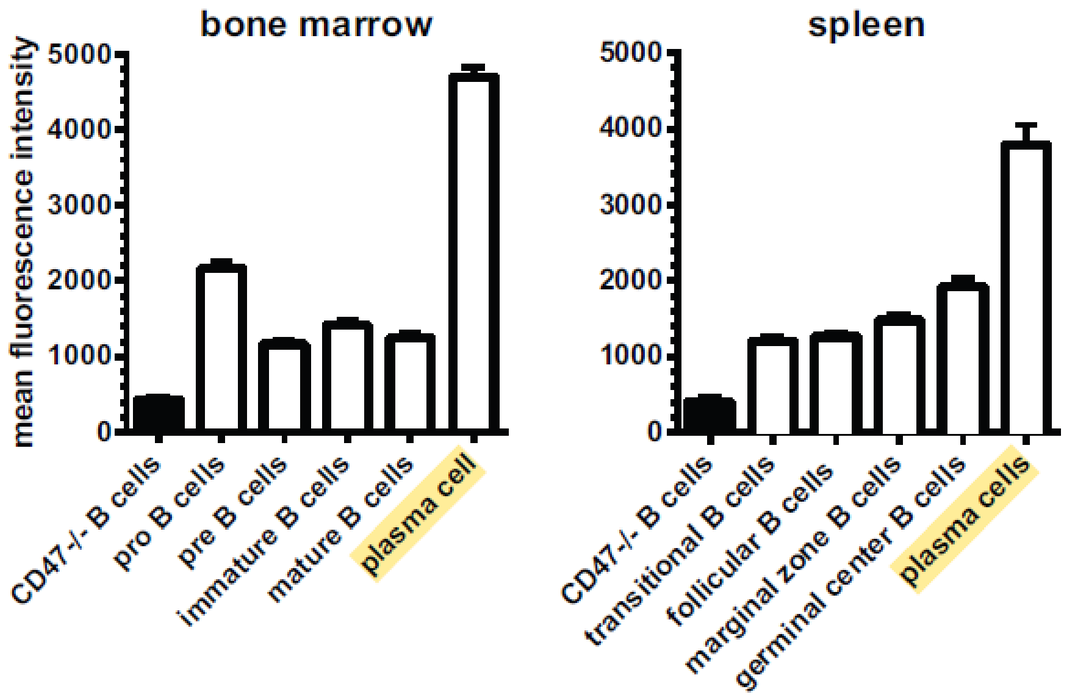
Left: Healthy PBMCs were incubated with serum from healthy controls (HC, n = 6) and SLE patients (n = 10), and fold changes in CD47 expression on monocytes were investigated by flow cytometry analysis.

Middle: Effect of serum from patients with low (n = 6) and high (n = 4) disease activity on CD47 expression was examined.

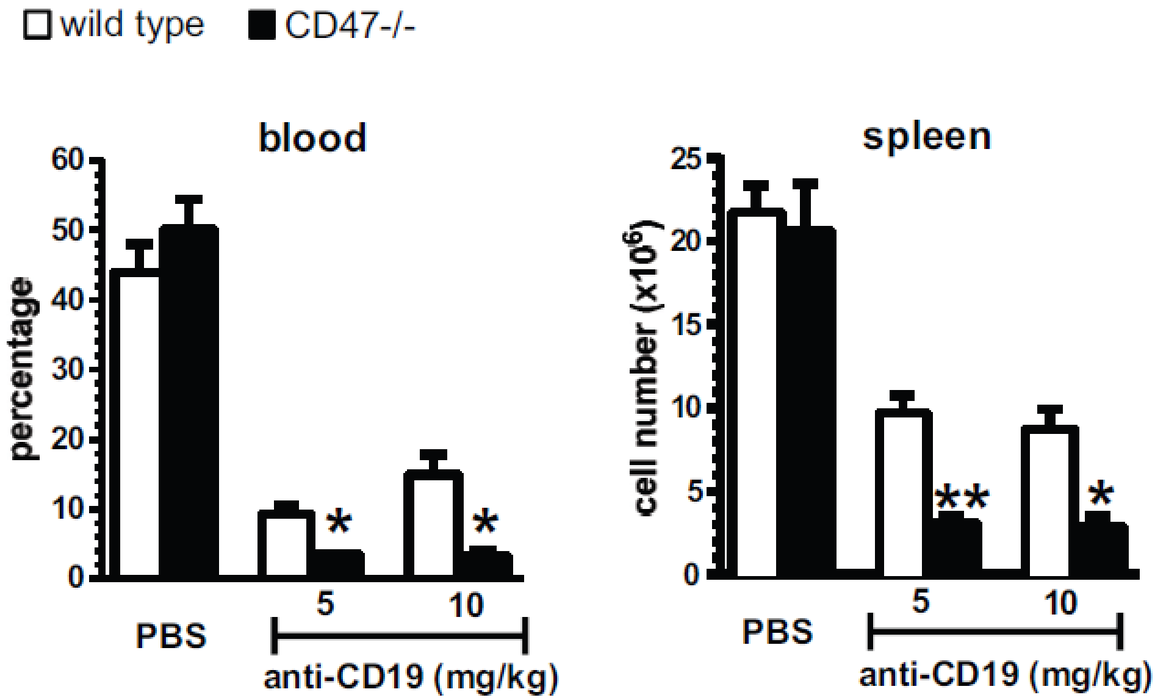
Right: Healthy PBMCs (n = 3) were incubated with increasing concentrations of interferon-alpha (IFN-α) and change in CD47 expression was examined by flow cytometry. Untreated samples served as a reference (i.e., 100%).

Enhanced B-Cell Depletion in CD47-Deficient Mice

Expression of CD47 on B cell subsets



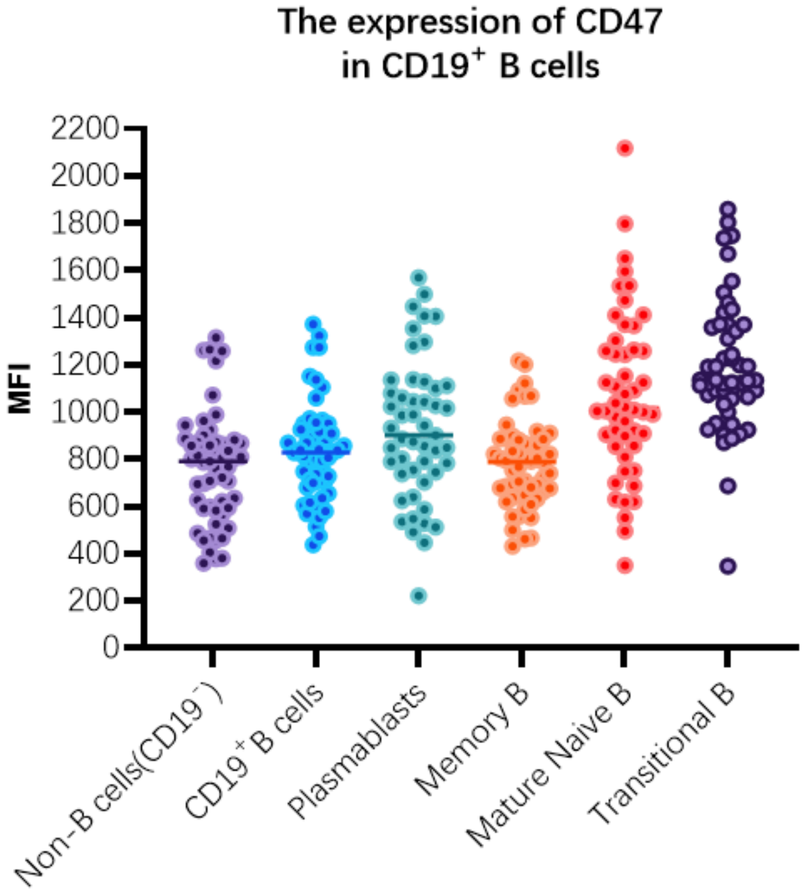
CD19^{+} B cell depletion is enhanced in CD47$^{-/-}$ mice

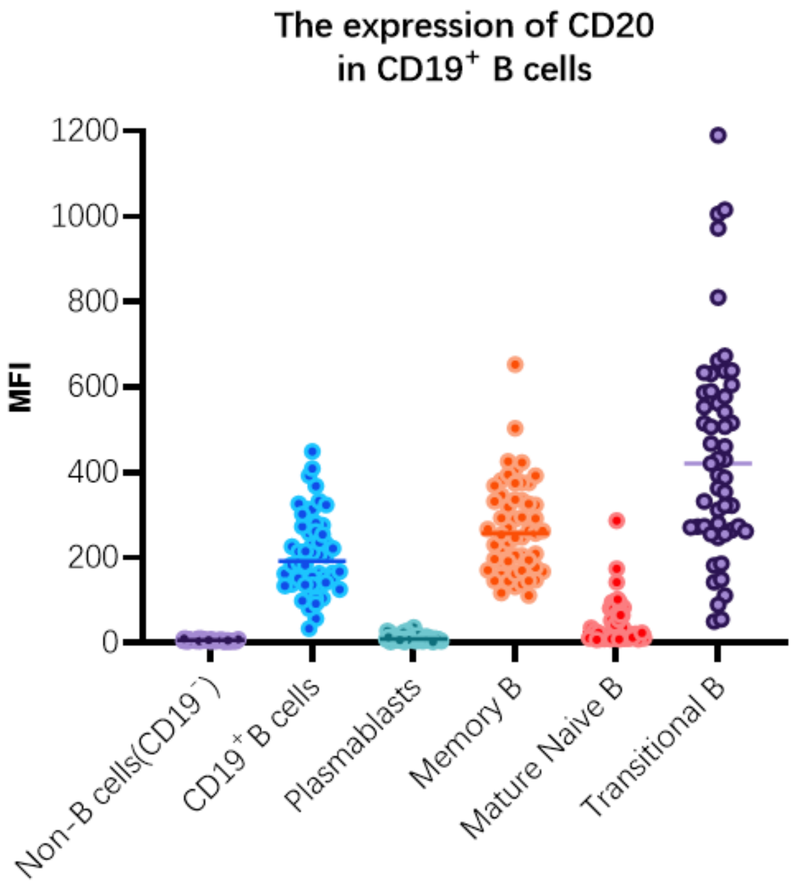


- Given its potent B-cell depletion ability, amulirafusp alfa (IMM0306) —a dual-targeting therapy against CD20 and CD47—shows strong potential as a promising treatment for autoimmune diseases.

Gallagher et al. Mol Immunol. 2017 May:85:57-65.

Dual Targeting of CD20 and CD47 Enhances Cell Lineage Coverage, Improving Therapeutic Potential





- In vitro analysis of SLE patient blood revealed:
  - CD47: High expression across B-cell subtypes (including plasmablasts), with no significant variation.
  - CD20: Minimal expression in plasmablasts and mature naïve B-cells vs other B-cell subsets.

Development Plan of Amulirafusp alfa (IMM0306) in Autoimmune Diseases

IND Approved in China

Systemic lupus erythematosus (SLE)
Phase Ib

Preliminary results presented here

Neuromyelitis optica spectrum disorder (NMOSD)
Phase Ib

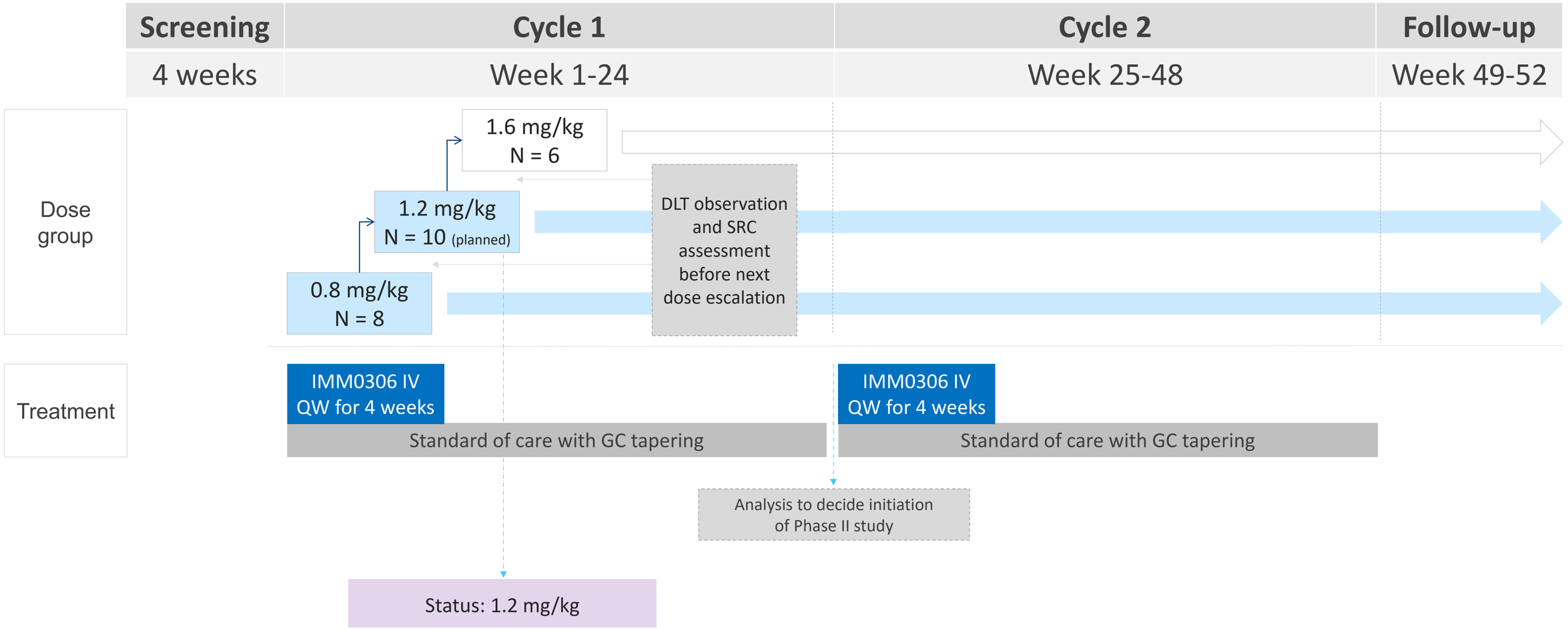
Lupus nephritis (LN)
Phase II

IND planned in US & China

Multiple sclerosis (MS)
China: Phase II
US: Phase Ib/II

Myasthenia gravis (MG)
China: Phase II
US: Phase Ib/II

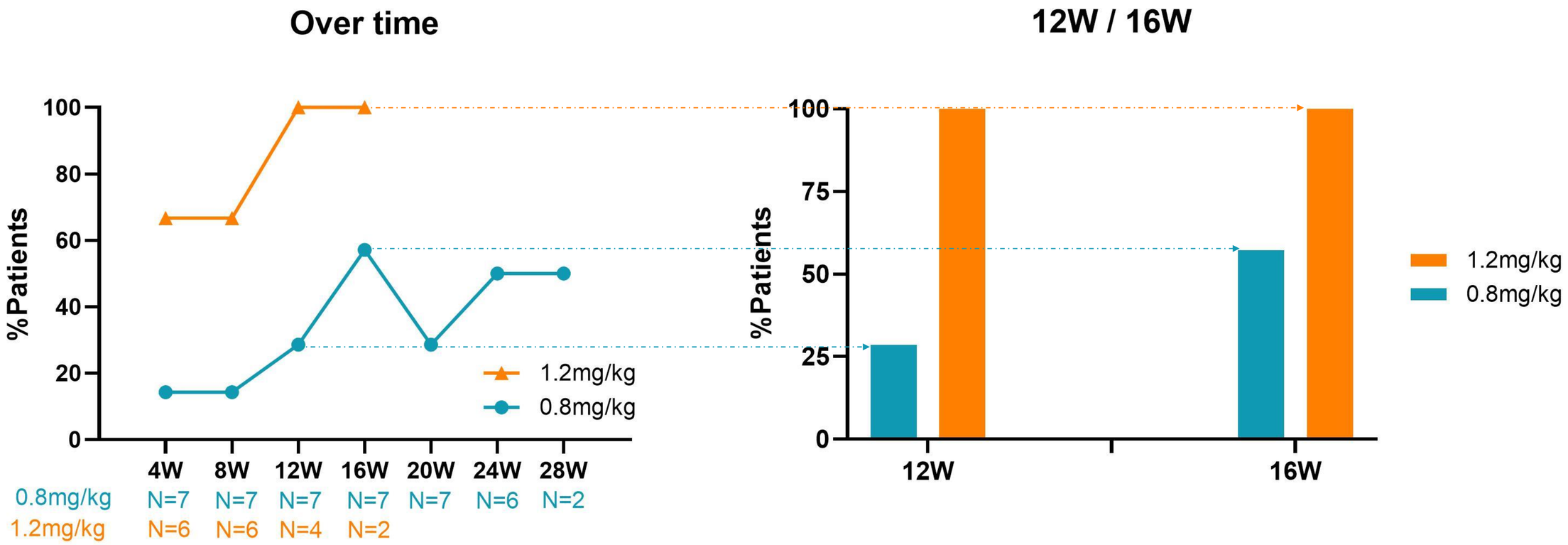
Amulirafusp alfa (IMM0306) - Phase Ib Trial Design in SLE



GC: glucocorticoids. QW: Once a week. DLT: dose limiting toxicity. SRC: safety review committee.

Amulirafusp alfa (IMM0306) Shows Rapid, Dose-Dependent SLEDAI-2K Improvement

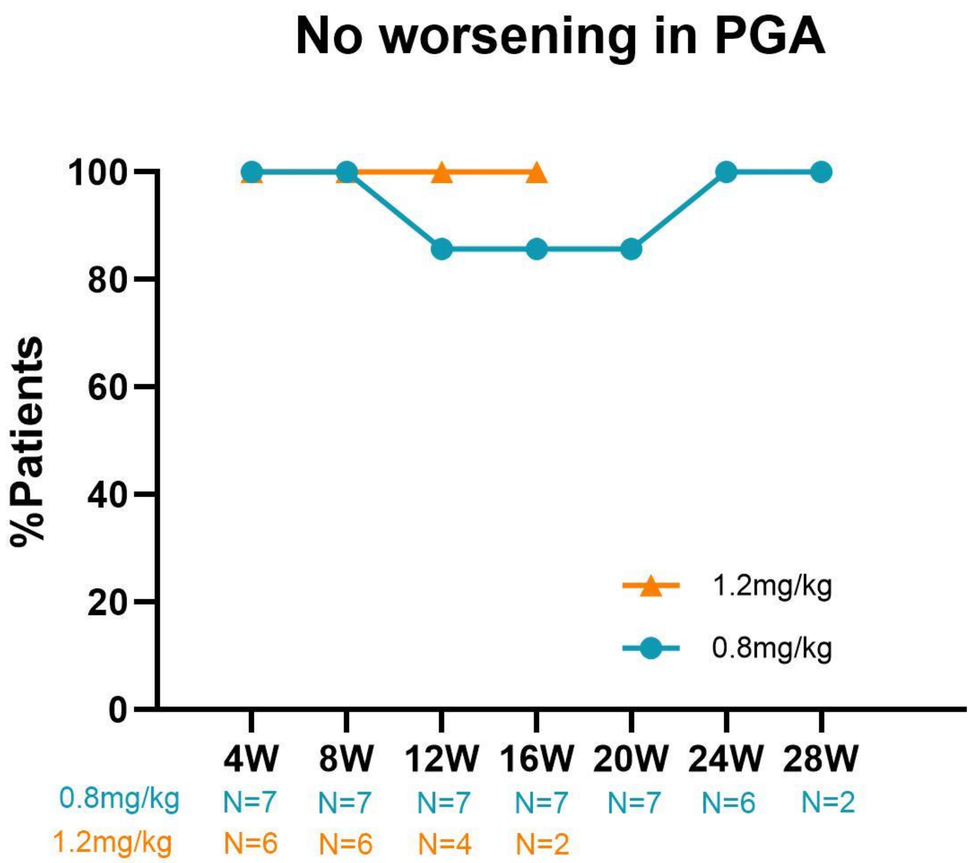
- Proportion of patients with ≥ 4 points reduction from baseline in SLEDAI-2K score



Data cut-off June 6, 2025.

Note: The patients included in the efficacy analysis had completed ≥4 doses and at least one efficacy evaluation (7 patients in 0.8mg/kg cohort, 6 patients in 1.2mg/kg cohort).

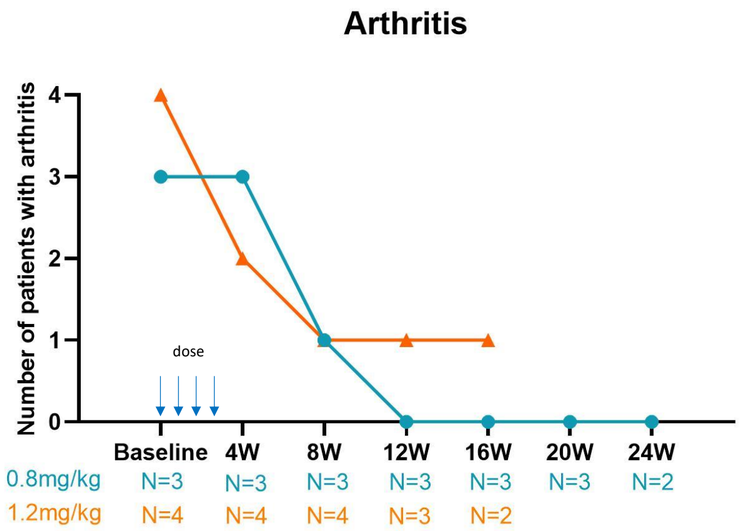
Strong Efficacy Signal in Preliminary Data of SLE Study

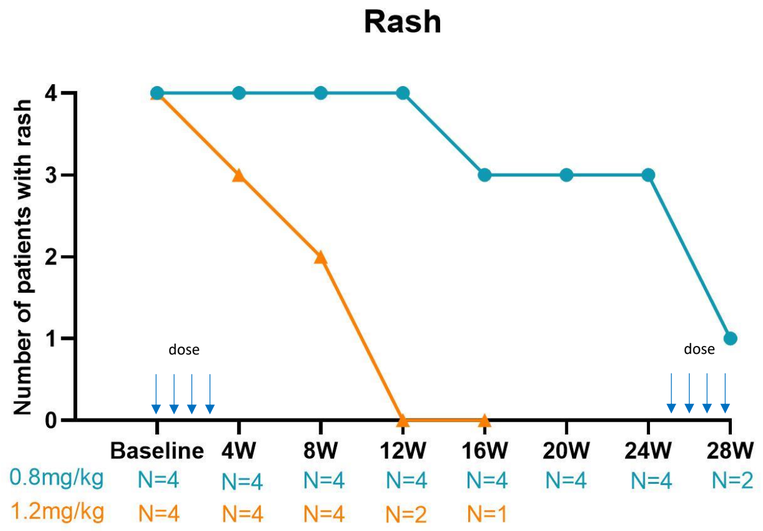


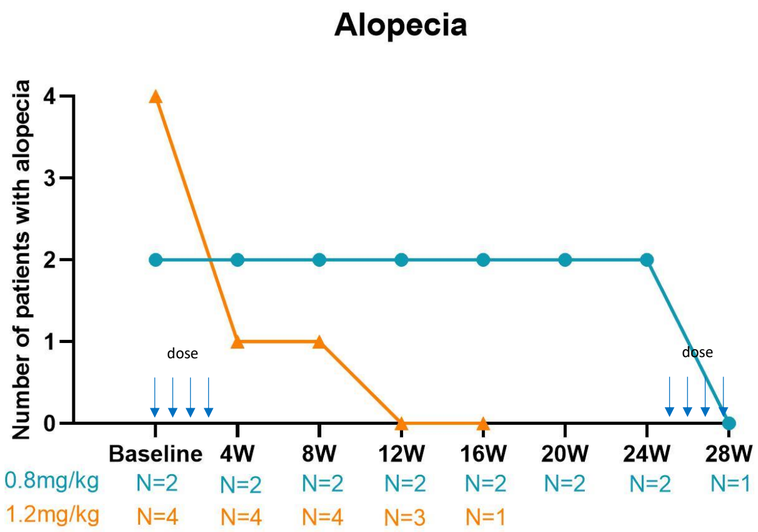
Data cut-off June 6, 2025.

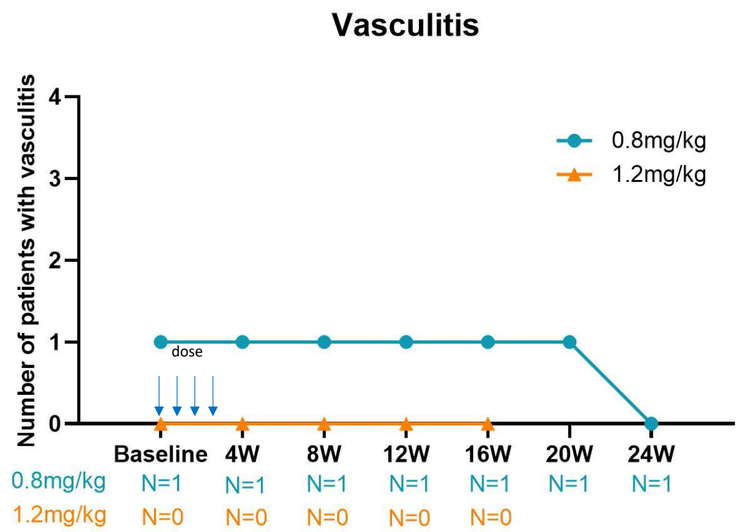
Note: The patients included in the efficacy analysis had completed ≥4 doses and at least one efficacy evaluation (7 patients in 0.8mg/kg cohort, 6 patients in 1.2mg/kg cohort).

Situation of Arthritis, Rash, Alopecia and Vasculitis are Improved



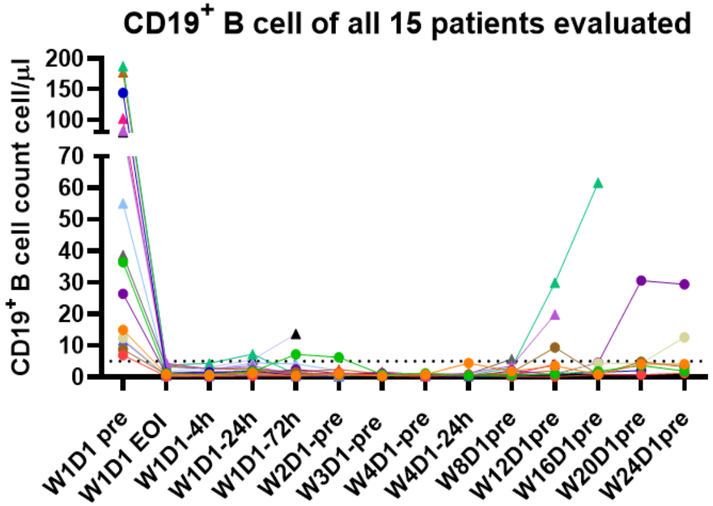




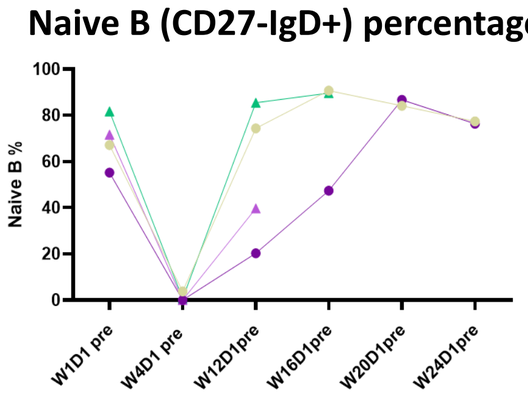


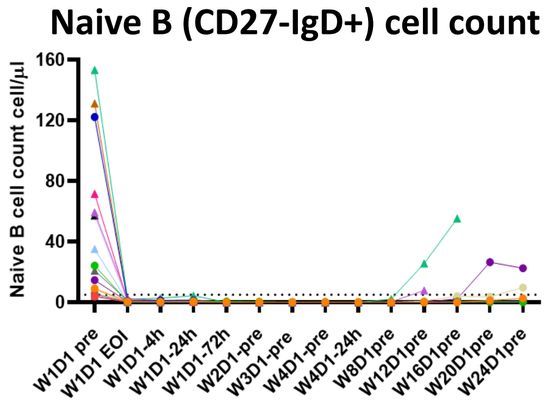
Data cut-off June 6, 2025.

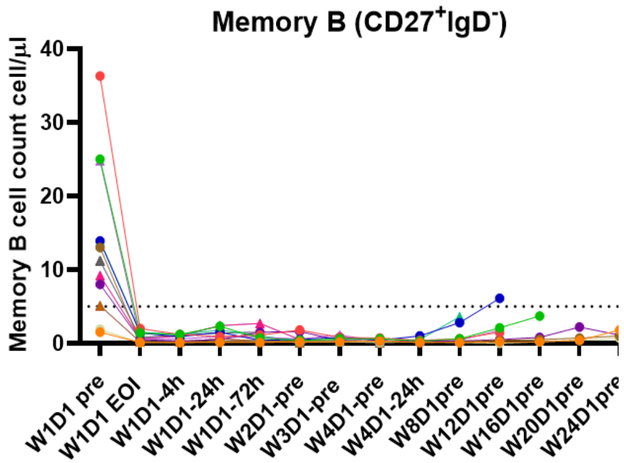
Efficient and Sustained B-cell Depletion with Immune Reconstitution Observed

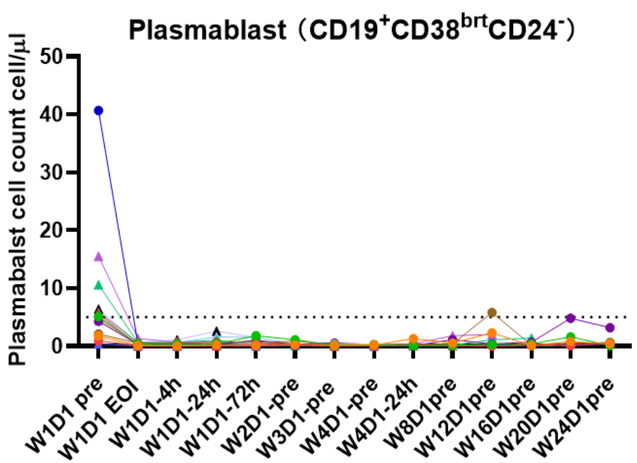


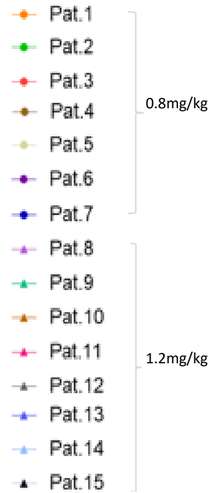
4 patients showed a trend of immune reconstitution from W12











Amulirafusp alfa (IMM0306) Shows Best-in-disease Potential in SLE

	Amulirafusp alfa (IMM0306)	Mosunetuzumab[2]	Telitacicept[3]	Belimumab[4]
Target	CD47xCD20	CD3xCD20	BLyS, APRIL	BLyS
≥4 points reduction from baseline in patients with SLEDAI-2K ≥8	83.3% (5/6) Week8-16 [1]	66.7% (4/6) Week52	77.8% (49/63) Week48[3.1]	46.5% (127/273) Week52[4.1]
B-cell depletion right after infusion	Yes	n.a.	n.a.	n.a.
Cytokine release syndrome	0	26.7% (4/15)	n.a.	n.a.
Dose step-up	Not required	Required	Not required	Not required
Stage	Phase Ib	Phase Ib	Approved in China	Approved by FDA

n.a. not available

1. 1.2 mg/kg. 2. Chindalore et al. EULAR2025 POS1160. 3. Wu et al. Ann Rheum Dis 2023;0:1–13. BLyS: B lymphocyte stimulator; APRIL: a proliferation inducing ligand. 4. Furie et al. Arthritis Rheum. 2011 Dec;63(12):3918-30.

3.1 Approved dose (160 mg). 4.1 Approved dose (10mg/kg), base line SLEDAI score ≥ 6.

Thank you

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