

PRODUCT

Therapeutic antibody targeting complement component 6 (C6)

INDICATION

Membrane Attack Complex (MAC)-mediated diseases including intravascular hemolysis

VALUE PROPOSITION

- More selective targeting over competitive therapeutics
- Potential for accelerated development and regulatory pathways
- Experienced biologic therapeutic development team

DEVELOPMENT STAGE

Antibody fully humanized and prepared for IND-enabling studies

INTELLECTUAL PROPERTY

[PCT/US2023/077377](https://pubmed.ncbi.nlm.nih.gov/38598870/)

RELATED PUBLICATIONS

Zhang L, ... **Lin F**. Mol Immunol. 2024 Jun;170:19-25.
[PMID: 38598870](https://pubmed.ncbi.nlm.nih.gov/38598870/)

Lin K, ... **Lin F**. Blood Adv. 2020 May 12;4(9):2049-2057.
[PMID: 32396613](https://pubmed.ncbi.nlm.nih.gov/32396613/)

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Anti-C6 Antibody Therapy for Complement Mediated Diseases

Feng Lin, PhD – Cleveland Clinic Lerner Research Institute

OPPORTUNITY

- Membrane attack complexes assembled after complement activation mediate various diseases.
- Eculizumab, a monoclonal antibody against complement component C5, is being used therapeutically to treat MAC-mediated disease.
- However, C5 is not a selective target for MAC inhibition, and some patients respond incompletely or not at all to the eculizumab treatment.
- C6, the next essential component in the terminal pathway of complement activation, may be an alternative target for the selective inhibition of MAC formation.

SOLUTION

- Feng Lin, PhD, a leader in the complement biology field, has developed a pipeline of antibody therapies target different proteins in the complement cascade.
- An anti-human C6 mAb demonstrated recognition of C6 both in free circulation and within C5b6 complexes.
- The candidate mAb has also demonstrated reduced human intravascular hemolysis in vivo in a preclinical animal model.
- The mAb has been fully humanized through collaboration with LifeArc and is prepared for IND-enabling studies

