

M43 - ENOXAPARIN (LOVENOX)

MEDICATION STANDING ORDER

HIGH ALERT MEDICATION 1

Version date: 2023-12-13 Effective date: 2024-01-16 (0700)

INDICATIONS

 Known or suspected ST elevation myocardial infarction (STEMI) if the patient is going directly to primary coronary intervention (PCI) and only after consultation with the Code-STEMI physician

CONTRAINDICATIONS

- Patient may be candidate for fibrinolysis with TNK
- Known hypersensitivity to enoxaparin
- Patient is known to be on an anticoagulant and has taken it that day
- History of heparin-induced thrombocytopenia (HIT) within the past 100 days
- Active bleeding that cannot be controlled by basic measures or at a non-compressible site

ROUTE (WORK SCOPE)	INITIAL DOSE	REPEAT DOSE
INTRAVENOUS (PCP & ABOVE)	Less than 75 years - 0.5 mg/kg (max = 50 mg)	
SUBCUTANEOUS (PCP & ABOVE)	Less than 75 years - 1 mg/kg (max = 100 mg) 2	None
	More than 75 years -0.75 mg/kg (max = 75 mg) 3	

NOTES

- 1. ERS HIGH-ALERT MEDICATION: Refer to Shared Health Provincial Clinical Standard Safety Controls for High-Alert Medications (refer to A03 HIGH ALERT MEDICATIONS).
- 2. There is an increased risk of intracranial bleeding with intravenous enoxaparin in patients over 75 years of age.
 - Patients under 75 years of age should receive subcutaneous heparin only f IV access cannot be obtained.
 - Patients over 75 years of age should r4eceive only subcutaneous enoxaparin at a reduced dose.
- 3. The dose should be rounded off to the nearest 10 mg

APPROVED BY		
Buftslevel	ffmanl.	
Medical Director - Provincial EMS/PT	Associate Medical Director - Provincial EMS/PT	

VERSION CHANGES (refer to X08 for change tracking)

• Addition of Shared Health Provincial Clinical Standard for high-alert medications