



Patient Name: \_\_\_\_\_

DOB: \_\_\_\_\_

## Ocrelizumab (Ocrevus®) Infusion Orders

Diagnosis (please provide ICD-10 code in space provided):

\_\_\_\_\_ Multiple Sclerosis  
(ICD-10)

\_\_\_\_\_ Other: \_\_\_\_\_  
(ICD-10)

- Hold infusion and notify provider for:
  - Signs/symptoms of infection, planned/recent surgical procedures, recent live vaccines, chance of pregnancy or new/worsening neurological symptoms.
- If infusion-related reaction occurs, stop infusion and initiate Hypersensitivity Reaction Management Protocol as clinically indicated.

**Pre-medications** (to be administered once *30 minutes prior* to infusion):

- Tylenol 1000 mg PO     
  Solu-medrol 125 mg IVP     
  Benadryl 50 mg IV  
 Loratadine 10 mg PO     
  Famotidine 20 mg PO     
  Other: \_\_\_\_\_

**Lab Orders:**

Induction <input type="checkbox"/>	Administer Ocrevus 300 mg in 250 ml 0.9% normal saline on Week 0 and Week 2. Administer Ocrevus 600 mg in 500 ml 0.9% normal saline 6 months after initial dose.
OR	
Maintenance <input type="checkbox"/>	Administer Ocrevus 600 mg in 500 ml 0.9% normal saline every 6 months

**Administration:**

- Use **0.2 or 0.22 micron in-line filter**
- Monitor **vital signs every 30 minutes**
- Titrate infusion rates as follows:

Hour	300 mg Infusion (duration at least 2.5 hours)	600 mg infusions (duration at least 3.5 hours)
0	30 ml/hr	40 ml/hr
0.5	60 ml/hr	80 ml/hr
1	90 ml/hr	120 ml/hr
1.5	120 ml/hr	160 ml/hr
2	150 ml/hr	200 ml/hr
2.5	180 ml/hr	No change
3	n/a (complete)	
3.5		

**Observation Period:**

- Monitor patient for hypersensitivity reaction for a period of **60 minutes following each infusion.**
- Record vital signs prior to discharge.

Provider (please print): \_\_\_\_\_

Provider signature: \_\_\_\_\_ Date: \_\_\_\_\_