

**Ruxience® (riTUXimab) 375 -
idelalisib 150 mg**

Regimen: Cycle I (Part I)

ARIA Protocol Name: Ruxience (rituximab) idelalisib

Adult Chemotherapy - Hematology Oncology

Chronic Lymphocytic Leukemia (CLL) or Small Lymphocytic Lymphoma (SLL)

Name: _____

HCN: _____

Date of Birth: _____



CC6320 0442 01/2024

Allergies:

No Known

Date: DD/MONTH/YYYY Planned Administration Date: DD/MONTH/YYYY

Cycle _____ of _____ **Cycle Duration: 28 days** Date of previous cycle: DD/MONTH/YYYY

MAY PROCEED WITH DOSES AS WRITTEN IF:

- ANC **greater than or equal to** 0.5 X 10⁹/L and platelets **greater than or equal to** 25 X 10⁹/L, otherwise notify Hematologist.
- LFTs and Bilirubin assessed.
- Creatinine clearance assessed.

PREMEDICATIONS (FOR HOSPITAL PHARMACY):

- acetaminophen 650 mg PO** pre riTUXimab on day 1
- diphenhydrAMINE 50 mg PO** pre riTUXimab on day 1
- allopurinol 300 mg PO** on day 1 (**for cycle 1 only**)
- Other: _____

HYDRATION/SUPPORTIVE CARE (FOR HOSPITAL PHARMACY):

- acetaminophen 650 mg PO** prn x 1 dose (Give during riTUXimab infusion if infusion lasts greater than 4 hours)
- diphenhydrAMINE 50 mg PO** prn x 1 dose (Give during riTUXimab infusion if infusion lasts greater than 4 hours)
- meperidine 25-50 mg IV** q1h prn x 2 doses (For chills and rigors associated with riTUXimab)
- Other: _____

PLEASE REFER TO CHEMOTHERAPY LETTER WHEN ORDERING SUPPORTIVE MEDICATIONS FOR THIS PATIENT

Authorized Prescriber: _____ Date: DD/MONTH/YYYY Time: _____

Authorized Prescriber's Signature: _____ ID #: _____

Nurse's Name: _____ Date: DD/MONTH/YYYY Time: _____

Nurse's Signature: _____

THIS IS A CONTROLLED DOCUMENT. PLEASE ENSURE THAT YOU ARE READING THE MOST RECENT VERSION.



Doctor's Order Sheet

**Ruxience® (riTUXimab) 375 -
idelalisib 150 mg**

Regimen: Cycle I (Part II)

ARIA Protocol Name: Ruxience (rituximab) idelalisib

Adult Chemotherapy - Hematology Oncology

Chronic Lymphocytic Leukemia (CLL) or Small Lymphocytic Lymphoma (SLL)

Name: _____

HCN: _____

Date of Birth: _____



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Weight: _____ kg Height: _____ cm Body Surface Area (BSA) = _____

CHEMOTHERAPY (FOR HOSPITAL PHARMACY):

Ruxience™ (riTUXimab) 375 mg/m² X BSA = _____ mg

IV diluted to 1mg/ml in normal saline and administer as per protocol on day 1 during first dose or patient who experienced a previous reaction. Subsequent infusions (if no previous reaction and not using subcutaneous rituximab option) dilute in 250 mL normal saline, infuse 20% of total volume over 30 minutes, then remaining volume over 60 minutes. Observe patient for 30 minutes after infusion is complete.

CHEMOTHERAPY (FOR COMMUNITY PHARMACY):

idelalisib 150 mg PO twice daily continuously

Dose modification: **idelalisib 100 mg PO** twice daily continuously

Mitte: 30 days

HYDRATION/SUPPORTIVE CARE (FOR COMMUNITY PHARMACY):

allopurinol 300 mg PO once daily on days 2 to 5 (for cycle 1 only)

metoclopramide 10-20 mg PO every 6 hours as needed

sulfamethoxazole/trimethoprim 400/80 mg PO once daily for duration of treatment and 2 to 6 months after completion of treatment

Other: _____

PLEASE REFER TO CHEMOTHERAPY LETTER WHEN ORDERING SUPPORTIVE MEDICATIONS FOR THIS PATIENT

Authorized Prescriber: _____ Date: DD/MONTH/YYYY Time: _____

Authorized Prescriber's Signature: _____ ID #: _____

Nurse's Name: _____ Date: DD/MONTH/YYYY Time: _____

Nurse's Signature: _____

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