

Provis Patient No.										

MEDICATION ORDER FORM

Rituximab (Rituxan®)											
Patient's Surname	Given Name & Initials		Date of Birth / / ddmm yyyy								
Referring Physician			yyyy								
Patient's Height: cm	Cycle:										
Weight: kg											
BSA: m²											
Pre-Medication ☐ Acetaminophen 650 mg PO 30-60 minutes pre-rituximab ☐ Diphenhydramine 50 mg PO/IV 30-60 minutes pre-rituximab ☐ Other											
 Hydration/IV solution: NS TKVO on day 1 of each cycle Monitor vitals (BP, pulse, respiration, temperature) every 15 mins. For the 1st hour or until stable and then every hour until infusion completed. Have an adverse reaction kit available. Keep IV line in and observe Pt for 1 hr after end of infusion. If complications occur during infusion, observe patient for 2 hrs. after the end of infusion. If Pt experiences transient fevers or rigors during infusion, STOP infusion and observe. Inform physician and treat as ordered. Once stable, restart infusion at ONE-HALF the previous rate. 											
Medication prescribed:											
Rituximab mg (375 mg/m ²) IV in NS (1 mg/mL) weekly for 4 weeks											
Or Rituximabmg (375 mg/m² ormg/m²) IV in NS (1 mg/mL) qdays (Please circle or indicate desired dose) Or Rituximab (Rituxan®) 1000 mg IV in NS (1 mg/mL) on Days 1 and 15. (Scheduling may dictate slight variation from day15) Please note: If treatment administered on q 3 wkly schedule a new medical order form is required each cycle.											
Scheduling (For Provis Use Only	<u>'</u>)										
Tx 1:		Tx 2:									
Tx 3:	(if necessary)	Tx 4:	(if necessary)								
Referring Physician's Signature		/	/ /								
Signature of Provis Physician		/	/ /								
Fax completed form to: 416-532-3635											



Information for Physicians regarding Rituxan® (Rituximab) Infusion at Provis Infusion Clinic

We would like to make the coordination of systemic therapy at the Provis Clinic and your facility as easy and seamless as possible for both you and your patient.

Rituxan® infusions are given by variable schedules and doses depending upon indication.

- 1. In all cases a CBC and chemistry are required within 2 weeks of commencing Rituxan®. In some instances high circulating white cell counts may alter planned schedule or date of initiation of Rituxan®.
- 2. Where there exist significant co-morbidities or high white cell counts, direct communication with the Provis Medical Director is important.

If there are any questions or concerns, please do not hesitate to contact our office at Tel. 416-595-0500.

The Provis Team