

# Neurology order form for biologic therapies

Infusion Pharmacy Phone: \_\_\_\_\_ Fax: \_\_\_\_\_

Page 1 of 4

Care specialist Name: \_\_\_\_\_ Phone: \_\_\_\_\_

## Patient information see attached PEDIATRIC (younger than 13 years or less than 45 kg in weight)

Patient first name: \_\_\_\_\_ Middle: \_\_\_\_\_ Last: \_\_\_\_\_

Gender:  M  F DOB: \_\_\_\_\_ Last 4 of SSN: \_\_\_\_\_

Address: \_\_\_\_\_ City: \_\_\_\_\_ State: \_\_\_\_\_ ZIP: \_\_\_\_\_

Phone: \_\_\_\_\_  Home  Work  Cell Alternate Phone: \_\_\_\_\_  Home  Work  Cell

Emergency contact: \_\_\_\_\_ Phone: \_\_\_\_\_ Relationship: \_\_\_\_\_

**Insurance:** Please attach copy of the front and back of insurance card(s).

Primary Insurance: \_\_\_\_\_ Phone: \_\_\_\_\_ Policy #: \_\_\_\_\_ Group: \_\_\_\_\_

Secondary Insurance: \_\_\_\_\_ Phone: \_\_\_\_\_ Policy #: \_\_\_\_\_ Group: \_\_\_\_\_

## Medical assessment

**Primary diagnosis:**  ICD10 Code: \_\_\_\_\_ Diagnosis: \_\_\_\_\_

**Weight in kg only:** \_\_\_\_\_ Date weight (in kg) obtained: \_\_\_\_\_

**Allergies:** \_\_\_\_\_

### Required documentation:

- Include with this order supportive documents of tried and failed therapies, a list of current medications, vaccination history, lab results, and clinical notes as relevant to this therapy.

- For Soliris and Ultomiris therapy, include documentation of meningococcal immunization dates, or clinical notes as to rationale for not completing, the plan for vaccination and the antibacterial drug regimen the patient will be taking until fully immunized.

**Prescriber opts to proceed with therapy before the patient receives full meningococcal vaccination.**

Selecting this box confirms the prescriber considers the risks to this patient of postponing start of care until at least 2 weeks following full immunization against meningococcal serotypes A, C, W, Y, and B per ACIP guidance as outweighing the risk of serious infection. The prescriber is requesting start of therapy to proceed. The prescriber has counseled the patient on the risks and is providing the patient with a prophylactic antibacterial drug regimen.

**Patient requires a first lifetime dose and is to receive the first dose in the home or Ambulatory Infusion Suite.**

**IV access** (if IV therapy is prescribed):  PIV  PICC  Port  Midline  Tunneled CVC; number of lumens: \_\_\_\_\_

Date of IV placement: \_\_\_\_\_ Date of last IV service (flush and/or dressing change): \_\_\_\_\_

## Medication prescriptions and orders

Medication	Dose and directions (Select or enter desired dose regimens.)
<b>Soliris</b> (eculizumab)	<input type="checkbox"/> INDUCTION: Soliris 600 mg IV once weekly for 4 weeks, then 900 mg for the 5th dose one week later, then 900 mg every 2 weeks thereafter.
	<input type="checkbox"/> INDUCTION: Soliris 900 mg IV once weekly for 4 weeks, then 1,200 mg for the 5th dose one week later, then 1,200 mg every 2 weeks thereafter.
	<input type="checkbox"/> INDUCTION: Soliris, Other (specify): _____
	<input type="checkbox"/> MAINTENANCE: Soliris <input type="checkbox"/> 900 mg <input type="checkbox"/> 1,200 mg <input type="checkbox"/> other dose _____ IV once every 2 weeks or indicate here if other frequency _____
	<input type="checkbox"/> SUPPLEMENTAL: Soliris dose and timing _____
Refills x 1 year	Prior to administration dilute dose in 0.9% Sodium Chloride to a final concentration of 5 mg / mL. For adults: Administer as an intravenous (IV) infusion over at least 35 minutes, but not to exceed 2 hours. For pediatrics: Administer IV over a 1 - 4 hour period, as tolerated. RN to monitor patient at minimum 60 minutes post infusion and to take a final set of vital signs before concluding visit.

**This form is not a valid prescription in New York.**

Continued on page 2 ↓

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Infusion Pharmacy Phone: \_\_\_\_\_ Fax: \_\_\_\_\_

Page 2 of 4

Patient first name: \_\_\_\_\_ Middle: \_\_\_\_\_ Last: \_\_\_\_\_ DOB: \_\_\_\_\_

## Medication prescriptions and orders

Medication	Dose and directions (Select or enter desired dose regimens.)
<p><b>Ultomiris</b> (ravulizumab-cwvz)</p> <p>Refills x 1 year</p>	<p><input type="checkbox"/> <b>LOADING DOSE</b> based on patient weight following the below specifications:</p> <p>Patient weight 40 kg to 59 kg: Ultomiris 2,400 mg in 0.9% Sodium Chloride for a final volume of 48 mL. Administer IV over no less than 48 minutes as tolerated by the patient at a rate not to exceed 60 mL / hr x 1 dose.</p> <p>Patient weight 60 kg to 99 kg: Ultomiris 2,700 mg in 0.9% Sodium Chloride for a final volume of 54 mL. Administer IV over no less than 36 minutes as tolerated by the patient at a rate not to exceed 90 mL / hr x 1 dose.</p> <p>Patient weight 100 kg or more: Ultomiris 3,000 mg in 0.9% Sodium Chloride for a final volume of 60 mL. Administer IV over no less than 24 minutes as tolerated by the patient at a rate not to exceed 150 mL / hr x 1 dose.</p> <p><input type="checkbox"/> <b>MAINTENANCE DOSE</b> based on patient weight following the below specifications. Administer IV 2 weeks following the Loading Dose and every 8 weeks, or indicate here if other frequency. _____</p> <p>Patient weight 40 kg to 59 kg: Ultomiris 3,000 mg in 0.9% Sodium Chloride for a final volume of 60 mL. Administer IV over no less than 54 minutes and at a rate no more than 67 mL / hr, as tolerated by the patient.</p> <p>Patient weight 60 kg to 99 kg: Ultomiris 3,300 mg in 0.9% Sodium Chloride for a final volume of 66 mL. Administer IV over no less than 42 minutes and at a rate no more than 95 mL / hr, as tolerated by the patient.</p> <p>Patient weight 100 kg or more: Ultomiris 3,600 mg in 0.9% Sodium Chloride for a final volume of 72 mL. Administer IV over no less than 30 minutes and at a rate no more than 144 mL / hr, as tolerated by the patient.</p> <p><input type="checkbox"/> SUPPLEMENTAL, Ultomiris dose and timing: _____</p> <p><input type="checkbox"/> OTHER, Ultomiris dose and schedule: _____</p> <p>_____</p> <p>RN to monitor patient at minimum for 60 minutes post infusion and to take a final set of vital signs before concluding visit.</p>
<p><b>Vyvgart</b> (efgartigimod alfa)</p> <p><b>For intravenous (IV) infusion</b></p> <p>Refills x 1 year.</p>	<p><input type="checkbox"/> Vyvgart dose based on patient weight following the below specifications.</p> <p>Patient weight 120 kg or less: Vyvgart dose is 10 mg / kg. Dose will be rounded to the nearest vial size available.</p> <p>Patients weight over 120 kg: Vyvgart dose is 1,200 mg.</p> <p>Dilute Vyvgart dose in 0.9% Sodium Chloride to a final volume of 125 mL.</p> <p>Administer as an intravenous (IV) infusion over 1 hour.</p> <p>Administration Schedule (select one):</p> <p><input type="checkbox"/> Administer Vyvgart dose IV once weekly x 4 weeks to complete a 28-day cycle. Repeat cycle beginning 50 days after the first dose of the previous cycle.</p> <p><input type="checkbox"/> Other _____</p> <p>RN to monitor patient at minimum for 60 minutes post infusion and to take a final set of vital signs before concluding visit.</p>
<p><b>Vyvgart Hytrulo</b> (efgartigimod alfa and hyaluronidase)</p> <p><b>For subcutaneous (SC) injection</b></p> <p>Refills x 1 year.</p>	<p><input type="checkbox"/> Vyvgart Hytrulo 1,008 mg and 11,200 units / 5.6 mL. RN to administer by slow subcutaneous push over 30 - 60 seconds using a single lumen needle set for subcutaneous infusion per specifications in product prescribing information.</p> <p>Administration schedule (select one):</p> <p><input type="checkbox"/> Administer Vyvgart Hytrulo SC once weekly x 4 weeks to complete a 28-day cycle. Repeat cycle beginning 50 days after the first dose of the previous cycle.</p> <p><input type="checkbox"/> Other _____</p> <p>RN to monitor patient at minimum for 30 minutes following administration and to take a final set of vital signs before concluding visit.</p>

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Infusion Pharmacy Phone: \_\_\_\_\_ Fax: \_\_\_\_\_

Page 3 of 4

Patient first name: \_\_\_\_\_ Middle: \_\_\_\_\_ Last: \_\_\_\_\_ DOB: \_\_\_\_\_

## Ancillary prescriptions and orders

Premedication (select below): Dispense PRN x 1 year.

	Drug	Patient Type	Dose	Dispense detail	Directions
<input type="checkbox"/>	DiphenhydrAMINE	Adult & Pediatric > 30 kg	50 mg (two 25 mg capsules or tablets)	Dispense 25 mg capsules or tablets #100	Administer orally 30 minutes prior to Biologic medication. May repeat once if symptoms occur.
		Pediatric 15 - 30 kg	25 mg (10 mL)	Dispense 2.5 mg / mL oral solution #120 mL (300 mg)	
		Pediatric < 15 kg	12.5 mg (5 mL)	Dispense 2.5 mg / mL oral solution #120 mL (300 mg)	
<input type="checkbox"/>	Acetaminophen	Adult & Pediatric > 30 kg	325 mg	Dispense 325 mg tablets or 325 mg (10.15 mL) unit dose oral solution #100.	Administer orally 30 minutes prior to Biologic medication. May repeat once if symptoms occur.
		Pediatric 15 - 30 kg	160 mg (5 mL)	Dispense 160 mg (5 mL) tablets #30 or 32 mg / mL oral solution 120 mL.	
		Pediatric < 15 kg	80 mg (2.5 mL)	Dispense 32 mg / mL oral solution 120 mL.	
<input type="checkbox"/>	Other, specify _____	_____	_____	_____	_____

### Lab Orders, x 1 year

CBC w/differential  CMP  Creatinine / BUN  CRP  ESR  Other \_\_\_\_\_  
 Frequency of labs: \_\_\_\_\_

Lab work to be obtained via IV access using aseptic technique. If RN is not able to draw labs from a central catheter, the labs may be drawn peripherally. RN to flush IV access after each blood draw with 0.9% Sodium Chloride 20 mL. As a final lock for patency, use 5 mL of heparin 10 units / mL, or if for a port use 5 mL of heparin 100 units / mL.

### Nursing Orders, x 1 year

RN to complete assessment and administer prescribed medication in the home or Ambulatory Infusion Suite. For therapies infused IV, RN to insert, maintain, and/or remove peripheral IV (PIVC) or access central venous catheter (CVC) as needed using aseptic technique. RN to rotate PIVC as needed for signs of infiltration or irritation. Flush catheter with 5 mL of 0.9% Sodium Chloride pre infusion and post infusion. Lock IV access with 3 mL of heparin 10 units / mL to maintain patency.

If port, RN to access with non-coring port needle using sterile technique. De-access after infusion and apply pressure with sterile gauze. Apply transparent dressing to site. RN to use 10 mL of sterile field 0.9% Sodium Chloride with needle change. To maintain catheter patency, following the post infusion flush, use 5 mL of heparin 100 units / mL. Discontinue port maintenance upon discontinuation of pharmacy services.

### Pharmacy Orders, x 1 year

Pharmacy to dispense flushes, needles, syringes and HME / DME quantity sufficient to complete therapy as prescribed.

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Infusion Pharmacy Phone: \_\_\_\_\_ Fax: \_\_\_\_\_

Page 4 of 4

Patient first name: \_\_\_\_\_ Middle: \_\_\_\_\_ Last: \_\_\_\_\_ DOB: \_\_\_\_\_

**Anaphylaxis/infusion reaction management orders:** Dispense PRN x 1 year

Drug	Patient Type	Dose	Dispense detail	Directions
<b>DiphenhydrAMINE</b>	Adult & Pediatric > 30 kg	50 mg (two 25 mg capsules or tablets)	Dispense 25 mg capsules or tablets #4	For mild* symptoms, slow infusion by 50% until symptoms resolve. Administer diphenhydrAMINE orally.  For moderate* to severe* symptoms, stop infusion.  Administer diphenhydrAMINE slow IV push not to exceed rate of 25 mg / minute. May repeat once if symptoms persist. For moderate* symptoms that resolve, resume infusion at 50% of the previous rate.
		50 mg (1 mL) injection	Dispense 50 mg vial for injection #1	
	Pediatric 15 - 30 kg	25 mg (10 mL) orally	Dispense 25 mg / 10 mL oral solution 120 mL	
		25 mg (0.5 mL)	Dispense 50 mg vial for injection #1	
	Pediatric < 15 kg	12.5 mg (5 mL) orally	Dispense 12.5 mg / 5 mL oral solution 120 mL	
		12.5 mg (0.25 mL)	Dispense 50 mg vial for injection #1	
<b>EPINEPHrine</b>	Adult & Pediatric > 30 kg	0.3 mg (0.3 mL) injection	Dispense 1 mg vial for injection #2	For severe* symptoms (anaphylaxis), stop infusion. Disconnect tubing from access device to prevent further administration. Activate 911. Administer EPINEPHrine IM into lateral thigh once. May repeat in 5 - 15 minutes if symptoms persist. Administer CPR if needed until EMS arrives. Contact prescriber to communicate patient status.
	Pediatric 15 - 30 kg	0.15 mg (0.15 mL) injection	Dispense 1 mg vial for injection #2	
	Pediatric 7.5 - 15 kg	0.1 mg (0.1 mL) injection	Dispense Autoinjector Pen 0.1 mg (PED) #2	
<b>0.9% Sodium Chloride Injection, USP</b>	Dispense 500 mL bag #1. For severe* symptoms, administer IV gravity bolus (1,000 mL / hour).			
<b>Other, specify</b>	_____			

\*Mild symptoms include flushing, dizziness, headache, apprehension, sweating, palpitations, nausea, pruritus, and/or throat itching.

Moderate symptoms include chest tightness, shortness of breath, >20 mmHg change in systolic blood pressure from baseline, and/or increase in temperature (>2°F).

Severe symptoms include >40 mmHg change in systolic blood pressure from baseline, increase in temperature with rigors, shortness of breath with wheezing, and/or stridor.

## Prescriber information

First Name: \_\_\_\_\_ Middle: \_\_\_\_\_ Last: \_\_\_\_\_ Practice: \_\_\_\_\_

Address: \_\_\_\_\_ City: \_\_\_\_\_ State: \_\_\_\_\_ ZIP: \_\_\_\_\_

Phone: \_\_\_\_\_ Fax: \_\_\_\_\_ NPI: \_\_\_\_\_ Contact: \_\_\_\_\_

By signing, I certify/recertify that the above therapy, products and services are medically necessary and that this patient is under my care. I have received authorization to release the above referenced information and medical and/or patient information relating to this therapy. Pharmacy has my permission to contact the insurance company on my behalf to obtain authorization for patient.

Substitution permissible signature OR Dispense as written signature Date

**Please fax:**  Completed form  Demographic sheet/insurance information  Clinical notes and labs

Please include ALL 4 pages of referral form and additional documentation when faxing.

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