Regimen Reference Order – HEME – luspatercept (MDS)

ARIA: HEME – [luspatercept]

Planned Course: **Every 21 days**
Indication for Use: **Transfusion-dependent anemia due to Myelodysplastic Syndrome (MDS)**

CVAD: Not Required

Proceed with treatment if:

Hemoglobin less than 115 g/L

- **Contact Hematologist if parameters not met**

SEQUENCE OF MEDICATION ADMINISTRATION

Pre-treatment Requirements

Drug	Dose	CCMB Administration Guideline
Not Applicable		

Treatment Regimen – HEME – luspatercept (MDS)

Drug	Dose	CCMB Administration Guideline
luspatercept	1 mg/kg*	Subcutaneous once every 3 weeks into the upper arm, thigh, and/or abdomen **Alert: Maximum volume at each injection site is 1.2 mL*
**Dose may be increased to 1.33 mg/kg at the hematologist's discretion if patient is not RBC transfusion-free after at least two consecutive doses (6 weeks) at the 1 mg/kg starting dose. Dose may be further increased to 1.75 mg/kg if patient is not RBC transfusion-free after at least two consecutive doses (6 weeks) at the 1.33 mg/kg dose* **If hemoglobin increases by greater than 20 g/L within 3 weeks, in the absence of a transfusion, dose reduction may be required at the hematologist's discretion*		

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

REQUIRED MONITORING

All Cycles

- CBC, serum creatinine, urea, electrolytes and liver enzymes as per Physician Orders
- Blood pressure at each clinic visit
- Full vital signs (temperature, heart rate, respiratory rate, blood pressure and O_2 saturation) prior to each dose and as clinically indicated
- No observation period required. Patient can be discharged from treatment room if stable whether they had a reaction or not

Recommended Support Medications

Drug	Dose	CCMB Administration Guideline
None required		

DISCHARGE INSTRUCTIONS

- Patients should be instructed to contact their cancer team immediately if symptoms of hypersensitivity reactions occur after discharge
- Confirm that females of childbearing potential received the manufacturer Patient Card (wallet card) at Cycle 1 by the clinic
- Instruct patient to report to clinic if they are experiencing injection site reactions that are bothersome

ADDITIONAL INFORMATION

- luspatercept may increase blood pressure
- luspatercept is teratogenic
- luspatercept is provided by an Access Program with requirements. Prescriber's Checklist must be completed and a Patient Card must be given to women of childbearing potential