	<b>Legacy Day Treatment Unit Provider's Orders</b>	<b>Patient Name:</b> _____ <b>Date of Birth:</b> _____ <b>Med. Rec. No (TVC MRN Only):</b> _____
	Adult Ambulatory Infusion Order INFLIXIMAB-dyyb (INFLECTRA) & INFLIXIMAB (REMICADE)	
ALL ORDERS MUST BE MARKED IN INK WITH A CHECKMARK (✓) TO BE ACTIVE		

**Anticipated Start Date:** \_\_\_\_\_ **Patient to follow up with provider on date:** \_\_\_\_\_

\*\*\*This plan will expire after 365 days, unless otherwise specified below\*\*\*

**Orders expire:** \_\_\_\_\_

**Weight:** \_\_\_\_\_ kg **Height:** \_\_\_\_\_ cm

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

**Diagnosis:** \_\_\_\_\_ **Diagnosis Code:** \_\_\_\_\_

#### GUIDELINES FOR PRESCRIBING:


1. Send **FACE SHEET, INSURANCE CARD and most recent provider chart or progress note.**
2. A Tuberculin test (PPD) or QuantiFERON Gold must have been read as negative prior to initiation  
If the QuantiFERON Gold is indeterminate a CHEST X-ray should be performed to rule out infection
3. Hepatitis B (Hep B surface antigen AND core antibody) screening must be completed prior to initiation of therapy and the patient should not be infected
4. Patient should not have an active ongoing infection, signs or symptoms of malignancy, or moderate to severe heart failure at the onset of INFLIXimab therapy. Baseline liver function tests should be normal
5. Patients should have regular monitoring for TB, Hepatitis B, infection, malignancy, and liver abnormalities throughout therapy

#### PRE-SCREENING: (Results must be available prior to initiation of therapy)

- |   |                    |                                   |                                   |
|---|--------------------|-----------------------------------|-----------------------------------|
| • Hepatitis B Surface AG                          | Result Date: _____ | <input type="checkbox"/> Positive | <input type="checkbox"/> Negative |
| • Hepatitis B Core AB Qual                        | Result Date: _____ | <input type="checkbox"/> Positive | <input type="checkbox"/> Negative |
| • Tuberculin Test                                 | Result Date: _____ | <input type="checkbox"/> Positive | <input type="checkbox"/> Negative |
| • QuantiFERON Gold Test                           | Result Date: _____ | <input type="checkbox"/> Positive | <input type="checkbox"/> Negative |
| • Chest X-ray (if QuantiFERON Gold indeterminate) | Result Date: _____ | <input type="checkbox"/> Positive | <input type="checkbox"/> Negative |
| • Baseline CMP/LFT                                | Result Date: _____ | <input type="checkbox"/> WNL      | <input type="checkbox"/> Negative |

#### LABS TO BE DRAWN (orders must be placed in TVC Epic by ordering provider if TVC provider):

- ☐ Complete Metabolic Panel, Routine, ONCE, every \_\_\_\_\_(visit)(days)(weeks)(months)- **Circle one**
- ☐ CBC with differential, Routine, ONCE, every \_\_\_\_\_(visit)(days)(weeks)(months)- **Circle one**
- ☐ Other: \_\_\_\_\_

	<b>Legacy Day Treatment Unit Provider's Orders</b>	<b>Patient Name:</b> <b>Date of Birth:</b> <b>Med. Rec. No (TVC MRN Only):</b>
	Adult Ambulatory Infusion Order INFliximab-dyyb (INflectra) & INFliximab (REmicide)	

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**PRE-MEDICATIONS:**

- ☐ acetaminophen (TYLENOL) tablet, 650 mg, oral, ONCE, every visit
- ☐ diphenhydramine (BENADRYL) tablet, 50 mg, oral, ONCE, every visit  
(Give either cetirizine or diphenhydramine, not both.)
- ☐ cetirizine (ZYRTEC) tablet, 10 mg, oral, ONCE, every visit  
(Give either cetirizine or diphenhydramine, not both.)
- ☐ methylprednisolone sodium succinate (SOLU-MEDROL), 40 mg, IV, ONCE, every visit  
(Choose if patient has required IV steroids for a reaction during prior TNF-alpha inhibitor infusion.)

**MEDICATIONS: (must check one):**

**Biosimilar selection (must check one) – applies to all orders below**


- ☐ INFLECTRA (inFLIXimab-dyyb) **\*\*formulary agent\*\***
- ☐ REMICADE (inFLIXimab) *Restricted ONLY to existing REMICADE patients for continuing therapy, or patients whose insurance will only cover REMICADE*
- ☐ RENFLEXIS (inFLIXimab-abda)
- ☐ AVSOLA (inFLIXimab-axxq)

**Dose:** (Pharmacist will use most recent weight and round dose to the nearest 100 mg vial – dose selected at initiation will be continued, unless new orders are received specifying a dose modification)

- ☐ 3 mg/kg in NaCl 0.9% IV, ONCE, every visit
- ☐ 5 mg/kg in NaCl 0.9% IV, ONCE, every visit
- ☐ 10 mg/kg in NaCl 0.9% IV, ONCE, every visit
- ☐ \_\_\_\_\_ mg/kg in NaCl 0.9% IV, ONCE, every visit
- ☐ \_\_\_\_\_ mg in NaCl 0.9% IV, ONCE, every visit

**Interval:**

- ☐ Once
- ☐ Three doses at Week 0, 2, and 6, then every \_\_\_\_\_ weeks
- ☐ Every \_\_\_\_\_ weeks
- ☐ Other: \_\_\_\_\_

	<b>Legacy Day Treatment Unit Provider's Orders</b>  Adult Ambulatory Infusion Order INFLIXIMAB-dyyb (INFLECTRA) & INFLIXIMAB (REMICADE)	<b>Patient Name:</b>  <b>Date of Birth:</b>  <b>Med. Rec. No (TVC MRN Only):</b>
ALL ORDERS MUST BE MARKED IN INK WITH A CHECKMARK (✓) TO BE ACTIVE		

#### AS NEEDED MEDICATIONS:


- acetaminophen 650 mg oral, EVERY 4 HOURS AS NEEDED for hypersensitivity or infusion reaction, chills, or malaise
- diphenhydramine 25 mg oral, may repeat x 1 EVERY 4 HOURS AS NEEDED for itching
- NaCl 0.9% 500 mL IV, AS NEEDED, ONCE, infusion tolerability. Give concurrently with inFLIXimab/inFLIXimab biosimilars

#### NURSING ORDERS (TREATMENT PARAMETERS):

1. Hold treatment and contact provider if Hepatitis B surface antigen or core antibody total test result is positive, TB test result is positive, or if screening has not been performed.
2. Hold infusion and contact provider if patient has signs or symptoms of infection.
3. Monitor and record vital signs, tolerance and presence of infusion-related reactions prior to infusion and at the end of the infusion.
4. Monitor vital signs every 15 minutes for the first hour, then every 30 minutes for remainder of infusion. Stop the infusion immediately if a reaction occurs.
5. For 1<sup>st</sup> infusion or for any patient with history of infusion-related adverse reactions, monitor patient for 30 minutes after completion of treatment.
6. Manage line per LH policy 904.4007 IV Catheter Insertion (Peripheral) and LH 904.4004 IV Access: Central Catheters
7. Manage hypersensitivity reactions per LH 906.6606

**HYPERSENSITIVITY MEDICATIONS:** Refer to LH policy 906.6606 Initiation of Emergency Measures for Adult Oncology, Radiation Oncology and Infusion Clinic Patients

1. hydrocortisone 100 mg IV, AS NEEDED x1 dose for hypersensitivity reaction
2. diphenhydramine 25-50 mg IV, EVERY 2 HOURS AS NEEDED for hypersensitivity reaction (Max dose: 50 mg)
3. famotidine 20 mg IV, AS NEEDED x1 dose for hypersensitivity reaction
4. EPINEPHrine 0.3 mg IM, AS NEEDED x1 dose for hypersensitivity reaction
5. naloxone (Narcan) 0.4 mg IV, AS NEEDED x1 dose for diminished respiratory rate
6. Nursing communication order, every visit: Please follow treatment algorithm for acute infusion reaction

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Please check the appropriate box for the patient's preferred clinic location:

- |   |   |
|---|---|
| <input type="checkbox"/> <b>Legacy Day Treatment Unit –<br/>The Vancouver Clinic Building</b><br><i>A department of Salmon Creek Medical Center</i><br>700 NE 87 <sup>th</sup> Avenue, Suite 360<br>Vancouver, WA 98664<br>Phone number: 360-896-7070<br>Fax number: 360-487-5773 | <input type="checkbox"/> <b>Legacy Emanuel Day Treatment Unit</b><br><i>A department of Emanuel Medical Center</i><br>501 N Graham Street, Suite 540<br>Portland, OR 97227<br>Phone number: 503-413-4608<br>Fax number: 503-413-4887        |
| <input type="checkbox"/> <b>Legacy Salmon Creek Day Treatment Unit</b><br>Legacy Salmon Creek Medical Center<br>2121 NE 139 <sup>th</sup> Street, Suite 110<br>Vancouver, WA 98686<br>Phone number: 360-487-1750<br>Fax number: 360-487-5773                                      | <input type="checkbox"/> <b>Legacy STEPS Clinic</b><br><i>A department of Silverton Medical Center</i><br>Legacy Woodburn Health Center<br>1475 Mt Hood Ave<br>Woodburn, OR 97071<br>Phone number: 503-982-1280<br>Fax number: 503-225-8723 |

**Provider signature:** \_\_\_\_\_ **Date/Time:** \_\_\_\_\_

**Printed Name:** \_\_\_\_\_ **Phone:** \_\_\_\_\_ **Fax:** \_\_\_\_\_

**Organization/Department:** \_\_\_\_\_