

Enoxaparin (Lovenox®) Considerations for Use*

Non-FDA Approved Indication: Alternative to Oral Anticoagulation for Stroke Prevention in Atrial Fibrillation

Black Box Warning*	Epidural and spinal hematomas may occur in patients who are anticoagulated and are receiving neuraxial anesthesia or undergoing spinal punctures.
Mechanism of Action	A low molecular weight heparin with antithrombotic properties
Dosing [†]	<p><u>Adult</u>: 1 mg/kg twice daily²</p> <p><u>Elderly</u>: No specific dosage adjustment</p> <p><u>Hepatic Impairment</u>: No specific dosage adjustment</p> <p><u>Renal Impairment</u>: <u>CrCl < 30 mL/min</u>: 1 mg/kg SQ daily</p>
Contraindications	Active pathological bleeding, thrombocytopenia, history of heparin-induced thrombocytopenia, pork allergy
Major Side Effects	Hemorrhagic event, heparin-induced thrombocytopenia
Dosage forms and Strengths	<u>SQ</u> : solution for injection
Reversal	<p>Discontinue enoxaparin. Anticoagulation effects should be minimized within 12 hours.</p> <p>Protamine, given as a slow IV infusion (1 % solution), may largely neutralize enoxaparin.</p> <ul style="list-style-type: none"> • If enoxaparin was given within the past 8 hours, the dose of protamine should match the dose of enoxaparin given (1 mg protamine neutralizes 1 mg of enoxaparin). If the aPTT remains prolonged after 2-4 hours, a second infusion of 0.5 mg protamine per 1 mg of enoxaparin may be given. • If enoxaparin was given more than 8 hours ago, an infusion of protamine 0.5 mg for every 1 mg of enoxaparin may be given.
Conversion to/from other drugs	<p><u>From enoxaparin to warfarin</u>:</p> <ul style="list-style-type: none"> • Initiate warfarin when appropriate (usually start on same day or within 72 hours of enoxaparin). • Continue enoxaparin until INR is within therapeutic range for at least 2 days.
Special Notes	<p>Monitor for signs and symptoms of bleeding.</p> <p>Monitor hemoglobin, hematocrit, platelets, stool for occult blood.</p> <p>Consider monitoring anti-Xa activity (goal 0.3 to 0.7 IU/mL antifactor Xa activity) with severe renal dysfunction, weight < 45 kg, or obese patients.</p> <p>Discontinue all heparin products when heparin induced thrombocytopenia is suspected or diagnosed.</p> <p>Not adequately studied in patients with mechanical heart valves.</p>
Counseling	Report signs and symptoms of bleeding (e.g., unexpected bleeding or bleeding that lasts a long time; red or black, tarry stool; pink or brown urine; unusual bruising; coughing up blood; vomiting blood or vomit that looks like coffee grounds; unexplained pain, swelling, or joint pain; unusual headaches, dizziness, or weakness; recurring nose bleeds)

* Refer to prescribing information for more complete information.

†Dosages given in the table may differ from those recommended by the manufacturers.

Sources:

1. American College of Cardiology (ACC), American Heart Association (AHA), and Heart Rhythm Society (HRS). *2011 ACCF/AHA/HRS Focused Update on the Management of Patients With Atrial Fibrillation (Update on Dabigatran)*. Washington, DC: American College of Cardiology Foundation. 2011.
2. Chest Supplement, Antithrombotic Therapy and Prevention of Thrombosis, 9th edition, American College of Chest Physicians.
3. Lovenox® prescribing information, 5/16/07.