

PATIENT INFORMATION
PATIENT NAME: _____ **PHONE:** _____

DATE OF BIRTH: _____ **SEX:** ☐ M ☐ F **HEIGHT:** _____ **WEIGHT:** _____ ☐ LBS ☐ KG

ALLERGIES: _____ **PREFERRED CLINIC:** _____

REFERRAL STATUS: ☐ NEW REFERRAL ☐ ORDER CHANGE ☐ ORDER RENEWAL

DIAGNOSIS & CLINICAL DOCUMENTATION
☐ G70.00 Myasthenia gravis w/o acute exacerbation (gMG)

☐ G70.01 Myasthenia gravis with acute exacerbation (gMG)

☐ ICD-10 CODE: _____ DESCRIPTION: _____

REQUIRED DOCUMENTATION
☐ Insurance Information

☐ List of Medications

☐ Tried & Failed Therapies

☐ Most Recent History & Physical

☐ MG-ADL Score / MGFA Classification

☐ Positive AChR Antibody (gMG)

MEDICATION ORDER
☐ **CYCLE:** VYVGART® (efgartigimod alfa) 10 mg/kg IV in 125 mL NS over 60 minutes once weekly for 4 weeks (max dose of 1200 mg for patients weighing ≥ 120 kg)

*Flush with 20 mL NS after each infusion

☐ Repeat cycle _____ week(s) from date of last infusion; patient to receive a total of _____ cycles

PRE-MEDICATIONS
PO
☐ Acetaminophen: 650 mg

☐ Cetirizine: 10 mg

☐ Diphenhydramine: 25 mg

IV
☐ Methylprednisolone: 125 mg

☐ Diphenhydramine: 25 mg

☐ **OTHER:** _____ ☐ PO ☐ IV

Patient to be observed for 30 minutes following each infusion.

In the event of an adverse reaction occurring in the infusion clinic, utilize the Immersiv Health adverse reaction protocol.

PRESCRIBER INFORMATION
PROVIDER NAME: _____ **NPI #:** _____

EMAIL: _____ **PHONE:** _____ **FAX:** _____

ADDRESS (INCLUDE CITY, STATE, ZIP): _____

SUPERVISING PHYSICIAN: _____ **CONTACT NAME:** _____
(IF APPLICABLE)
SIGNATURE: _____ **DATE:** _____
(NO STAMPS)

SUBSTITUTION PERMITTED

DISPENSE AS WRITTEN