



Doctor's Order Sheet

Ruxience® (riTUXimab) 500 - idelalisib 150 mg

Regimen: Cycles 2-8 (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: _____



CC6330 0443 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** $0.5 \times 10^9/L$ and platelets **greater than or equal to** $25 \times 10^9/L$, 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
- 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: _____

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Doctor's Order Sheet

**Ruxience® (riTUXimab) 500 -
idelalisib 150 mg**

Regimen: Cycles 2-8 (Part II)

ARIA Protocol Name: Ruxience (rituximab) idelalisib

Adult Chemotherapy - Hematology Oncology

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



CC6330 0443 01/2024

Name: _____

HCN: _____

Date of Birth: _____

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

CHEMOTHERAPY (FOR HOSPITAL PHARMACY):

Ruxience™ (riTUXimab) 500 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):

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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