

Protein A immunoAdsorption for the Treatment of Acute Episodes of Neuromyelitis Optica Spectrum Disorder

NCT06763848

Status	RECRUITING
Phase	Not Applicable
Sponsor	Third Affiliated Hospital, Sun Yat-Sen University
Enrollment	144 participants

Key Eligibility Criteria

Inclusion (4)

- clinical diagnosis of acute Neuromyelitis Optica Spectrum Disorders (NMOSD)
- Age and Gender: Participants must be between 18 and 65 years old, inclusive, with no gender restrictions.
- Serological Marker: Participants must test positive for AQP4-IgG using the cell-based assay (CBA) method.
- Understanding and Consent: Participants or their legal representatives must be able to understand the study's purpose, demonstrate sufficient compliance with the study protocol, and sign the informed consent form.

Exclusion (8)

- Women who are pregnant or breastfeeding.
- Participants who cannot establish peripheral or central venous access, or have a history of allergic reactions to plasmapheresis.
- Participants with contraindications to intravenous methylprednisolone treatment.
- Participants who have used monoclonal antibodies in the last 6 months, or FcRn antagonists in the last 3 months.
- Participants who must use ACE inhibitors (ACEI) within 1 week before the start of treatment or during the study, and cannot discontinue their use.

... and 3 more (see full listing online)

Locations (1 total)

The Third Affiliated Hospital of Sun Yat-sen University, Guangzhou, Guangdong, China