

**FAX TO: 855.694.4656**

## 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

\*PLEASE COMPLETE ICD-10 FOR SPECIFIC DIAGNOSIS

- ☐ M05. \_\_\_\_ Rheumatoid arthritis w/ rheumatoid factor
- ☐ M06. \_\_\_\_ Rheumatoid arthritis w/o rheumatoid factor
- ☐ M31.30 Granulomatosis w/ Polyangiitis (GPA/Wegener's)
- ☐ M31.7 Microscopic Polyangiitis (MPA)
- ☐ ICD-10 CODE: \_\_\_\_\_ DESCRIPTION: \_\_\_\_\_

## REQUIRED DOCUMENTATION

- ☐ Insurance Information ☐ List of Medications ☐ Tried & Failed Therapies ☐ Most Recent History & Physical ☐ Full Hep B Panel

## MEDICATION ORDER

Rituxan® or biosimilar (Ruxience®, Riabni®, Truxima®) may be used according to payor guidelines

### FOR RA

- ☐ Rituximab 1000mg IV in 500ml NS per protocol day 0 and day 14 every 4 months (16 weeks)
- ☐ Rituximab 1000mg IV in 500ml NS per protocol day 0 and day 14 every 6 months (24 weeks)

### FOR GPA/MPA

- ☐ Rituximab 375mg/m<sup>2</sup> IV per protocol once weekly for 4 weeks
- ☐ OTHER: \_\_\_\_\_

### LAB ORDERS

LAB: \_\_\_\_\_ FREQUENCY: \_\_\_\_\_

☒ REFILL X 12 MONTHS UNLESS OTHERWISE NOTED HERE: \_\_\_\_\_

Patient to be observed for 60 minutes following the first 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