

# Multiple Sclerosis IV/SC Infusion REFERRAL FORM

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## 1 PATIENT'S INFORMATION (Complete or include demographic sheet)

Patient Name: \_\_\_\_\_ DOB: \_\_\_\_\_ Gender: ☐ MALE ☐ FEMALE  
Address: \_\_\_\_\_ City/State/Zip: \_\_\_\_\_  
Primary Phone: \_\_\_\_\_ Email: \_\_\_\_\_ SSN: \_\_\_\_\_ Primary Language: \_\_\_\_\_  
Emergency Contact: \_\_\_\_\_ Relationship to Patient: \_\_\_\_\_ Phone: \_\_\_\_\_

## 2 INSURANCE (Please attach If Available a copy of the Patients' Insurance card(s) Front/ Back)

Primary Insurance: \_\_\_\_\_ Policy No: \_\_\_\_\_ Group: \_\_\_\_\_  
Primary RX Insurance: \_\_\_\_\_ ID No: \_\_\_\_\_ RX GRP: \_\_\_\_\_ RX BIN: \_\_\_\_\_ RX PCN: \_\_\_\_\_

## 3 DIAGNOSIS AND CLINICAL INFORMATION

### Diagnosis (ICD-10) and Patient Clinical Information:

Drug Allergies: \_\_\_\_\_ ☐ NKDA Ht: \_\_\_\_\_ in/cm Wt: \_\_\_\_\_ lbs/Kg

Diagnosis (ICD-10): ☐ G35 Multiple Sclerosis (MS) ☐ Other: \_\_\_\_\_ Description: \_\_\_\_\_

If MS, please

indicate type: ☐ Primary progressive MS (PPMS) ☐ Relapsing-remitting MS (RRMS) ☐ Progressive-relapsing MS (PRMS)  
☐ Secondary progressive MS (SPMS); If SPMS, does the patient have documented relapses? ☐ YES ☐ NO  
☐ Secondary progressive MS (SPMS); If SPMS, does the patient have documented relapses? ☐ YES ☐ NO

MS drug(s) Tried and Failed: Drug: \_\_\_\_\_ ☐ Inadequate response, trial duration \_\_\_\_\_  
Drug: \_\_\_\_\_ ☐ Intolerance, specify: \_\_\_\_\_  
Drug: \_\_\_\_\_ ☐ Inadequate response, trial duration \_\_\_\_\_  
Drug: \_\_\_\_\_ ☐ Intolerance, specify: \_\_\_\_\_

If Applicable, please indicate Pregnancy Results: ☐ POSITIVE (+) ☐ NEGATIVE (-)

Labs: Please fax copy of the following labs dated within 1 year: ☐ CMP ☐ CBCw/ diff ☐ IgG/ IgG Subclasses ☐ Hep-B ☐ MRI

## 4 PRESCRIPTION INFORMATION

MEDICATION	STRENGTH	DOSE & DIRECTION	QUANTITY	REFILLS
<input type="checkbox"/> Briumvi	150 mg/6 mL vial	Administer 150 mg in 250 mL NS IV over 4 hours minimum. For second and subsequent infusion; administer 450 mg IV in 250 mL NS over 1 hour minimum at week 2, Then repeat every 24 Weeks, following day 1 starting dose.	Quantity: <b>Q.S</b>	Refills: _____
<input type="checkbox"/> Ocrevus	300 mg/10 mL	Infuse 300 mg in 250 mL NS IV over 2.5 hours minimum at weeks 0 and 2. Then Infuse 600 mg in 500 mL NS IV over 2 hours minimum every 6 months.	Quantity: <b>Q.S</b>	Refills: _____
<input type="checkbox"/> Ocrevus Zunovo	920 mg ocrelizumab + 23,000U hyaluronidase /23 mL	Administer 23 mL of OCREVUS ZUNOVO subcutaneously in the abdomen over approximately 10 minutes every 6 months	Quantity: <b>Q.S</b>	Refills: _____

### PRE/POST ORDERS:

MEDICATION	DIRECTION	QUANTITY	REFILLS
<input type="checkbox"/> Diphenhydramine: <input type="checkbox"/> PO <input type="checkbox"/> IV	Administer _____ mg 30-60 min Prior to Infusion. May repeat every _____ hours PRN.	Quantity: <b>Q.S</b>	Refills: <b>PRN</b>
<input type="checkbox"/> Acetaminophen (PO)	Administer _____ mg PO 30-60 min Prior to Infusion. May repeat every _____ hours PRN.	Quantity: <b>Q.S</b>	Refills: <b>PRN</b>

### ANAPHYLAXIS PROTOCOL:

MEDICATION	DIRECTION	QUANTITY	REFILLS
<input checked="" type="checkbox"/> EPINEPHRINE (vial or autoinjector)	Administer IM for severe anaphylactic reaction. May repeat in 5-15 minutes <input type="checkbox"/> 0.3 mg (Wt > 30 kg) <input type="checkbox"/> 0.15 mg (Wt 15-30 kg) <input type="checkbox"/> 0.1 mg (Wt 7.5-15 kg)	Quantity: (x1) Vial or (x2) Pen(s)	Refills: <b>PRN</b>
<input checked="" type="checkbox"/> DIPHENHYDRAMINE (50mg/mL)	Administer via IM or slow IV push for severe anaphylactic reaction. <input type="checkbox"/> 50 mg (Wt > 30 kg) <input type="checkbox"/> 25 mg (Wt 15-30 kg) <input type="checkbox"/> 12.5mg (Wt 7.5-15kg)	Quantity: (x1) Vial	Refills: <b>PRN</b>
<input checked="" type="checkbox"/> SODIUM CHLORIDE 0.9% (IV)	Infuse 500 mL IV as directed for severe anaphylactic reaction.	Quantity: <b>500 mL</b>	Refills: <b>PRN</b>

### NURSING/ LABS/ INFUSION SUPPLIES:

☒ **NURSING:** Nursing visits with each infusion to establish venous access, administer medication, assess and monitor patient, provide education, and complete lab draws.

☒ **INFUSION SUPPLIES:** Infusion supplies and infusion pump PRN for the administration and disposal of medication.

**FLUSHING PROTOCOL:** ☒ Sodium chloride 0.9%, Upt to 10 mL before/after medication, and/or PRN to maintain patency.

☐ Heparin: ☐ 10 Units/mL ☐ 100 Units/mL, \_\_\_\_\_ as final flush and/or PRN to maintain patency

☒ **LABS:** [Dx code: Z79.899]; Labs to be drawn by RN prior to infusion every \_\_\_\_\_ ☐ Week ☐ Month(s), as followed:

☐ CMP ☐ CBC w/ diff ☐ IgG Total ☐ IgG Subclasses 1-4 ☐ Hepatitis (B) ☐ Other: \_\_\_\_\_

## 5 PRESCRIBER INFORMATION

Prescriber Name: \_\_\_\_\_ Title: ☐ MD ☐ DO ☐ ND ☐ PA ☐ APRN NPI: \_\_\_\_\_

Address: \_\_\_\_\_ City/ State/ Zip: \_\_\_\_\_

Phone: \_\_\_\_\_ Fax: \_\_\_\_\_ Contact Person: \_\_\_\_\_

The information provided above is true and accurate to the best of my knowledge, with supporting documentation in the patient's medical record.

I authorize IV Solutions RX and its representatives to act as an agent to initiate and execute the insurance prior authorization process for this prescription and any future refills of the same prescription for the patient listed above. I understand that I can revoke the designation at any time by providing written notice to IV Solutions RX.

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Physicians signature \_\_\_\_\_

Date \_\_\_\_\_

**THANK YOU FOR YOU TRUSTING US IN YOUR PATIENTS SPECIALTY CARE**

**#welcome to our family**