ADULT Updated: February 8, 2023

Regimen Reference Order - LEUK - GIMEMA IT (ALL Ph+) IT Prophylaxis

Intrathecal therapy (IT) Prophylaxis is prescribed in combination with induction oral dasatinib,

and with each cycle of blinatumomab post induction

ARIA: LEUK - [GIMEMA IT induction]
LEUK - [GIMEMA IT post-blin]

Planned Course: Intrathecal therapy (IT) on Day +14, +22, +43, +57, +85 of induction with oral

dasatinib, followed by

Intrathecal therapy (IT) on Day +29 at the end of each cycle of blinatumomab

(post induction)

Indication for Use: Acute Lymphoblastic Leukemia, Philadelphia Chromosome Positive (Ph+);

Newly Diagnosed

CVAD: At Provider's Discretion

SEQUENCE OF MEDICATION ADMINISTRATION

| Pre-treatment Requirements | | | | |
|----------------------------|------|------|-------------------------------|--|
| | Drug | Dose | CCMB Administration Guideline | |
| Not Applicable | | | | |

Intrathecal Therapy (IT) during Induction ARIA: LEUK - [GIMEMA IT induction]

Proceed with treatment if:

- Platelets equal to or greater than 50 x 10⁹/L
- INR less than 1.5
- Fibrinogen greater than 1
 - Contact Leukemia/BMT (L/BMT) Physician if parameters not met

Days +14, +43 and +85 of induction

| Drug | Dose | CCMB Administration Guideline |
|--------------|-------|--|
| methotrexate | 12 mg | Intrathecal in 6 mL preservative free normal saline administered in L/BMT Clinic |

Days +22 and + 57 of induction

| Drug | Dose | CCMB Administration Guideline |
|------------|-------|--|
| cytarabine | 50 mg | Intrathecal in 6 mL preservative free normal saline administered in L/BMT Clinic |



Intrathecal Therapy (IT) at the end of each cycle of blinatumomab ARIA: LEUK - [GIMEMA IT post-blin]

Proceed with treatment if:

- Platelets equal to or greater than 50 x 10⁹/L
- INR less than 1.5
- Fibrinogen greater than 1
 - Contact Leukemia/BMT (L/BMT) Physician if parameters not met

Day +29 of Cycles 1, 3, and 5 of blinatumomab

| Drug | Dose | CCMB Administration Guideline |
|------------|-------|--|
| cytarabine | 50 mg | Intrathecal in 6 mL preservative free normal saline administered in L/BMT Clinic |

Day +29 of Cycles 2 and 4 of blinatumomab

| Drug | Dose | CCMB Administration Guideline |
|--------------|-------|--|
| methotrexate | 12 mg | Intrathecal in 6 mL preservative free normal saline administered in L/BMT Clinic |

In the event of an infusion-related hypersensitivity reaction, refer to the 'Hypersensitivity Reaction Standing Order'

REQUIRED MONITORING

All Cycles

· CBC and biochemistry as per Physician Orders

| Recommended Support Medications | | | |
|---------------------------------|---------------|------|-------------------------------|
| | Drug | Dose | CCMB Administration Guideline |
| | None required | | |

DISCHARGE INSTRUCTIONS

Not applicable

ADDITIONAL INFORMATION

- This Regimen Reference Order is intended for patients receiving intrathecal (IT) therapy on Days +14, +22, +43, +57 and +85 during induction with oral dasatinib and on Day +29 after each cycle of blinatumomab post induction
- IT therapy (induction) is ordered as a separate Support protocol under GIMEMA IT Induction in the "Leukemia" folder
- IT therapy (post blinatumomab) is ordered as a separate Support protocol under GIMEMA IT post blin in the "Leukemia" folder

