

POLICY NAME

Enspryng (satralizumab)

POLICY #

2794P

Criteria

Exclusion Criteria – Any of the following prevents coverage

- ☐ 2.1 Enspryng will not be approved for use in combination with Uplizna or Soliris

Coverage Criteria

- ☐ 1.1 Documented diagnosis of neuromyelitis optica spectrum disorder (NMOSD) with chart notes indicating the member exhibits at least one of the core clinical characteristics:
- Optic neuritis (inflammation of optic nerve)
 - Acute myelitis (a type of inflammation of the spinal cord)
 - Area postrema syndrome (episode of otherwise unexplained hiccups or nausea and vomiting)
 - Acute brainstem syndrome (lesions of the brain stem causing symptoms such as dizziness, vertigo, headache, facial pain, vision disturbances)
 - Symptomatic narcolepsy or acute diencephalic clinical syndrome with NMOSD-typical diencephalic MRI lesions (resulting from a rare type of central nervous system lesion)
 - Symptomatic cerebral syndrome with NMOSD-typical brain lesions
- ☐ 1.2 Documentation that the patient is anti-aquaporin-4 (AQP4) antibody positive
- ☐ 1.3 Ordered by a neuro-ophthalmologist or specialist in the treatment of NMOSD
- ☐ 1.4 Documentation that the member has been on a stable dose of immunosuppressive therapy (i.e., azathioprine, mycophenolate mofetil, oral corticosteroids, etc.)
- ☐ 1.5 Review of chart notes documenting diagnosis and confirming that patient has met all of the above requirements for treatment with Enspryng by both a pharmacist and a medical director