HIGHLIGHTS OF PRESCRIBING INFORMATION These highlights do not include all the information needed to use ENOXAPARIN SODIUM INJECTION safely and effectively. See full prescribing information for **ENOXAPARIN SODIUM INJECTION.** ENOXAPARIN SODIUM injection, for subcutaneous use Hypersensitivity to enoxaparin sodium (4 Initial U.S. Approval: 1993 · Hypersensitivity to heparin or pork products (4)

WARNING: SPINAL/EPIDURAL HEMATOMAS

See full prescribing information for complete boxed warning.

Epidural or spinal hematomas may occur in patients who are anticoagulated with low molecular weight heparins (LMWH) or heparinoids and are receiving neuraxia anesthesia or undergoing spinal puncture. These hematomas may result in long-term or permanent paralysis. Consider these risks when scheduling patients for spinal rocedures. Factors that can increase the risk of developing epidural or spinal hematomas in these patients include: Use of indwelling epidural catheters $Concomitant \ use \ of \ other \ drugs \ that \ affect \ hemostasis, such as \ non-steroidal \ anti-inflammatory \ drugs \ (NSAIDs), \ platelet \ inhibitors, \ and \ other \ anti-coagulants$

A history of traumatic or repeated epidural or spinal punctures A history of spinal deformity or spinal surgery
 Optimal timing between the administration of enoxaparin sodium injection and neuraxial procedures is not known Monitor patients frequently for signs and symptoms of neurological impairment. If neurological compromise is noted, urgent treatment is necessary. (5.1, 7)

Enoxaparin sodium injection is a low molecular weight heparin [LMWH] indicated for • Prophylaxis of deep vein thrombosis (DVT) in abdominal surgery, hip replacement surgery, knee replacement surgery, or medical patients with severely restricted mobility during acute illness (1.1) Inpatient treatment of acute DVT with or without pulmonary embolism (1.2) Outpatient treatment of acute DVT without pulmonary embolism (1.2)

Prophylaxis of ischemic complications of unstable angina and non-Q-wave myocardial infarction [MI] (1.3) Treatment of acute ST-segment elevation myocardial infarction [STEMI] managed medically or with subsequent percutaneous coronary intervention [PCI] (1.4)

- DOSAGE AND ADMINISTRATION

See full prescribing information for dosing and administration information. (2) — DOSAGE FORMS AND STRENGTHS

100 mg/mL concentration (3):

Single-dose prefilled syringes: 30 mg/0.3 mL, 40 mg/0.4 mL
 Single-dose graduated prefilled syringes: 60 mg/0.6 mL, 80 mg/0.8 mL, 100 mg/mL

150 mg/mL concentration (3):

Single-dose graduated prefilled syringes: 120 mg/0.8 mL, 150 mg/mL

FULL PRESCRIBING INFORMATION: CONTENTS

WARNING: SPINAL/EPIDURAL HEMATOMAS 1 INDICATIONS AND USAGE

Prophylaxis of Deep Vein Thrombosis Treatment of Acute Deep Vein Thrombosis Prophylaxis of Ischemic Complications of Unstable Angina and Non-Q-Wave Myocardial Infarction

2 DOSAGE AND ADMINISTRATION Pretreatment Evaluatio

Treatment of Acute ST-Segment Elevation Myocardial Infarction

2.2 Adult Dosage

Dose Reduction for Patients with Severe Renal Impairme Recommended Dosage for Geriatric Patients with Acute ST-Segment Elevation Myocardial Infarction Administration Monitoring for Safety

3 DOSAGE FORMS AND STRENGTHS **4 CONTRAINDICATIONS**

5 WARNINGS AND PRECAUTIONS Increased Risk of Hemorrhage

Increased Risk of Bleeding following Percutaneous Coronary Revascularization Procedures Increased Risk of Bleeding in Patients with Concomitant Medical Conditions 5.3 Risk of Heparin-Induced Thrombocytopenia with or without Thrombosis Thrombocytopenia Interchangeability with Other Heparins
Increased Risk of Thrombosis in Pregnant Women with Mechanical Prosthetic Heart Valves

6 ADVERSE REACTIONS Clinical Trials Experience Postmarketing Experience

7 DRUG INTERACTIONS

WARNING: SPINAL/ EPIDURAL HEMATOMAS Epidural or spinal hematomas may occur in patients who are anticoagulated with low molecular weight heparins (LMWH) or heparinoids and are receiving neuraxial anesthesia or undergoing spinal puncture. These hematomas may result in long-term or permanent paralysis. Consider these risks when scheduling patients for spinal rocedures. Factors that can increase the risk of developing epidural or spinal hematomas in these patients include Use of indwelling epidural catheters

Concomitant use of other drugs that affect hemostasis, such as non-steroidal anti-inflammatory drugs (NSAIDs), platelet inhibitors, and other anticoagulants A history of traumatic or repeated epidural or spinal punctures A history of spinal deformity or spinal surgery Optimal timing between the administration of enoxaparin sodium injection and neuraxial procedures is not known Monitor patients frequently for signs and symptoms of neurological impairment. If neurological compromise is noted, urgent treatment is necessary

FULL PRESCRIBING INFORMATION

(5.1) and Drug Interactions (7)]. 1 INDICATIONS AND USAGE 1.1 Prophylaxis of Deep Vein Thrombosis

Enoxaparin sodium injection is indicated for the prophylaxis of deep vein thrombosis (DVT), which may lead to pulmonary embolism (PE):

 in patients undergoing abdominal surgery who are at risk for thromboembolic complications [see Clinical Studies (14.1)]
 in patients undergoing hip replacement surgery, during and following hospitalization
 in patients undergoing knee replacement surgery · in medical patients who are at risk for thromboembolic complications due to severely restricted mobility during acute illness

1.2 Treatment of Acute Deep Vein Thrombosis Enoxaparin sodium injection is indicated for

 the inpatient treatment of acute deep vein thrombosis with or without pulmonary embolism, when administered in conjunction with warfarin sodium
 the outpatient treatment of acute deep vein thrombosis without pulmonary embolism, when administered in conjunction with warfarin sodium 1.3 Prophylaxis of Ischemic Complications of Unstable Angina and Non-Q-Wave Myocardial Infarction

Enoxaparin sodium injection is indicated for the prophylaxis of ischemic complications of unstable angina and non-Q-wave myocardial infarction, when concurrently administere with aspirin.

1.4 Treatment of Acute ST-Segment Elevation Myocardial Infarction Enoxaparin sodium injection, when administered concurrently with aspirin, has been shown to reduce the rate of the combined endpoint of recurrent myocardial infarction or deat

in patients with acute ST-segment elevation myocardial infarction (STEMI) receiving thrombolysis and being managed medically or with percutaneous coronary intervention (PCI).

2.1 Pretreatment Evaluatio Evaluate all patients for a bleeding disorder before starting enoxaparin sodium injection treatment, unless treatment is urgently needed.

Consider the benefits and risks before neuraxial intervention in patients anticoagulated or to be anticoagulated for thromboprophylaxis [see Warnings and Precautic

2.2 Adult Dosage Abdominal Surgery nded dose of enoxaparin sodium injection is 40 mg by subcutaneous injection once a day (with the initial dose given 2 hours prior to surgery) in patients undergoing

abdominal surgery who are at risk for thromboembolic complications. The usual duration of administration is 7 to 10 days [see Clinical Studies (14.1)]. Hip or Knee Replacement Surgery he recommended dose of enoxaparin sodium injection is 30 mg every 12 hours administered by subcutaneous injection in patients undergoing hip or knee replacement surger Administer the initial dose 12 to 24 hours after surgery, provided that hemostasis has been established. The usual duration of administration is 7 to 10 days [see Clinical Studies (14.2)].

A dose of enoxaparin sodium injection of 40 mg once a day subcutaneously may be considered for hip replacement surgery for up to 3 weeks. Administer the initial dose 12 (±3)

Medical Patients during Acute Illness The recommended dose of enoxaparin sodium injection is 40 mg once a day administered by subcutaneous injection for medical patients at risk for thromboembolic complications due to severely restricted mobility during acute illness. The usual duration of administration is 6 to 11 days [see Clinical Studies (14.3)]. Treatment of Deep Vein Thrombosis with or without Pulmonary Embolism

parin sodium injection is 1 mg/kg every 12 hours administered subcutaneously in patients with acute deep vein thrombosis without pulmonary embolism, who can be treated at home in an outpatient setting. The recommended dose of enoxaparin sodium injection is 1 mg/kg every 12 hours administered subcutaneously or 1.5 mg/kg once a day administered subcutaneously at the same time every day for inpatient (hospital) treatment of patients with acute deep vein thrombosis with pulmonary embolism or patients with acute deep vein thrombosis without

ulmonary embolism (who are not candidates for outpatient treatment). In both outpatient and inpatient (hospital) treatments, initiate warfarin sodium therapy when appropriate (usually within 72 hours of enoxaparin sodium injection). Continue enoxaparin sodium injection for a minimum of 5 days and until a therapeutic oral anticoagulant effect has been achieved (International Normalization Ratio 2 to 3). The average duration of administration is 7 days [see Clinical Studies (14.4)]. Unstable Angina and Non-Q-Wave Myocardial Infarction nded dose of enoxaparin sodium injection is 1 mg/kg administered subcutaneously every 12 hours in conjunction with oral aspirin therapy (100 to 325 mg once daily)

in patients with unstable angina or non-Q-wave myocardial infarction. Treat with enoxaparin sodium injection for a minimum of 2 days and continue until clinical stabilization. The usual duration of treatment is 2 to 8 days [see Warnings and Precautions (5.2) and Clinical Studies (14.5)]. Treatment of Acute ST-Segment Elevation Myocardial Infarction The recommended dose of enoxaparin sodium injection is a **single intravenous bolus of 30 mg** plus a 1 mg/kg subcutaneous dose followed by 1 mg/kg administered subcutaneously every 12 hours (maximum 100 mg for the first two doses only, followed by 1 mg/kg dosing for the remaining doses) in patients with acute ST-segment elevation myocardial infarction.

Reduce the dosage in patients >75 years of age [see Dosage and Administration (2.4)]. Unless contraindicated, administer aspirin to all patients as soon as they are identified as having STEMI and continue dosing with 75 to 325 mg once daily.

When administered in conjunction with a thrombolytic (fibrin specific or non-fibrin specific), administer enoxaparin sodium injection between 15 minutes before and 30 minutes after the start of fibrinolytic therapy. The usual duration of enoxaparin sodium injection therapy is 8 days or until hospital discharge. For patients managed with percutaneous coronary intervention (PCI), if the last enoxaparin sodium injection subcutaneous administration was given less than 8 hours before balloon inflation, no additional dosing is needed. If the last enoxaparin sodium injection subcutaneous administration was given more than 8 hours before balloon inflation, administer an intravenous bolus of 0.3 mg/kg of enoxaparin sodium injection [see Warnings and Precautions (5.2)].

2.3 Dose Reduction for Patients with Severe Renal Impairment The recommended prophylaxis and treatment dosage regimens for patients with severe renal impairment (creatinine clearance <30 mL/min) are described in Table 1 [see Use in

Specific Populations (8.7) and Clinical Pharmacology (12.3)]. Table 1 Dosage Regimens for Patients with Severe Renal Impairment (creatinine clearance <30 mL/minute)

30 mg administered subcutaneously once daily

Prophylaxis in hip or knee replacement surgery 30 mg administered subcutaneously once daily Prophylaxis in medical patients during acute illness 30 mg administered subcutaneously once daily Inpatient treatment of acute deep vein thrombosis with or without pulmonary embolism, when administered 1 mg/kg administered subcutaneously once daily n conjunction with warfarin sodium Outpatient treatment of acute deep vein thrombosis without pulmonary embolism, when administered in 1 mg/kg administered subcutaneously once daily conjunction with warfarin sodium Prophylaxis of ischemic complications of unstable angina and non-Q-wave myocardial infarction, when mg/kg administered subcutaneously once daily Treatment of acute ST-segment elevation myocardial infarction in patients <75 years of age, when 30 mg single intravenous bolus plus a 1 mg/kg subcutaneous dose administered in conjunction with aspirin followed by 1 mg/kg administered subcutaneously once daily Treatment of acute ST-segment elevation myocardial infarction in geriatric patients ≥75 years of age, wher 1 mg/kg administered subcutaneously once daily (no initial bolus administered in conjunction with aspirin Although no dose adjustment is recommended in patients with creatinine clearance 30 to 50 mL/min and creatinine clearance 50 to 80 mL/min, observe these patients frequently

for signs and symptoms of bleeding. 2.4 Recommended Dosage for Geriatric Patients with Acute ST-Segment Elevation Myocardial Infarction For treatment of acute ST-segment elevation myocardial infarction in geriatric patients ≥75 years of age, do not use an initial intravenous bolus. Initiate dosing with 0.75 mg/kg subcutaneously every 12 hours (maximum 75 mg for the first two doses only, followed by 0.75 mg/kg dosing for the remaining doses) [see Use in Specific Populations (8.5)]

and Clinical Pharmacology (12.3)] No dose adjustment is necessary for other indications in geriatric patients unless kidney function is impaired [see Dosage and Administration (2.2)].

2.5 Administration Do not administer enoxaparin sodium injection by intramuscular injection Administer enoxaparin sodium injection by intravenous or subcutaneous injection only.

Enoxaparin sodium injection is a clear, colorless to pale yellow sterile solution, and as with other parenteral drug products, should be inspected visually for particulate matter and

Patients may self-inject by the subcutaneous route of administration only after their physicians determine that it is appropriate and with medical follow-up, as necessary. Provide proper training in subcutaneous injection technique before allowing self-injection (with or without the assistance of an injection device). Subcutaneous Injection Technique Position patients in a supine position for enoxaparin sodium injection administration by deep subcutaneous injection
 Do not expel the air bubble from the prefilled syringes before the injection, to avoid the loss of drug.

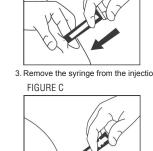
 Do not inject into skin that has bruises or scars. Do not inject through clothes. Alternate injection sites between the left and right anterolateral and left and right posterolateral abdominal wall. Introduce the whole length of the needle into a skin fold held between the thumb and forefinger; hold the skin fold throughout the injection. To minimize bruising, do not rub the

injection site after completion of the injection. Enoxaparin sodium injection prefilled syringes and graduated prefilled syringes are for single, one-time use only and are available with a system that shields the needle after injection Remove the prefilled syringe from the packaging by peeling at the arrow as directed on the lid. Do not remove by pulling on the plunger as this may damage the syringe 1. Remove the needle shield by pulling it straight off the syringe (see Figure A). If less than the full syringe volume is needed to administer the prescribed dose, elect syringe contents

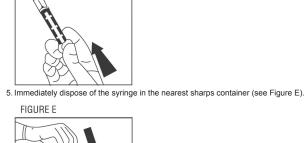
FIGURE A

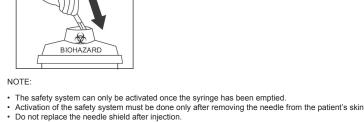
Prophylaxis in abdominal surgery

2. Inject using standard technique, pushing the plunger to the bottom of the syringe (see Figure B). FIGURE B



4. Orient the needle away from you and others and activate the safety system by firmly pushing the plunger rod. The protective sleeve will automatically cover the needle and an audible "click" will be heard to confirm shield activation (see Figure D).





· The safety system should not be sterilized. Activation of the safety system may cause minimal splatter of fluid. For optimal safety, activate the system while orienting it downwards away from yourself and others.

2.6 Monitoring for Safety During therapy monitor complete blood counts including platelets and stool occult blood. Assess for signs and symptoms of bleeding

In patients with renal impairment anti-Factor Xa levels may be used to monitor the anticoagulant effects of enoxaparin sodium injection If during enoxaparin sodium injection therapy abnormal coagulation parameters or bleeding should occur, anti- Factor Xa levels may be used to monitor the anticoagulant effects of enoxaparin sodium injection [see Clinical Pharmacology (12.3)].

Prothrombin Time (PT) and Activated Partial Thromboplastin Time (aPTT) are not adequate for monitoring the anticoagulant effects of enoxaparin sodium injection 3 DOSAGE FORMS AND STRENGTHS

Enoxaparin sodium injection, USP is a clear, colorless to pale-yellow sterile solution available in two concentrations. 100 mg/mL Concentratio Single-dose Prefilled Syringes 30 mg/0.3 mL, 40 mg/0.4 m

150 mg/mL Concentration - Single-dose Graduated Prefilled Syringes 120 mg/0.8 mL, 150 mg/mL 4 CONTRAINDICATIONS

Enoxaparin sodium injection is contraindicated in patients with Active major bleeding History of immune-mediated heparin-induced thrombocytopenia (HIT) within the past 100 days or in the presence of circulating antibodies [see Warnings and Precautions (5.4)]
 Known hypersensitivity to enoxaparin sodium (e.g., pruritus, urticaria, anaphylactic/anaphylactoid reactions) [see Adverse Reactions (6.2)]

Single-dose Graduated Prefilled Syringes 60 mg/0.6 mL, 80 mg/0.8 mL, 100 mg/ml

· Known hypersensitivity to heparin or pork products

5 WARNINGS AND PRECAUTIONS

5.1 Increased Risk of Hemorrhage Cases of epidural or spinal hemorrhage and subsequent hematomas have been reported with the use of enoxaparin sodium injection and epidural or spinal anesthesia/analgesia

or spinal puncture procedures, resulting in long-term or permanent paralysis. The risk of these events is higher with the use of post-operative indwelling epidural catheters, with the concomitant use of additional drugs affecting hemostasis such as NSAIDs, with traumatic or repeated epidural or spinal puncture, or in patients with a history of spinal surgery or spinal deformity [see Boxed Warning, Adverse Reactions (6.2) and Drug Interactions (7)].

To reduce the potential risk of bleeding associated with the concurrent use of enoxaparin sodium injection and epidural or spinal anesthesia/analgesia or spinal puncture, consider the pharmacokinetic profile of enoxaparin [see Clinical Pharmacology (12.3)]. Placement or removal of an epidural catheter or lumbar puncture is best performed when the anticoagulant effect of enoxaparin is low; however, the exact timing to reach a sufficiently low anticoagulant effect in each patient is not known. Placement or removal of a catheter should be delayed for at least 12 hours after administration of lower doses (30 mg once or twice daily or 40 mg once daily) of enoxaparin sodium injection and at least 24 hours after the administration of higher doses (0.75 mg/kg twice-daily, 1 mg/kg twice-daily, or 1.5 mg/kg once daily) of enoxaparin sodium injection. Anti-Xa levels are still detectable at these time points, and these delays are not a guarantee that neuraxial hematoma will be avoided. Patients receiving the 0.75 mg/kg twice daily dose, or the 1 mg/kg twice-daily dose should not receive the second enoxaparin dose in the twice daily regimen to allow a longer delay before catheter placement or removal. Likewise,

although a specific recommendation for timing of a subsequent enoxaparin sodium injection dose after catheter removal cannot be made, consider delaying this next dose for at

least four hours, based on a benefit-risk assessment considering both the risk for thrombosis and the risk for bleeding in the context of the procedure and patient risk factors. For

patients with creatinine clearance <30 mL/minute, additional considerations are necessary because elimination of enoxaparin is more prolonged; consider doubling the timing of removal of a catheter, at least 24 hours for the lower prescribed dose of enoxaparin sodium injection (30 mg once daily) and at least 48 hours for the higher dose (1 mg/kg/day) [see Clinical Pharmacology (12.3)]. - CONTRAINDICATIONS

Most common adverse reactions (>1%) were bleeding, anemia, thrombocytopenia, elevation of serum aminotransferase, diarrhea, nausea, ecchymosis, fever, edema, predema, dyspnea, confusion, and injection site pain (6.1)

— DRUG INTERACTIONS —

- USE IN SPECIFIC POPULATIONS -

To report SUSPECTED ADVERSE REACTIONS, contact Fresenius Kabi USA, LLC, at 1-800-551-7176 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch

Discontinue agents which may enhance hemorrhage risk prior to initiation of enoxaparin sodium injection or conduct close clinical and laboratory monitoring (2.6, 7)

Prophylaxis of Deep Vein Thrombosis following Abdominal Surgery in Patients at Risk for Thromboembolic Complications Prophylaxis of Deep Vein Thrombosis following Hip or Knee Replacement Surgery Prophylaxis of Deep Vein Thrombosis in Medical Patients with Severely Restricted Mobility during Acute Illness

Should the physician decide to administer anticoagulation in the context of epidural or spinal anesthesia/analgesia or lumbar puncture, frequent monitoring must be exercised to

detect any signs and symptoms of neurological impairment such as midline back pain, sensory and motor deficits (numbness or weakness in lower limbs), and bowel and/or bladder dysfunction. Instruct patients to report immediately if they experience any of the above signs or symptoms. If signs or symptoms of spinal hematoma are suspected, initiate urgent diagnosis and treatment including consideration for spinal cord decompression even though such treatment may not prevent or reverse neurological sequelae.

Use enoxaparin sodium injection with extreme caution in conditions with increased risk of hemorrhage, such as bacterial endocarditis, congenital or acquired bleeding disorders, active ulcerative and angiodysplastic gastrointestinal disease, hemorrhagic stroke, or shortly after brain, spinal, or ophthalmological surgery, or in patients treated concomitantly

Bleeding can occur at any site during therapy with enoxaparin sodium injection. An unexplained fall in hematocrit or blood pressure should lead to a search for a bleeding site.

To minimize the risk of bleeding following the vascular instrumentation during the treatment of unstable angina, non-Q-wave myocardial infarction and acute ST-segment elevation myocardial infarction, adhere precisely to the intervals recommended between enoxaparin sodium injection doses. It is important to achieve hemostasis at the puncture site after PCI.

In case a closure device is used, the sheath can be removed immediately. If a manual compression method is used, sheath should be removed 6 hours after the last intravenous/ subcutaneous enoxaparin sodium injection. If the treatment with enoxaparin sodium is to be continued, the next scheduled dose should be given no sooner than 6 to 8 hours after sheath removal. The site of the procedure should be observed for signs of bleeding or hematoma formation [see Dosage and Administration (2.1)].

Because HIT may still occur in these circumstances, the decision to use enoxaparin sodium injection in such a case must be made only after a careful benefit-risk assessment and

Platelet counts less than 50,000/mm3 occurred at a rate of 0.1% in patients given enoxaparin sodium injection, in 0.2% of patients given heparin, and 0.4% of patients given placebo

Enoxaparin sodium injection cannot be used interchangeably (unit for unit) with heparin or other low molecular weight heparins as they differ in manufacturing process, molecular weight distribution, anti-Xa and anti-Ila activities, units, and dosage. Each of these medicines has its own instructions for use.

Use of enoxaparin sodium injection for thromboprophylaxis in pregnant women with mechanical prosthetic heart valves may result in valve thrombosis. In a clinical study of pregnant

women with mechanical prosthetic heart valves given enoxaparin sodium injection (1 mg/kg twice daily) to reduce the risk of thromboembolism, 2 of 8 women developed clots resulting in blockage of the valve and leading to maternal and fetal death. No patients in the heparin/warfarin group (0 of 4 women) died. There also have been isolated postmarketing

reports of valve thrombosis in pregnant women with mechanical prosthetic heart valves while receiving enoxaparin sodium injection for thromboprophylaxis. Women with mechanical prosthetic heart valves may be at higher risk for thromboembolism during pregnancy and, when pregnant, have a higher rate of fetal loss from stillbirth, spontaneous abortion,

and premature delivery. Therefore, frequent monitoring of peak and trough anti-Factor Xa levels and adjusting of dosage may be needed [see Use in Specific Populations (8.6)].

Because clinical trials are conducted under widely varying conditions, adverse reaction rates observed in the clinical trials of a drug cannot be directly compared to rates in the clinical

During clinical development for the approved indications, 15,918 patients were exposed to enoxaparin sodium. These included 1,228 for prophylaxis of deep vein thrombosis

ollowing abdominal surgery in patients at risk for thromboembolic complications, 1,368 for prophylaxis of deep vein thrombosis following hip or knee replacement surgery, 711 for

prophylaxis of deep vein thrombosis in medical patients with severely restricted mobility during acute illness, 1,578 for prophylaxis of ischemic complications in unstable angina

and non-Q-wave myocardial infarction, 10,176 for treatment of acute ST-elevation myocardial infarction, and 857 for treatment of deep vein thrombosis with or without pulmonary embolism. Enoxaparin sodium doses in the clinical trials for prophylaxis of deep vein thrombosis following abdominal or hip or knee replacement surgery or in medical patients with

severely restricted mobility during acute illness ranged from 40 mg subcutaneously once daily to 30 mg subcutaneously twice daily. In the clinical studies for prophylaxis of ischemic complications of unstable angina and non-Q-wave myocardial infarction doses were 1 mg/kg every 12 hours and in the clinical studies for treatment of acute ST-segment elevation

Table 2 Major Bleeding Episodes following Abdominal and Colorectal Surgery*

Enoxaparin Sodium Injection

23 (4%)

n = 673

Bleeding complications were considered major: (1) if the hemorrhage caused a significant clinical event, or (2) if accompanied by a hemoglobin decrease ≥ 2 g/dL or transfusion of 2 or more units of blood products. Retroperitoneal, intraocular, and intracranial hemorrhages were always considered major.

Enoxaparin Sodium Injection

30 mg q12h

subcutaneously

n = 786

31 (4%)

n = 294

Dosing Regime

Enoxaparin Sodium Injection

40 mg daily subcutaneously

3 (<1%)

Enoxaparin Sodium Injection

1 mg/kg q12h subcutaneously

9 (2%)

Dosing Regimen

Dosing Regimen

Table 3 Major Bleeding Episodes following Hip or Knee Replacement Surgery*

* Bleeding complications were considered major: (1) if the hemorrhage caused a significant clinical event, or (2) if accompanied by a hemoglobin decrease ≥ 2 g/dL or transfusion of 2 or more units of blood products. Retroperitoneal and intracranial hemorrhages were always considered major. In the knee replacement surgery trials, intraocular hemorrhages

NOTE: At no time point were the 40 mg once a day pre-operative and the 30 mg every 12 hours post-operative hip replacement surgery prophylactic regimens compared in clinical trials

Injection site hematomas during the extended prophylaxis period after hip replacement surgery occurred in 9% of the enoxaparin sodium injection patients versus 1.8% of the

Table 4 Major Bleeding Episodes in Medical Patients with Severely Restricted Mobility during Acute Illness*

* Bleeding complications were considered major: (1) if the hemorrhage caused a significant clinical event, (2) if the hemorrhage caused a decrease in hemoglobin of ≥ 2 g/dL or transfusion of 2 or more units of blood products. Retroperitoneal and intracranial hemorrhages were always considered major although none were reported during the trial.

Table 5 Major Bleeding Episodes in Deep Vein Thrombosis with or without Pulmonary Embolism Treatment *

* Bleeding complications were considered major: (1) if the hemorrhage caused a significant clinical event, or (2) if accompanied by a hemoglobin decrease ≥ 2 g/dL or transfusion of 2 or

All patients also received warfarin sodium (dose-adjusted according to PT to achieve an INR of 2.0 to 3.0) commencing within 72 hours of enoxaparin sodium injection or standard heparin

Table 6 Major Bleeding Episodes in Unstable Angina and Non-Q-Wave Myocardial Infarction

Enoxaparin Sodium Injection

‡ Bleeding complications were considered major: (1) if the hemorrhage caused a significant clinical event, or (2) if accompanied by a hemoglobin decrease by ≥ 3 g/dL or transfusion of 2 or more units of blood products. Intraocular, retroperitoneal, and intracranial hemorrhages were always considered major.

Table 7 Major Bleeding Episodes in Acute ST-Segment Elevation Myocardial Infarction

Enoxaparin Sodium Injection* Initial 30 mg intravenous bolus followed by

1 mg/kg q 12h subcutaneously

n = 10176

n (%)

211 (2.1)

84 (0.8)

Other adverse reactions that were thought to be possibly or probably related to treatment with enoxaparin sodium injection, heparin, or placebo in clinical trials with patients undergoing hip or knee replacement surgery, abdominal or colorectal surgery, or treatment for DVT and that occurred at a rate of at least 2% in the enoxaparin sodium injection

Table 8 Adverse Reactions Occurring at ≥2% Incidence in Enoxaparin Sodium Injection-Treated Patients Undergoing Abdominal or Colorectal Surgery

Enoxaparin Sodium Injection 40 mg daily subcuta n = 1228

Table 9 Adverse Reactions Occurring at ≥2% Incidence in Enoxaparin Sodium Injection-Treated Patients Undergoing Hip or Knee Replacement Surgery

<1

n = 131‡

Total

Tota

Dosing Regimen

Enoxaparin Sodium Injectio

n = 1080

Total

Dosing Regimen

1 mg/kg q12h subcutaneously

n = 559

Total

Severe

30 mg q12h subcuta

Severe

Data represent enoxaparin sodium injection 40 mg subcutaneously once a day initiated up to 12 hours prior to surgery in 288 hip replacement surgery patients who received

enoxaparin sodium injection peri-operatively in an unblinded fashion in one clinical trial.

† Data represent enoxaparin sodium injection 40 mg subcutaneously once a day given in a blinded fashion as extended prophylaxis at the end of the peri-operative period in 131 of the original 288 hip replacement surgery patients for up to 21 days in one clinical trial.

Table 10 Adverse Reactions Occurring at ≥2% Incidence in Enoxaparin Sodium Injection-Treated Medical Patients with Severely Restricted Mobility during Acute Illness

Enoxaparin Sodium Injection

40 mg daily subcutaneously

n = 360

3.3

2.8

2.2

2.5

Enoxaparin Sodium Injection

1.5 mg/kg daily subcutaneously

n = 298

Severe

Table 11 Adverse Reactions Occurring at ≥2% Incidence in Enoxaparin Sodium Injection-Treated Patients Undergoing Treatment of Deep Vein Thrombosis with or without Pulmonary Embolism

Total

40 mg daily subcut

Enoxaparin Sodium Injection

40 mg daily subcutaneously

4 (2%)

0 (0%)

Enoxaparin sodium injection 30 mg every 12 hours subcutaneously initiated 12 to 24 hours after surgery and continued for up to 14 days after surgery Enoxaparin sodium injection 40 mg subcutaneously once a day initiated up to 12 hours prior to surgery and continued for up to 7 days after surgery

Enoxaparin Sodium Injection

20 mg daily subcutaneously

1 (<1%)

Enoxaparin Sodium Injection

1.5 mg/kg daily subcutaneously

5 (2%)

5000 U q8h subcutaned

n = 560 16 (3%)

n = 67421 (3%)

15,000 U/24h

subcutaneously

n = 541

32 (6%)

n = 225

Placebo[†]

n = 362 2 (<1%)

Heparin

Therapy

aPTT Adjusted Intravenous

9 (2%)

Heparin*

aPTT Adjusted Intravenous Therapy

18 (1%)

Heparin*

n = 10151

n (%)

138 (1.4)

66 (0.7)

5000 U q8h subcuta

<1

15,000 U/24h

n = 766

Total

daily subcutaneously

n = 362

5.2

1.7

Severe

aPTT Adjusted Intrav

n = 544

Total

Severe

n = 1234

Tota

Placebo

n = 115

Total

q12h subcutaneo

Severe

myocardial infarction enoxaparin sodium doses were a 30 mg intravenous bolus followed by 1 mg/kg every 12 hours subcutaneously.

The following rates of major bleeding events have been reported during clinical trials with enoxaparin sodium injection [see Tables 2 to 7].

Thrombocytopenia of any degree should be monitored closely. If the platelet count falls below 100,000/mm³, enoxaparin sodium injection should be discontinued.

Treatment of Deep Vein Thrombosis with or without Pulmonary Embolism
Prophylaxis of Ischemic Complications in Unstable Angina and Non-Q-Wave Myocardial Infarction

Major hemorrhages including retroperitoneal and intracranial bleeding have been reported. Some of these cases have been fatal.

Severe Renal Impairment: Adjust dose for patients with creatinine clearance <30 mL/min (2.3, 8.7)

See 17 for PATIENT COUNSELING INFORMATION and FDA- approved patient labeling

Patients with Mechanical Prosthetic Heart Valves

Carcinogenesis, Mutagenesis, Impairment of Fertility Animal Toxicology and/or Pharmacology

Treatment of Acute ST-Segment Elevation Myocardial Infarction

5.2 Increased Risk of Bleeding following Percutaneous Coronary Revascularization Procedures

5.3 Increased Risk of Bleeding in Patients with Concomitant Medical Conditions

5.4 Risk of Heparin-Induced Thrombocytopenia with or without Thrombosis

limb ischemia, or death. Monitor thrombocytopenia of any degree closely.

after non-heparin alternative treatments are considered.

heparin, and 0.7% in patients given placebo in clinical trials.

5.6 Interchangeability with other Heparins

6.1 Clinical Trials Experience

Abdominal Surgery

Colorectal Surgery

Indications

Prophylaxis[†]

Hip Replacement Surgery without Extended

Peri-operative Period

Enoxaparin sodium injection 40 mg subcutaneously once a day for up to 21 days after discharge

The rates represent major bleeding on study medication up to 24 hours after last dose.

The rates represent major bleeding on study medication up to 12 hours after dose. Aspirin therapy was administered concurrently (100 to 325 mg per day).

drugs like enoxaparin sodium injection should be interpreted with caution

more units of blood products. Retroperitoneal, intraocular, and intracranial hemorrhages were always considered major

Local irritation, pain, hematoma, ecchymosis, and erythema may follow subcutaneous injection of enoxaparin sodium injection

Adverse Reactions in Patients Receiving Enoxaparin Sodium Injection for Prophylaxis or Treatment of DVT, PE

Enoxaparin Sodium Injection

Severe

Peri-operative Period

n = 288†

Severe Total

Extended Prophylaxis Period

Hip Replacement Surgery with Extended

Knee Replacement Surgery without

were also considered major hemorrhages.

Indication

Medical Patients during Acute Illness

Treatment of DVT and PE

therapy and continuing for up to 90 days.

Unstable Angina and Non-Q-Wave MI†

Acute ST-Segment Elevation

Major bleeding (including ICH)

Intracranial hemorrhages (ICH)

group, are provided below [see Tables 8 to 11].

Hemorrhage Anemia

Ecchymosis

Adverse Reaction

nemia

Edema

Dyspnea

Confusion

Diarrhea

Nausea

hrombocytopenia hrombocytopenia

Injection Site Hemorrhage

njection Site Pain

ematuria

Adverse Reaction

Peripheral edema

Mvocardial Infarction

Contraindications (4)]. Circulating antibodies may persist for several years.

Thrombocytopenia can occur with the administration of enoxaparin sodium injection.

5.7 Increased Risk of Thrombosis in Pregnant Women with Mechanical Prosthetic Heart Valves

The following serious adverse reactions are also discussed in other sections of the labeling:

Spinal/epidural hematomas [see Boxed Warning and Warnings and Precautions (5.1)]

Increased Risk of Hemorrhage [see Warnings and Precautions (5.1)]
 Thrombocytopenia [see Warnings and Precautions (5.5)]

rials of another drug and may not reflect the rates observed in clinical practice.

Reproductive and Developmental Toxicology

*Sections or subsections omitted from the full prescribing information are not listed.

Geriatric Patients: Monitor for increased risk of bleeding (8.5)

Low-Weight Patients: Observe for signs of bleeding (8.8)

adjust dosage as needed. (5.7)

8 USE IN SPECIFIC POPULATIONS

Lactation

Pediatric Use

Geriatric Use

Renal Impairment

Low-Weight Patients Obese Patients

Mechanism of Action Pharmacodynamics

Pharmacokinetics

16 HOW SUPPLIED/STORAGE AND HANDLING

17 PATIENT COUNSELING INFORMATION

8.2

8.5

8.7

8.8 8.9

10 OVERDOSAGE

11 DESCRIPTION

12.1

12.3

13.3

14.3

14 4

14.6

with platelet inhibitors

14 CLINICAL STUDIES

12 CLINICAL PHARMACOLOGY

13 NONCLINICAL TOXICOLOGY

Thrombocytopenia: Monitor platelet count closely. (5.5) • Interchangeability with other heparins: Do not exchange with heparin or other LMWHs (5.6) Increased Risk of Thrombosis in Pregnant Women with Mechanical Prosthetic Heart Valves: Women and their fetuses may be at increased risk. Monitor more frequently and - ADVERSE REACTIONS -

Non-major hemorrhagic events, primarily injection site ecchymosis and hematomas, were more frequently reported in patients treated with subcutaneous enoxaparin sodium injection WARNINGS AND PRECAUTIONS -

History of heparin-induced thrombocytopenia (HIT) within the past 100 days or in the presence of circulating antibodies (4)

Increased Risk of Hemorrhage: Monitor for sign of bleeding (5.1, 5.2, 5.3) Risk of Heparin-Induced Thrombocytopenia with or without Thrombosis. (5.4)

at least 0.5% in the enoxaparin sodium injection group are provided below [see Table 12]. Table 12 Serious Adverse Events Occurring at ≥0.5% Incidence in Enoxaparin Sodium Injection-Treated Patients with Unstable Angina or Non-Q-Wave Myocardial Infarction

Adverse Events in Enoxaparin Sodium Injection-Treated Patients with Unstable Angina or Non-Q-Wave Myocardial Infarctio

Non-hemorrhagic clinical events reported to be related to enoxaparin sodium injection therapy occurred at an incidence of ≤1%.

Serious adverse events with enoxaparin sodium injection or heparin in a clinical trial in patients with unstable angina or non-Q-wave myocardial infarction that occurred at a rate of

Adverse Event	Enoxaparin Sodium Injection 1 mg/kg q12h subcutaneously n = 1578 n (%)	Heparin aPTT Adjusted Intravenous Therapy n = 1529 n (%)
Atrial fibrillation	11 (0.70)	3 (0.20)
Heart failure	15 (0.95)	11 (0.72)
Lung edema	11 (0.70)	11 (0.72)
Pneumonia	13 (0.82)	9 (0.59)
Adverse Reactions in Enoxaparin Sodium Injection	n-Treated Patients with Acute ST-Segment Elevation Myoc	cardial Infarction
In a clinical trial in patients with acute ST-segment ele-	vation myocardial infarction, thrombocytopenia occurred at a r	ate of 1.5%.

Dosing Regimer

6.2 Postmarketing Experienc

The following adverse reactions have been identified during post approval use of enoxaparin sodium injection. Because these reactions are reported voluntarily from a population of uncertain size, it is not always possible to reliably estimate their frequency or establish a causal relationship to drug exposure

There have been reports of epidural or spinal hematoma formation with concurrent use of enoxaparin sodium injection and spinal/epidural anesthesia or spinal puncture. The majority of patients had a post-operative indwelling epidural catheter placed for analgesia or received additional drugs affecting hemostasis such as NSAIDs. Many of the epidural or spinal hematomas caused neurologic injury, including long-term or permanent paralysis. Local reactions at the injection site (e.g. nodules, inflammation, oozing), systemic allergic reactions (e.g. pruritus, urticaria, anaphylactic/anaphylactoid reactions including shock), vesiculobullous rash, cases of hypersensitivity cutaneous vasculitis, purpura, skin necrosis (occurring at either the injection site or distant from the injection site), thrombocytosis and thrombocytopenia with thrombosis [see Warnings and Precautions (5.5)] have been reported.

Cases of hyperkalemia have been reported. Most of these reports occurred in patients who also had conditions that tend toward the development of hyperkalemia (e.g., renal dysfunction, concomitant potassium-sparing drugs, administration of potassium, hematoma in body tissues). Very rare cases of hyperlipidemia have also been reported, with one case of hyperlipidemia, with marked hypertriglyceridemia, reported in a diabetic pregnant woman; causality has not been determined.

Cases of headache, hemorrhagic anemia, eosinophilia, alopecia, hepatocellular and cholestatic liver injury have been reported.

Osteoporosis has also been reported following long-term therapy. 7 DRUG INTERACTION

Whenever possible, agents which may enhance the risk of hemorrhage should be discontinued prior to initiation of enoxaparin sodium injection therapy. These agents include medications such as: anticoagulants, platelet inhibitors including acetylsalicylic acid, salicylates, NSAIDs (including ketorolac tromethamine), dipyridamole, or sulfinpyrazone. If coadministration is essential, conduct close clinical and laboratory monitoring [see Warnings and Precautions (5.1)].

8.1 Pregnancy Placental transfer of enoxaparin was observed in the animal studies. Human data from a retrospective cohort study, which included 693 live births, suggest that enoxaparin does not increase the risk of major developmental abnormalities (see Data). Based on animal data, enoxaparin sodium injection is not predicted to increase the risk of major developmental

8 USE IN SPECIFIC POPULATIONS

from the underlying maternal condition.

tudied for long-term use in this patient po

8.4 Pediatric Use

Adverse outcomes in pregnancy occur regardless of the health of the mother or the use of medications. The estimated background risk of major birth defects and miscarriage for the indicated populations is unknown. In the U.S. general population, the estimated background risk of major birth defects and miscarriage in clinically recognized pregnancies is 2% to 4% and 15% to 20%, respectively.

Clinical Considerations Pregnancy alone confers an increased risk for thromboembolism that is even higher for women with thromboembolic disease and certain high-risk pregnancy conditions. While no adequately studied, pregnant women with mechanical prosthetic heart valves may be at even higher risk for thrombosis [see Warnings and Precautions (5.7) and Use in Specific Populations (8.6)]. Pregnant women with thromboembolic disease, including those with mechanical prosthetic heart valves and those with inherited or acquired thrombophilias, have an increased risk of other maternal complications and fetal loss regardless of the type of anticoagulant used.

All patients receiving anticoagulants, including pregnant women, are at risk for bleeding. Pregnant women receiving enoxaparin sodium injection should be carefully monitored for evidence of bleeding or excessive anticoagulation. Consideration for use of a shorter acting anticoagulant should be specifically addressed as delivery approaches [see Boxed Warning]. Hemorrhage can occur at any site and may lead to death of mother and/or fetus. Pregnant women should be apprised of the potential hazard to the fetus and the mother

if enoxaparin sodium injection is administered during pregnancy. It is not known if monitoring of anti-Factor Xa activity and dose adjustment (by weight or anti-Factor Xa activity) of enoxaparin sodium injection affect the safety and the efficacy of the drug during pregnancy

• Human Data - There are no adequate and well-controlled studies in pregnant women. A retrospective study reviewed the records of 604 women who used enoxaparin sodium injection during pregnancy. A total of 624 pregnancies resulted in 693 live births. There were 72 hemorrhagic events (11 serious) in 63 women. There were 14 cases of neonatal hemorrhage. Major congenital anomalies in live births occurred at rates (2.5%) similar to background rates. There have been postmarketing reports of fetal death when pregnant women received enoxaparin sodium injection. Causality for these cases has not been determined. Insufficient data, the underlying disease, and the possibility of inadequate anticoagulation complicate the evaluation of these cases

A clinical study using enoxaparin sodium injection in pregnant women with mechanical prosthetic heart valves has been conducted [see Warnings and Precautions (5.7)]. Animal Data - Teratology studies have been conducted in pregnant rats and rabbits at subcutaneous doses of enoxaparin up to 15 times the recommended human dose (by mparison with 2 mg/kg as the maximum recommended daily dose). There was no evidence of teratogenic effects or fetotoxicity due to enoxaparin. Because animal reproduction

studies are not always predictive of human response, this drug should be used during pregnancy only if clearly needed. 8.2 Lactation

It is unknown whether enoxaparin sodium injection is excreted in human milk. In lactating rats, the passage of enoxaparin or its metabolites in the milk is very limited. There is no

information available on the effect of enoxaparin or its metabolites on the breastfed child, or on the milk production. The developmental and health benefits of breastfeeding should be considered along with the mother's clinical need for enoxaparin sodium injection and any potential adverse effects on the breastfed child from enoxaparin sodium injection or

Prevention of Deep Vein Thrombosis in Hip, Knee and Abdominal Surgery; Treatment of Deep Vein Thrombosis, Prevention of Ischemic Complications of Unstable Angina and Non-Q-Wave Myocardial Infarction Over 2800 patients, 65 years and older, have received enoxaparin sodium injection in clinical trials. The efficacy of enoxaparin sodium injection in the geriatric (≥65 years) was

and interactive reports) has not revealed additional interactives in the safety of the and Precautions (2.6) and Clinical Pharmacology (12.3)]. Treatment of Acute ST-Segment Elevation Myocardial Infarction In the clinical study for treatment of acute ST-segment elevation myocardial infarction, there was no evidence of difference in efficacy between patients ≥75 years of age (n = 1241)

have received enoxaparin sodium injection for thromboprophylaxis. Some of these cases were pregnant women in whom thrombosis led to maternal and fetal deaths. Insufficien data, the underlying disease and the possibility of inadequate anticoagulation complicate the evaluation of these cases. Pregnant women with mechanical prosthetic heart valves may be at higher risk for thromboembolism [see Warnings and Precautions (5.7)]. n patients with renal impairment, there is an increase in exposure of enoxaparin sodium. All such patients should be observed carefully for signs and symptoms of bleeding

The use of enoxaparin sodium injection has not been adequately studied for thromboprophylaxis in patients with mechanical prosthetic heart valves and has not been adequately

8.8 Low-Weight Patients An increase in exposure of enoxaparin sodium with prophylactic dosages (non-weight adjusted) has been observed in low-weight women (<45 kg) and low-weight men (<57 kg). Observe low-weight patients frequently for signs and symptoms of bleeding [see Clinical Pharmacology (12.3)].

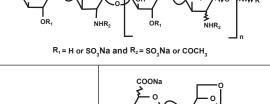
Obese patients are at higher risk for thromboembolism. The safety and efficacy of prophylactic doses of enoxaparin sodium injection in obese patients (BMI >30 kg/m²) has not been

fully determined and there is no consensus for dose adjustment. Observe these patients carefully for signs and symptoms of thromboemboli 10 OVERDOSAGE Accidental overdosage following administration of enoxagarin sodium injection may lead to hemorrhagic complications. Injected enoxagarin sodium injection may be largely neutralized by the slow intravenous injection of protamine sulfate (1% solution). The dose of protamine sulfate should be equal to the dose of enoxaparin sodium injection injected:

1 mg protamine sulfate should be administered to neutralize 1 mg enoxaparin sodium injection if enoxaparin sodium was administered in the previous 8 hours. An infusion of 0.5 mg protamine per 1 mg of enoxaparin sodium may be administered if enoxaparin sodium was administered greater than 8 hours previous to the protamine administration, or if it has been determined that a second dose of protamine is required. The second infusion of 0.5 mg protamine sulfate per 1 mg of enoxaparin sodium injection may be administered If at least 12 hours have elapsed since the last enoxaparin sodium injection, protamine administration may not be required; however, even with higher doses of protamine, the aPTT

often resembling anaphylaxis, have been reported with protamine sulfate, it should be given only when resuscitation techniques and treatment of anaphylactic shock are readily available. For additional information consult the labeling of protamine sulfate injection products.

structure contains an 1,6 anhydro derivative on the reducing end of the polysaccharide chain. The drug substance is the sodium salt. The average molecular weight is about 4500 daltons. The molecular weight distribution is: 2000 to 8000 daltons ≥68%



100 - X n =1 to 21 * X = Percent of polysaccharide chain containing 1,6 anhydro derivative on the reducing end Enoxaparin sodium injection 100 mg/mL concentration contains 10 mg enoxaparin sodium (approximate anti-Factor Xa activity of 1000 IU [with reference to the W.H.O. First cular Weight Heparin Reference Standard]) per 0.1 mL Water for Injection Enoxaparin sodium injection 150 mg/mL concentration contains 15 mg enoxaparin sodium (approximate anti-Factor Xa activity of 1500 IU [with reference to the W.H.O. First International Low Molecular Weight Heparin Reference Standard]) per 0.1 mL Water for Injection.

The enoxaparin sodium injection, USP prefilled syringes and graduated prefilled syringes are preservative-free and intended for use only as a single-dose injection [see Dosage and

In humans, enoxaparin given at a dose of 1.5 mg/kg subcutaneously is characterized by a higher ratio of anti-Factor Xa to anti-Factor Ila activity (mean ± SD, 14.0 ± 3.1) (based on areas under anti-Factor activity versus time curves) compared to the ratios observed for heparin (mean ± SD, 1.22 ± 0.13). Increases of up to 1.8 times the control values were

Absorption Pharmacokinetic trials were conducted using the 100 mg/mL formulation. Maximum anti-Factor Xa and anti-thrombin (anti-Factor IIa) activities occur 3 to 5 hours after subcutaneous

A 30 mg intravenous bolus immediately followed by 1 mg/kg subcutaneously every 12 hours provided initial peak anti-Factor Xa levels of 1.16 IU/mL (n=16) and average exposure responding to 84% of steady-state levels. Steady state is achieved on the second day of treatment. Enoxaparin pharmacokinetics appears to be linear over the recommended dosage ranges [see Dosage and Administration (2)]. After repeated subcutaneous administration of 40 mg once daily and 1.5 mg/kg once-daily regimens in healthy volunteers, the steady-state is reached on day 2 with an average exposure ratio about 15% higher than after a single dose. Steady state enoxaparin activity levels are well predicted by single-dose pharmacokinetics. After repeated subcutaneous administration of the 1 mg/kg twice-daily regimer the steady state is reached from day 4 with mean exposure about 65% higher than after a single dose and mean peak and trough levels of about 1.2 and 0.52 IU/mL, respectively. Based on enoxaparin sodium pharmacokinetics, this difference in steady state is expected and within the therapeutic range.

1.37 (±0.23) 19 (±5) 100 mg/mL 0.23 (±0.05) 105 (±17) Γ_{max} (IU/mL or Δ sec) 1.45 (±0.22) 0.26 (±0.05) 111 (±17) 22 (±7) 200 mg/mL 90% CI 3 (2-6) 2.5 (2-4.5) 100 mg/mL 4 (2-5) 3(2-4.5)

		200 mg/mL	3.5 (2–6)	4.5 (2.5–6)	3.3 (2–5)	3 (2–5)
	AUC (ss)	100 mg/mL	14.26 (±2.93)	1.54 (±0.61)	1321 (±219)	
	(h*IU/mL or h* Δ sec)	200 mg/mL	15.43 (±2.96)	1.77 (±0.67)	1401 (±227)	
-		90% CI	105–112%		103–109%	
	* Means ± SD at Day 5 and 90% † Median (range)	6 Confidence Interval (CI) of th	e ratio			
	Distribution					
r.	The volume of distribution of anti-	-Factor Xa activity is about 4.3	L.			
	Elimination					
nce	Following intravenous dosing, the					

Enoxaparin sodium is primarily metabolized in the liver by desulfation and/or depolymerization to lower molecular weight species with much reduced biological potency. Renal clearance of active fragments represents about 10% of the administered dose and total renal excretion of active and non-active fragments 40% of the dose.

Apparent clearance and A derived from anti-Factor Xa values following single subcutaneous dosing (40 mg and 60 mg) were slightly higher in males than in females. The source

of the gender difference in these parameters has not been conclusively identified; however, body weight may be a contributing factor Apparent clearance and A_{max} derived from anti-Factor Xa values following single and multiple subcutaneous dosing in geriatric subjects were close to those observed in young subjects. Following once a day subcutaneous dosing of 40 mg enoxaparin, the Day 10 mean area under anti-Factor Xa activity versus time curve (AUC) was approximately 15% greater than the mean Day 1 AUC value [see Dosage and Administration (2.4) and Use in Specific Populations (8.5)].

A linear relationship between anti-Factor Xa plasma clearance and creatinine clearance at steady state has been observed, which indicates decreased clearance of enoxaparin

sodium in patients with reduced renal function. Anti-Factor Xa exposure represented by AUC, at steady state, is marginally increased in patients with creatinine clearance 50 to 80 mL/min and patients with creatinine clearance 30 to <50 mL/min after repeated subcutaneous 40 mg once-daily doses. In patients with severe renal impairment (creatinine clearance <30 mL/min), the AUC at steady state is significantly increased on average by 65% after repeated subcutaneous 40 mg once-daily doses [see Dosage and Administration]

(2.3) and Use in Specific Populations (8.7)].

Studies with enoxaparin in patients with hepatic impairment have not been conducted and the impact of hepatic impairment on the exposure to enoxaparin is unknown

After repeated subcutaneous 1.5 mg/kg once-daily dosing, mean AUC of anti-Factor Xa activity is marginally higher at steady state in obese healthy volunteers (BMI 30 to 48 kg/m²) compared to non-obese control subjects, while A_{max} is not increased.

No pharmacokinetic interaction was observed between enoxaparin and thrombolytics when administered concomitantly

found to have no effect on fertility or reproductive performance of male and female rats at subcutaneous doses up to 20 mg/kg/day or 141 mg/m²/day. The maximum human dose in clinical trials was 2.0 mg/kg/day or 78 mg/m²/day (for an average body weight of 70 kg, height of 170 cm, and body surface area of 1.8 m²).

Abdominal surgery patients at risk include those who are over 40 years of age, obese, undergoing surgery under general anesthesia lasting longer than 30 minutes or who have additional risk factors such as malignancy or a history of deep vein thrombosis (DVT) or pulmonary embolism (PE).

Enoxaparin Sodium Injection 40 mg daily subcut 5000 U q8h subcut n (%) n (%) All Treated Abdominal Surgery Patients 555 (100) 560 (100) Treatment Failures Total VTE* (%) 56 (10.1) 63 (11.3)

(95% CI: 7 to 12)

Table 14 Efficacy of Enoxaparin Sodium Injection in the Prophylaxis of Deep Vein Thrombosis following Abdominal Surgery

(95% CI[†]: 8 to 13) (95% CI: 9 to 14) DVT Only (%) (95% CI: 8 to 13)

when enoxaparin sodium injection was administered at doses of 1.5 mg/kg once a day or 1 mg/kg every 12 hours. The risk of enoxaparin sodium injection-associated bleeding increased with age. Serious adverse events increased with age for patients receiving enoxaparin sodium injection. Other clinical experience (including postmarketing surveillance and literature reports) has not revealed additional differences in the safety of enoxaparin sodium injection between geriatric and younger patients. Careful attention to dosing intervals

may remain more prolonged than following administration of heparin. In all cases, the anti-Factor Xa activity is never completely neutralized (maximum about 60%). Particular care should be taken to avoid overdosage with protamine sulfate. Administration of protamine sulfate can cause severe hypotensive and anaphylactoid reactions. Because fatal reactions,

>8000 daltons

X*= 15 to 25% n=0 to 20

Administration (2) and How Supplied/Storage and Handling (16)]. 12 CLINICAL PHARMACOLOGY 12.1 Mechanism of Action Enoxaparin is a low molecular weight heparin which has antithrombotic properties

seen in the thrombin time (TT) and the activated partial thromboplastin time (aPTT). Enoxaparin at a 1 mg/kg dose (100 mg/mL concentration), administered subcutaneously every 12 hours to patients in a large clinical trial resulted in aPTT values of 45 seconds or less in the majority of patients (n = 1607). A 30 mg intravenous bolus immediately followed by a 1 mg/kg subcutaneous administration resulted in aPTT post injection values of 50 seconds. The average aPTT prolongation value on Day 1 was about 16% higher than on Day 4. 12.3 Pharmacokinetics

injection of enoxaparin. Mean peak anti-Factor Xa activity was 0.16 IU/mL (1.58 mcg/mL) and 0.38 IU/mL (3.83 mcg/mL) after the 20 mg and the 40 mg clinically tested subcutaneous doses, respectively. Mean (n = 46) peak anti-Factor Xa activity was 1.1 IU/mL at steady state in patients with unstable angina receiving 1 mg/kg subcutaneously every 12 hours for 14 days. Mean absolute bioavailability of enoxaparin, after 1.5 mg/kg given subcutaneously, based on anti-Factor Xa activity is approximately 100% in healthy subjects

Although not studied clinically, the 150 mg/mL concentration of enoxaparin sodium is projected to result in anticoagulant activities similar to those of 100 mg/mL and 200 mg/mL concentrations at the same enoxaparin dose. When a daily 1.5 mg/kg subcutaneous injection of enoxaparin sodium was given to 25 healthy male and female subjects using a 100 mg/mL or a 200 mg/mL concentration the following pharmacokinetic profiles were obtained [see Table 13]. Table 13 Pharmacokinetic Parameters* After 5 Days of 1.5 mg/kg Subcutaneously Once Daily Doses of Enoxaparin Sodium Using 100 mg/mL or 200 mg/mL Concentrations Concentration aPTT

* The rates represent major bleeding (including ICH) up to 30 days.

† Bleedings were considered major if the hemorrhage caused a significant clinical event associated with a hemoglobin decrease by ≥ 5 g/dL. ICH were always considered major Asymptomatic increases in aspartate (AST [SGOT]) and alanine (ALT [SGPT]) aminotransferase levels greater than three times the upper limit of normal of the laboratory reference range have been reported in up to 6.1% and 5.9% of patients, respectively, during treatment with enoxaparin sodium injection. and 8 to 20% of anti-Factor Xa activity were recovered in urine in 24 hours. Elimination half-life based on anti-Factor Xa activity was 4.5 hours after a single subcutaneous dose to about 7 hours after repeated dosing. Significant anti-Factor Xa activity persists in plasma for about 12 hours following a 40 mg subcutaneous once a day dose. Since aminotransferase determinations are important in the differential diagnosis of myocardial infarction, liver disease, and pulmonary emboli, elevations that might be caused by Following subcutaneous dosing, the apparent clearance (CL/F) of enoxaparin is approximately 15 mL/mir

In a single study, elimination rate appeared similar, but AUC was two-fold higher than control population, after a single 0.25 or 0.5 mg/kg intravenous dose.

When non-weight-adjusted dosing was administered, it was found after a single-subcutaneous 40 mg dose, that anti-Factor Xa exposure is 52% higher in low-weight women (<45 kg) and 27% higher in low-weight men (<57 kg) when compared to normal weight control subjects [see Use in Specific Populations (8.8)].

No long-term studies in animals have been performed to evaluate the carcinogenic potential of enoxaparin. Enoxaparin was not mutagenic in *in vitro* tests, including the Ames test, mouse lymphoma cell forward mutation test, and human lymphocyte chromosomal aberration test, and the *in vivo* rat bone marrow chromosomal aberration test. Enoxaparin was

A single subcutaneous dose of 46.4 mg/kg enoxaparin was lethal to rats. The symptoms of acute toxicity were ataxia, decreased motility, dyspnea, cyanosis, and coma.

Teratology studies have been conducted in pregnant rats and rabbits at subcutaneous doses of enoxaparin up to 30 mg/kg/day corresponding to 211 mg/m²/day and 410 mg/m²/day in rats and rabbits respectively. There was no evidence of teratogenic effects or fetotoxicity due to enoxaparin. 14.1 Prophylaxis of Deep Vein Thrombosis following Abdominal Surgery in Patients at Risk for Thromboembolic Complications

In a double-blind, parallel group study of patients undergoing elective cancer surgery of the gastrointestinal, urological, or gynecological tract, a total of 1116 patients were enrolled in the study, and 1115 patients were treated. Patients ranged in age from 32 to 97 years (mean age 67 years) with 52.7% men and 47.3% women. Patients were 98% Caucasian, 1.1% Black, 0.4% Asian and 0.4% others. Enoxaparin sodium injection 40 mg subcutaneously, administered once a day, beginning 2 hours prior to surgery and continuing for a maximum of 12 days after surgery, was comparable to heparin 5000 U every 8 hours subcutaneously in reducing the risk of DVT. The efficacy data are provided below [see Table 14].

VTE = Venous thromboembolic events which included DVT, PE, and death considered to be thromboembolic in origin. CI = Confidence Interval

Safety and effectiveness of enoxaparin sodium injection in pediatric patients have not been established. Enoxaparin sodium injection should be used with care in patients with a bleeding diathesis, uncontrolled arterial hypertension or a history of recent gastrointestinal ulceration, diabetic Enoxaparin sodium injection is not approved for use in neonates or infants. Enoxaparin sodium injection may cause heparin-induced thrombocytopenia (HIT) or heparin-induced thrombocytopenia with thrombosis (HITTS). HITTS may lead to organ infarction, Use of enoxaparin sodium injection in patients with a history of immune-mediated HIT within the past 100 days or in the presence of circulating antibodies is contraindicated [see similar to that seen in younger patients (<65 years). The incidence of bleeding complications was similar between geriatric and younger patients when 30 mg every 12 hours or 40 mg once a day doses of enoxaparin sodium injection were employed. The incidence of bleeding complications was higher in geriatric patients as compared to younger patients Only use enoxaparin sodium injection in patients with a history of HIT if more than 100 days have elapsed since the prior HIT episode and no circulating antibodies are present.

Moderate thrombocytopenia (platelet counts between 100,000/mm³ and 50,000/mm³) occurred at a rate of 1.3% in patients given enoxaparin sodium injection, 1.2% in patients given and patients less than 75 years of age (n=9015). Patients ≥75 years of age did not receive a 30 mg intravenous bolus prior to the normal dosage regimen and had their subcutaneous dose adjusted to 0.75 mg/kg every 12 hours [see Dosage and Administration (2.4)]. The incidence of bleeding complications was higher in patients ≥65 years of age as compared to younger patients (<65 years) 8.6 Patients with Mechanical Prosthetic Heart Valves

> Because exposure of enoxaparin sodium is significantly increased in patients with severe renal impairment (creatinine clearance <30 mL/min), a dosage adjustment is recommended for therapeutic and prophylactic dosage ranges. No dosage adjustment is recommended in patients with creatinine clearance 30 to <50 mL/min and creatinine clearance 50 to 80 mL/min [see Dosage and Administration (2.3) and Clinical Pharmacology (12.3)]. In patients with renal failure, treatment with enoxaparin has been associated with the development of hyperkalemia [see Adverse Reactions (6.2)].

if the aPTT measured 2 to 4 hours after the first infusion remains prolonged

11 DESCRIPTION Enoxaparin sodium injection is a sterile aqueous solution containing enoxaparin sodium, a low molecular weight heparin. The pH of the injection is 5.5 to 7.5. Enoxaparin sodium is obtained by alkaline depolymerization of heparin benzyl ester derived from porcine intestinal mucosa. Its structure is characterized by a 2-O-sulfo-4-enepyranosuronic acid group at the non-reducing end and a 2-N,6-O-disulfo-D-glucosamine at the reducing end of the chain. About 20% (ranging between 15% and 25%) of the enoxaparin

STRUCTURAL FORMULA

Table 15 Efficacy of Enoxaparin Sodium Injection in the Prophylaxis of Deep Vein Thrombosis following Colorectal Surgery			
	Dosing Regimen		
Indication	Enoxaparin Sodium Injection 40 mg daily subcutaneously n (%)	Heparin 5000 U q8h subcutaneously n (%)	
All Treated Colorectal Surgery Patients	673 (100)	674 (100)	
Treatment Failures			
Total VTE* (%)	48 (7.1) (95% Cl [†] : 5 to 9)	45 (6.7) (95% CI: 5 to 9)	
DVT Only (%)	47 (7.0) (95% CI: 5 to 9)	44 (6.5) (95% CI: 5 to 8)	

* VTE = Venous thromboembolic events which included DVT, PE, and death considered to be thromboembolic in origin

14.2 Prophylaxis of Deep Vein Thrombosis following Hip or Knee Replacement Surgery

Enoxaparin sodium injection has been shown to reduce the risk of post-operative deep vein thrombosis (DVT) following hip or knee replacement surgery

In a double-blind study, enoxaparin sodium injection 30 mg every 12 hours subcutaneously was compared to placebo in patients with hip replacement. A total of 100 patients were randomized in the study and all patients were treated. Patients ranged in age from 41 to 84 years (mean age 67.1 years) with 45% men and 55% women. After hemostasis was established, treatment was initiated 12 to 24 hours after surgery and was continued for 10 to 14 days after surgery. The efficacy data are provided below [see Table 16].

Table 16 Efficacy of Enoxaparin	Sodium Injection in the Prophylaxis of Deep Vein Thrombosis	following Hip Replacement Surgery		
	Dosing Regimen			
	Enoxaparin Sodium Injection	Placebo		
	30 mg q12h subcutaneously	q12h subcutaneously		
Indication	n (%)	n (%)		

Indication	Enoxaparin Sodium Injection 30 mg q12h subcutaneously n (%)	Placebo q12h subcutaneously n (%)
All Treated Hip Replacement Patients	50 (100)	50 (100)
Treatment Failures		
Total DVT (%)	5 (10)*	23 (46)
Proximal DVT (%)	1 (2) [†]	11 (22)

A double-blind, multicenter study compared three dosing regimens of enoxaparin sodium injection in patients with hip replacement. A total of 572 patients were randomized in the study and 568 patients were treated. Patients ranged in age from 31 to 88 years (mean age 64.7 years) with 63% men and 37% women. Patients were 93% Caucasian, 6% Black, <1% Asian, and 1% others. Treatment was initiated within two days after surgery and was continued for 7 to 11 days after surgery. The efficacy data are provided below [see Table 17].

Table 17 Efficacy of Enoxaparin Sodium Injection in the Prophylaxis of Deep Vein Thrombosis following Hip Replacement Surgery				
	Dosing Regimen			
Indication	10 mg daily subcutaneously n (%)	30 mg q12h subcutaneously n (%)	40 mg daily subcutaneously n (%)	
All Treated Hip Replacement Patients	161 (100)	208 (100)	199 (100)	
Treatment Failures				
Total DVT (%)	40 (25)	22 (11)*	27 (14)	

17 (11)

* p value versus enoxaparin sodium injection 10 mg once a day = 0.0008

There was no significant difference between the 30 mg every 12 hours and 40 mg once a day regimens. In a double-blind study, enoxaparin sodium injection 30 mg every 12 hours subcutaneously was compared to placebo in patients undergoing knee replacement surgery. A total of 132 patients were randomized in the study and 131 patients were treated, of which 99 had total knee replacement and 32 had either unicompartmental knee replacement or tibial osteotomy. The 99 patients with total knee replacement ranged in age from 42 to 85 years (mean age 70.2 years) with 36.4% men and 63.6% women. After hemostasis was established, treatment was initiated 12 to 24 hours after surgery and was continued up to 15 days after surgery. The incidence of proximal and total DVT after surgery was significantly lower for enoxaparin sodium injection compared to placebo. The efficacy data are provided below [see Table 18].

8 (4)†

9 (5)

Table 18 Efficacy of Enoxaparin Sodium Injection in the Prophylaxis of Deep Vein Thrombosis Following Total Knee Replacement Surgery

	Dosing Regimen		
Indication	Enoxaparin Sodium Injection 30 mg q12h subcutaneously n (%)	Placebo q12h subcutaneously n (%)	
All Treated Total Knee Replacement Patients	47 (100)	52 (100)	
Treatment Failures			
Total DVT (%)	5 (11)* (95% CI ⁺ : 1 to 21)	32 (62) (95% CI: 47 to 76)	
Proximal DVT (%)	0 (0)‡ (95% Upper CL§: 5)	7 (13) (95% CI: 3 to 24)	

† CI = Confidence Interval p value versus placebo = 0.013

§ CL = Confidence Limit

Proximal DVT (%)

Additionally, in an open-label, parallel group, randomized clinical study, enoxaparin sodium injection 30 mg every 12 hours subcutaneously in patients undergoing elective knee replacement surgery was compared to heparin 5000 U every 8 hours subcutaneously. A total of 453 patients were randomized in the study and all were treated. Patients ranged in age from 38 to 90 years (mean age 68.5 years) with 43.7% men and 56.3% women. Patients were 92.5% Caucasian, 5.3% Black, and 0.6% others. Treatment was initiated after surgery and continued up to 14 days. The incidence of deep vein thrombosis was lower for enoxaparin sodium injection compared to heparin.

Extended Prophylaxis of Deep Vein Thrombosis following Hip Replacement Surgery: In a study of extended prophylaxis for patients undergoing hip replacement surgery, patients were treated, while hospitalized, with enoxaparin sodium injection 40 mg subcutaneously, initiated up to 12 hours prior to surgery for the prophylaxis of post-operative DVT. At the end of the peri-operative period, all patients underwent bilateral venography. In a double-blind design, those patients with no venous thromboembolic disease were randomized to a post-discharge regimen of either enoxaparin sodium injection 40 mg (n = 90) once a day subcutaneously or to placebo (n = 89) for 3 weeks. A total of 179 patients were randomized in the double-blind phase of the study and all patients were treated. Patients ranged in age from 47 to 87 years (mean age 69.4 years) with 57% men and 43% women In this population of patients, the incidence of DVT during extended prophylaxis was significantly lower for enoxaparin sodium injection compared to placebo. The efficacy data are provided below [see Table 19].

Table 19 Efficacy of Enoxaparin Sodium Injection in the Extended Prophylaxis of Deep Vein Thrombosis following Hip Replacement Surgery

	Post-Discharge Dosing Regimen		
Indication (Post-Discharge)	Enoxaparin Sodium Injection 40 mg daily subcutaneously n (%)	Placebo daily subcutaneously n (%)	
All Treated Extended Prophylaxis Patients	90 (100)	89 (100)	
Treatment Failures			
Total DVT (%)	6 (7)* (95% Cl [†] : 3 to 14)	18 (20) (95% CI: 12 to 30)	
Proximal DVT (%)	5 (6)‡ (95% CI: 2 to 13)	7 (8) (95% CI: 3 to 16)	

[‡] p value versus placebo = 0.537

In a second study, patients undergoing hip replacement surgery were treated, while hospitalized, with enoxaparin sodium injection 40 mg subcutaneously, initiated up to 12 hours prior to surgery. All patients were examined for clinical signs and symptoms of venous thromboembolic (VTE) disease. In a double-blind design, patients without clinical signs and symptoms of VTE disease were randomized to a post-discharge regimen of either enoxaparin sodium injection 40 mg (n = 131) once a day subcutaneously or to placebo (n = 131) for 3 weeks. A total of 262 patients were randomized in the study double-blind phase and all patients were treated. Patients ranged in age from 44 to 87 years (mean age 68.5 years) with 43.1% men and 56.9% women. Similar to the first study the incidence of DVT during extended prophylaxis was significantly lower for enoxaparin sodium injection compared to placebo, with a statistically significant difference in both total DVT (enoxaparin sodium injection 21 [16%] versus placebo 45 [34%]; p = 0.001) and proximal DVT (enoxaparin sodium injection 21 [16%] versus placebo 45 [34%]; p = 0.001) sodium injection 8 [6%] versus placebo 28 [21%]; p = <0.001).

14.3 Prophylaxis of Deep Vein Thrombosis in Medical Patients with Severely Restricted Mobility during Acute Illness

In a double blind multicenter, parallel group study, enoxaparin sodium injection 20 mg or 40 mg once a day subcutaneously was compared to placebo in the prophylaxis of deep vein thrombosis (DVT) in medical patients with severely restricted mobility during acute illness (defined as walking distance of <10 meters for ≤3 days). This study included patients with heart failure (NYHA Class III or IV); acute respiratory failure or complicated chronic respiratory insufficiency (not requiring ventilatory support); acute infection (excluding septic shock); or acute rheumatic disorder [acute lumbar or sciatic pain, vertebral compression (due to osteoporosis or tumor), acute arthritic episodes of the lower extremities]. A total of 1102 patients were enrolled in the study, and 1073 patients were treated. Patients ranged in age from 40 to 97 years (mean age 73 years) with equal proportions of men and women. Treatment continued for a maximum of 14 days (median duration 7 days). When given at a dose of 40 mg once a day subcutaneously, enoxaparin sodium injection significantly reduced the incidence of DVT as compared to placebo. The efficacy data are provided below [see Table 20].

Table 20 Efficacy of Enoxaparin Sodium Injection in the Prophylaxis of Deep Vein Thrombosis in Medical Patients with Severely Restricted Mobility during Acute Illness

Indication	Enoxaparin Sodium Injection 20 mg daily subcutaneously n (%)	Enoxaparin Sodium Injection 40 mg daily subcutaneously n (%)	Placebo n (%)
All Treated Medical Patients during Acute Illness	351 (100)	360 (100)	362 (100)
Treatment Failure*			
Total VTE [†] (%)	43 (12.3)	16 (4.4)	43 (11.9)
Total DVT (%)	43 (12.3) (95% CI [‡] 8.8 to 15.7)	16 (4.4) (95% CI [‡] 2.3 to 6.6)	41 (11.3) (95% CI [‡] 8.1 to 14.6)
Proximal DVT (%)	13 (3.7)	5 (1.4)	14 (3.9)

Treatment failures during therapy, between Days 1 and 14 VTE = Venous thromboembolic events which included DVT, PE, and death considered to be thromboembolic in origin ‡ CI = Confidence Interval

At approximately 3 months following enrollment, the incidence of venous thromboembolism remained lower in the enoxaparin sodium injection 40 mg treatment group versus the

14.4 Treatment of Deep Vein Thrombosis with or without Pulmonary Embolism

In a multicenter, parallel group study, 900 patients with acute lower extremity deep vein thrombosis (DVT) with or without pulmonary embolism (PE) were randomized to an inpatient (hospital) treatment of either (i) enoxaparin sodium injection 1.5 mg/kg once a day subcutaneously, (ii) enoxaparin sodium injection 1 mg/kg every 12 hours subcutaneously, or (iii) heparin intravenous bolus (5000 IU) followed by a continuous infusion (administered to achieve an aPTT of 55 to 85 seconds). A total of 900 patients were randomized in the study and all patients were treated. Patients ranged in age from 18 to 92 years (mean age 60.7 years) with 54.7% men and 45.3% women. All patients also received warfarin sodium (dose adjusted according to PT to achieve an International Normalization Ratio [INR] of 2.0 to 3.0), commencing within 72 hours of initiation of enoxaparin sodium injection or standard heparin therapy, and continuing for 90 days. Enoxaparin sodium injection or standard heparin therapy was administered for a minimum of 5 days and until the targeted warfarin sodium INR was achieved. Both enoxaparin sodium injection regimens were equivalent to standard heparin therapy in reducing the risk of recurrent venous thromboembolism (DVT and/or PE). The efficacy data are provided below [see Table 21].

Table 21 Efficacy of Enoxaparin Sodium Injection in Treatment of Deep Vein Thrombosis with or without Pulmonary Embolism

Enoxaparin Sodium Injection 1.5 mg/kg daily subcutaneously n (%)	Enoxaparin Sodium Injection 1 mg/kg q12h subcutaneously n (%)	Heparin aPTT Adjusted Intravenous Therapy n (%)
298 (100)	312 (100)	290 (100)
13 (4.4) [‡]	9 (2.9) [‡]	12 (4.1)
11 (3.7)	7 (2.2)	8 (2.8)
9 (3.0)	6 (1.9)	7 (2.4)
2 (0.7)	2 (0.6)	4 (1.4)
	1.5 mg/kg daily subcutaneously n (%) 298 (100) 13 (4.4) [‡] 11 (3.7) 9 (3.0)	1.5 mg/kg daily subcutaneously n (%) 1 mg/kg q12h subcutaneously n (%) 298 (100) 312 (100) 13 (4.4) [‡] 9 (2.9) [‡] 11 (3.7) 7 (2.2) 9 (3.0) 6 (1.9)

VTE = venous thromboembolic event (DVT and/or PE). The 95% Confidence Intervals for the treatment differences for total VTE were: Enoxaparin sodium injection once a day versus heparin (-3.0 to 3.5) Enoxaparin sodium injection every 12 hours versus heparin (-4.2 to 1.7)

Similarly, in a multicenter, open-label, parallel group study, patients with acute proximal DVT were randomized to enoxaparin sodium injection or heparin. Patients who could not receive outpatient therapy were excluded from entering the study. Outpatient exclusion criteria included the following: inability to receive outpatient heparin therapy because of associated co-morbid conditions or potential for non-compliance and inability to attend follow-up visits as an outpatient because of geographic inaccessibility. Eligible patients could be treated in the hospital, but ONLY enoxaparin sodium injection patients were permitted to go home on therapy (72%). A total of 501 patients were randomized in the study and all patients were treated. Patients ranged in age from 19 to 96 years (mean age 57.8 years) with 60.5% men and 39.5% women. Patients were randomized to either enoxaparin sodium injection 1 mg/kg every 12 hours subcutaneously or heparin intravenous bolus (5000 IU) followed by a continuous infusion administered to achieve an aPTT of 60 to 85 seconds (in-patient treatment). All patients also received warfarin sodium as described in the previous study. Enoxaparin sodium injection or standard heparin therapy was administered for a minimum of 5 days. Enoxaparin sodium injection was equivalent to standard heparin therapy in reducing the risk of recurrent venous thromboembolism. The efficacy data are provided below [see Table 22].

Table 22 Efficacy of Enoxaparin Sodium Injection in Treatment of Deep Vein Thrombosis Dosina Regimen

Indication	Enoxaparin Sodium Injection 1 mg/kg q12h subcutaneously n (%)	Heparin aPTT Adjusted Intravenous Therapy n (%)
All Treated DVT Patients	247 (100)	254 (100)
Patient Outcome		
Total VTE† (%)	13 (5.3) [‡]	17 (6.7)
DVT Only (%)	11 (4.5)	14 (5.5)
Proximal DVT (%)	10 (4.0)	12 (4.7)
PE (%)	2 (0.8)	3 (1.2)

All patients were also treated with warfarin sodium commencing on the evening of the second day of enoxaparin sodium injection or standard heparin therapy † VTE = venous thromboembolic event (deep vein thrombosis [DVT] and/or pulmonary embolism [PE]) ‡ The 95% Confidence Intervals for the treatment difference for total VTE was: enoxaparin sodium injection versus heparin (- 5.6 to 2.7).

14.5 Prophylaxis of Ischemic Complications in Unstable Angina and Non-Q-Wave Myocardial Infarction In a multicenter, double-blind, parallel group study, patients who recently experienced unstable angina or non-Q-wave myocardial infarction were randomized to either enoxaparin sodium injection 1 mg/kg every 12 hours subcutaneously or heparin intravenous bolus (5000 U) followed by a continuous infusion (adjusted to achieve an aPTT of 55 to 85 seconds). A total of 3171 patients were enrolled in the study, and 3107 patients were treated. Patients ranged in age from 25 to 94 years (median age 64 years), with 33.4% of patient's female and 66.6% male. Race was distributed as follows: 89.8% Caucasian, 4.8% Black, 2.0% Asian, and 3.5% other. All patients were also treated with aspirin 100 to 325 mg per day. Treatment was initiated within 24 hours of the event and continued until clinical stabilization, revascularization procedures, or hospital discharge, with a maximal duration of 8 days of therapy. The combined incidence of the triple endpoint of death, myocardial infarction, or recurrent angina was lower for enoxaparin sodium injection compared with heparin therapy at 14 days after initiation of treatment. The lower incidence of the triple endpoint was sustained up to 30 days after initiation of treatment. These results were observed in an analysis of both all-randomized and all-treated patients. The efficacy data are provided below [see Table 23].

Table 23 Efficacy of Enoxaparin Sodium Injection in the Prophylaxis of Ischemic Complications in Unstable Angina and Non-Q-Wave Myocardial Infarction nbined endpoint of death, myocardial infarction, or recurrent angina)

	Dosing Re	gimen*		İ	
Indication	Enoxaparin Sodium Injection 1mg/kg q12h subcutaneously n (%)	Heparin aPTT Adjusted Intravenous Therapy n (%)	Reduction (%)	p Value	
All Treated Unstable Angina and Non-Q-Wave MI Patients	1578 (100)	1529 (100)			
Time point [†]					
48 Hours	96 (6.1)	112 (7.3)	1.2	0.120	
14 Days	261 (16.5)	303 (19.8)	3.3	0.017	
30 Days	313 (19.8)	358 (23.4)	3.6	0.014	

[†] Evaluation time points are after initiation of treatment. Therapy continued for up to 8 days (median duration of 2.6 days). The combined incidence of death or myocardial infarction at all time points was lower for enoxaparin sodium injection compared to standard heparin therapy but did not achieve statistical significance. The efficacy data are provided below [see Table 24].

Table 24 Ellicacy of Elloxapariii of	odium Injection in the Prophylaxis of Ische (combined endpoint of dea Dosing Re	Angina and Non-Q-Wave	In your diam in arction		
Indication	Enoxaparin Sodium Injection 1mg/kg q12h subcutaneously n (%) Heparin aPTT Adjusted Intravenous Therapy n (%)		Reduction (%)	p Value	
All Treated Unstable Angina and Non-Q-Wave MI Patients	1578 (100)	1529 (100)			
Γime point [†]					
48 Hours	16 (1.0)	20 (1.3)	0.3	0.126	
14 Days	76 (4.8)	93 (6.1)	1.3	0.115	
30 Days	96 (6.1)	118 (7.7)	1.6	0.069	

30 Days All patients were also treated with aspirin 100 to 325 mg per day. Evaluation time points are after initiation of treatment. Therapy continued for up to 8 days (median duration of 2.6 days). In a survey one year following treatment, with information available for 92% of enrolled patients, the combined incidence of death, myocardial infarction, or recurrent angina remained

lower for enoxaparin sodium injection versus heparin (32.0% vs 35.7%). Urgent revascularization procedures were performed less frequently in the enoxaparin sodium injection group as compared to the heparin group, 6.3% compared to 8.2% at 30 days (p = 0.047).

14.6 Treatment of Acute ST-Segment Elevation Myocardial Infarction

enoxaparin sodium injection was 6.6 days. The mean treatment duration of unfractionated heparin was 54 hours.

In a multicenter, double-blind, double-dummy, parallel-group study, patients with acute ST-segment elevation myocardial infarction (STEMI) who were to be hospitalized within 6 hours of onset and were eligible to receive fibrinolytic therapy were randomized in a 1:1 ratio to receive either enoxaparin sodium injection or unfractionated heparin.

Study medication was initiated between 15 minutes before and 30 minutes after the initiation of fibrinolytic therapy. Unfractionated heparin was administered beginning with an intravenous bolus of 60 U/kg (maximum 4000 U) and followed with an infusion of 12 U/kg per hour (initial maximum 1000 U per hour) that was adjusted to maintain an aPTT of 1.5 to 2 times the control value. The intravenous infusion was to be given for at least 48 hours. The enoxaparin sodium injection dosing strategy was adjusted according to the patient's age and renal function. For patients younger than 75 years of age, enoxaparin sodium injection was given as a single 30 mg intravenous bolus plus a 1 mg/kg subcutaneous dose followed by a subcutaneous injection of 1 mg/kg every 12 hours. For patients at least 75 years of age, the intravenous bolus was not given and the subcutaneous dose was reduced to 0.75 mg/kg every 12 hours. For patients with severe renal insufficiency (estimated creatinine clearance of less than 30 mL per minute), the dose was to be modified to 1 mg/kg every 24 hours. The subcutaneous injections of enoxaparin sodium injection were given until hospital discharge or for a maximum of eight days (whichever came first). The mean treat

the PCI was to be performed on enoxaparin sodium injection (no switch) using the regimen established in previous studies, i.e. no additional dosing, if the last subcutaneous administration was less than 8 hours before balloon inflation, intravenous bolus of 0.3 mg/kg enoxaparin sodium injection if the last subcutaneous administration was more than 8 hours before balloon inflation All patients were treated with aspirin for a minimum of 30 days. Eighty percent of patients received a fibrin-specific agent (19% tenecteplase, 5% reteplase and 55% alteplase) and

When percutaneous coronary intervention was performed during study medication period, patients received antithrombotic support with blinded study drug. For patients on enoxaparin,

Among 20,479 patients in the ITT population, the mean age was 60 years, and 76% were male. Racial distribution was: 87% Caucasian, 9.8% Asian, 0.2% Black, and 2.8% other. Medical history included previous MI (13%), hypertension (44%), diabetes (15%) and angiographic evidence of CAD (5%). Concomitant medication included aspirin (95%), betablockers (86%), ACE inhibitors (78%), statins (70%) and clopidogrel (27%). The MI at entry was anterior in 43%, non-anterior in 56%, and both in 1%.

The primary efficacy endpoint was the composite of death from any cause or myocardial re-infarction in the first 30 days after randomization. Total follow-up was one year.

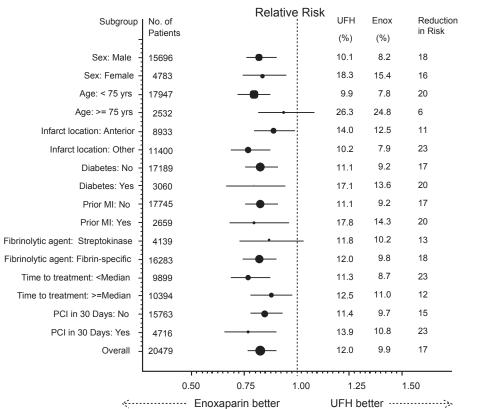
The rate of the primary efficacy endpoint (death or myocardial re-infarction) was 9.9% in the enoxaparin sodium injection group, and 12% in the unfractionated heparin group, a 17% reduction in the relative risk, (P=0.000003) [see Table 25]. Table 25 Efficacy of Enoxaparin Sodium Injection in the Treatment of Acute ST-Segment Elevation Myocardial Infarction

Enoxaparin Relative Risk P Value (N=10.256) (N=10,223) (95% CI) Outcome at 48 hours n (%) n (%) Death or Myocardial Re-infarction 478 (4.7) 531 (5.2) 0.90 (0.80 to 1.01) 0.08 Death 383 (3.7) 390 (3.8) 0.98 (0.85 to 1.12) 0.76 102 (1.0) 0.65 (0.51 to 0.84) Myocardial Re-infarction 156 (1.5) < 0.001 Urgent Revascularization 74 (0.7) 96 (0.9) 0.77 (0.57 to 1.04) 0.09 Death or Myocardial Re-infarction or Urgent Revascularization 548 (5.3) 622 (6.1) 0.88 (0.79 to 0.98) 0.02 Outcome at 8 Days 740 (7.2) Death or Myocardial Re-infarction 954 (9.3) 0.77 (0.71 to 0.85) < 0.001 559 (5.5) 605 (5.9) Death 0.92 (0.82 to 1.03) 0.15 Myocardial Re-infarction 204 (2.0) 379 (3.7) 0.54 (0.45 to 0.63) < 0.001 145 (1.4) 0.59 (0.48 to 0.72) Urgent Revascularization 247 (2.4) < 0.001 Death or Myocardial Re-infarction or Urgent Revascularization 874 (8.5) 1181 (11.6) 0.74 (0.68 to 0.80) < 0.001 Outcome at 30 Days Primary efficacy endpoint 1017 (9.9) 1223 (12.0) 0.83 (0.77 to 0.90) 0.000003 (Death or Myocardial Re-infarction) 708 (6.9) 765 (7.5) 0.92 (0.84 to 1.02) 0.11 508 (5.0) Myocardial Re-infarction 352 (3.4) 0.69 (0.60 to 0.79) < 0.001 286 (2.8) 0.74 (0.62 to 0.88) < 0.001 1479 (14.5) 1199 (11.7) 0.81 (0.75 to 0.87)

Note: Urgent revascularization denotes episodes of recurrent myocardial ischemia (without infarction) leading to the clinical decision to perform coronary revas

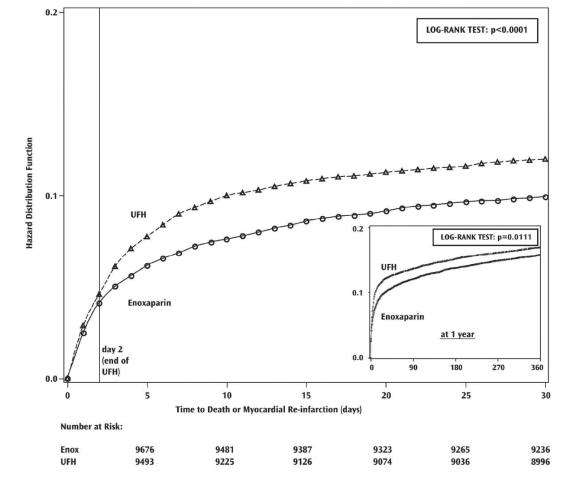
the same hospitalization. CI denotes confidence intervals.

The beneficial effect of enoxaparin sodium injection on the primary endpoint was consistent across key subgroups including age, gender, infarct location, history of diabetes, history of prior myocardial infarction, fibrinolytic agent administered, and time to treatment with study drug (see Figure 1); however, it is necessary to interpret such subgroup analyses with caution. Figure 1. Relative Risks of and Absolute Event Rates for the Primary Endpoint at 30 Days in Various Subgroups



* The primary efficacy end point was the composite of death from any cause or myocardial re-infarction in the first 30 days. The overall treatment effect of enoxaparin sodium injection as compared to the unfractionated heparin (UFH) is shown at the bottom of the figure. For each subgroup, the circle is proportional to the number and represents the point estimate of the treatment effect and the horizontal lines represent the 95% confidence intervals. Fibrin-specific fibrinolytic agents included alteplase, tenecteplase, and reteplase. Time to nent indicates the time from the onset of symptoms to the administration of study drug (median: 3.2 hours). The beneficial effect of enoxaparin sodium injection on the primary endpoint observed during the first 30 days was maintained over a 12-month follow-up period (see Figure 2).

Figure 2 - Kaplan-Meier Plot -Death or Myocardial Re-infarction at 30 days -ITT Population



There is a trend in favor of enoxaparin sodium injection during the first 48 hours, but most of the treatment difference is attributed to a step increase in the event rate in the UFH group at 48 hours (seen in Figure 2), an effect that is more striking when comparing the event rates just prior to and just subsequent to actual times of discontinuation. These results provide evidence that UFH was effective and that it would be better if used longer than 48 hours. There is a similar increase in endpoint event rate when enoxaparin sodium injection was discontinued, suggesting that it too was discontinued too soon in this study. The rates of major hemorrhages (defined as requiring 5 or more units of blood for transfusion, or 15% drop in hematocrit or clinically overt bleeding, including intracranial hemorrhage) at 30 days were 2.1% in the enoxaparin sodium injection group and 1.4% in the unfractionated heparin group. The rates of intracranial hemorrhage at 30 days were 0.8% in the enoxaparin sodium injection group and 0.7% in the unfractionated heparin group. The 30-day rate of the composite endpoint of death, myocardial re-infarction or ICH (a measure

16 HOW SUPPLIED/STORAGE AND HANDLING Enoxaparin sodium injection, USP is available in two concentrations [see Tables 26 and 27]:

of net clinical benefit) was significantly lower in the enoxaparin sodium injection group (10.1%) as compared to the heparin group (12.2%)

Table 26 100 mg/mL Concentration

Anti-Xa Activity†	Unit of Sale	Syringe Label Color	Each
3000 IU	NDC# 63323-559-93 10 syringes per carton	Medium Blue	NDC# 63323-559-21
4000 IU	NDC# 63323-564-97 10 syringes per carton	Yellow	NDC# 63323-564-21
6000 IU	NDC# 63323-566-98 10 syringes per carton	Orange	NDC# 63323-566-21
8000 IU	NDC# 63323-584-99 10 syringes per carton	Brown	NDC# 63323-584-21
10,000 IU	NDC# 63323-586-96 10 syringes per carton	Black	NDC# 63323-586-21
	3000 IU 4000 IU 6000 IU 8000 IU	3000 IU NDC# 63323-559-93 10 syringes per carton 4000 IU NDC# 63323-564-97 10 syringes per carton NDC# 63323-566-98 10 syringes per carton NDC# 63323-584-99 10 syringes per carton NDC# 63323-586-96	Anti-Xa Activity†

100 mg graduated prefilled syringes each contain 10 mg enoxaparin sodium per 0.1 mL Water for Injection Approximate anti-Factor Xa activity based on reference to the W.H.O. First International Low Molecular Weight Heparin Reference Standard. Each enoxaparin sodium injection, USP prefilled syringe is for single, one-time use only and is affixed with a 27 gauge × 1/2 inch needle.

	Table 2	7 150 mg/mL Concentration		
Dosage Unit / Strength*	Anti-Xa Activity [†]	Unit of Sale	Syringe Label Color	Each
Graduated Prefilled Syringes [‡]				
120 mg/0.8 mL	12,000 IU	NDC# 63323-655-99 10 syringes per carton	Purple	NDC# 63323-655-21
150 mg/mL	15,000 IU	NDC# 63323-589-94 10 syringes per carton	Navy Blue	NDC# 63323-589-21
* Strength represents the number of million 15 mg enoxaparin sodium per 0.1 ml		ter for Injection. Enoxaparin sodium inje	ction, USP 120 and 150 mg g	raduated prefilled syringes contain

Approximate anti-Factor Xa activity based on reference to the W.H.O. First International Low Molecular Weight Heparin Reference Standard. Each enoxaparin sodium injection, USP graduated prefilled syringe is for single, one-time use only and is affixed with a 27 gauge × 1/2 inch needle Store at 20°C to 25°C (68°F to 77°F) [see USP Controlled Room Temperature]. Store in the original carton or packaging until ready to use

Keep out of reach of children 17 PATIENT COUNSELING INFORMATION

Advise the patient to read the FDA-approved patient labeling (Instructions for Use).

If patients have had neuraxial anesthesia or spinal puncture, and particularly, if they are taking concomitant NSAIDs, platelet inhibitors, or other anticoagulants, advise them to watch for signs and symptoms of spinal or epidural hematoma, such as tingling, numbness (especially in the lower limbs) and muscular weakness. Instruct the patient to seek immediate medical attention if any of these symptoms occur.

· of the instructions for injecting enoxaparin sodium injection if they continue enoxaparin sodium injection therapy after discharge from the hospital that it may take them longer than usual to stop bleeding. • that they may bruise and/or bleed more easily when they use enoxaparin sodium injection.

• that they should report any unusual bleeding, bruising, or signs of thrombocytopenia (such as a rash of dark red spots under the skin) to their physician [see Warnings and Precautions (5.1, 5.5)]. • to tell their physicians and dentists they are taking enoxaparin sodium injection and/or any other product known to affect bleeding before any surgery is scheduled and before any new drug is taken [see Warnings and Precautions (5.1, 5.3)]. • to tell their physicians and dentists of all medications they are taking, including those obtained without a prescription, such as aspirin or other NSAIDs [see Drug Interactions (7)].

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INSTRUCTIONS FOR USE **Enoxaparin Sodium Injection** for subcutaneous use

Single-dose prefilled syringe This Instructions for Use contains information on how to prepare and inject enoxaparin sodium injection prefilled syringe

Read this Instructions for Use before using the enoxaparin sodium injection prefilled syringe and each time you get a new prescription. There may be new information. Do not inject yourself or someone else until you have been shown how to inject enoxaparin sodium injection. Your healthcare provider can show you or your caregiver how to prepare and inject a dose of enoxaparin sodium injection. Call your healthcare provider if you have any questions.

· Your healthcare provider will tell you the prescribed dose that you should take and how often you will need to inject enoxaparin sodium injection. If your dose is based on your

body weight, your dose of enoxaparin sodium injection might be less than what is in the prefilled syringe.

Enoxaparin sodium injection is injected as a subcutaneous (under the skin) injection only. **Do not** inject enoxaparin sodium injection into muscle. · Do not use the enoxaparin sodium injection prefilled syringe if the needle cap is missing or not securely attached.

 Do not remove the needle cap until just before you give the injection Do not touch the syringe plunger rod until you are ready to inject. Do not pull back on the plunger rod at any time

To protect from needle-stick injuries, each prefilled syringe has a safety system that covers the needle after injection.

Do not get rid of any air bubble(s) in the enoxaparin sodium injection prefilled syringe. This can lead to a loss of the medicine.
To avoid bruising, do not rub the injection site after you have injected yourself.

 Throw away (dispose of) the used enoxaparin sodium injection prefilled syringe and needle cap right away after use, even if there is medicine left in the prefilled syringe. See "Step 13: Dispose of used enoxaparin sodium injection prefilled syringes and needle caps" below. · Enoxaparin sodium injection is for one time use only. Do not reuse an enoxaparin sodium injection prefilled syringe.

Storing enoxaparin sodium injection prefilled syringes:

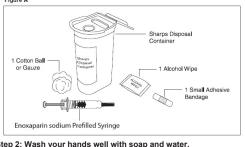
 Store enoxaparin sodium injection prefilled syringes at 77°F (25°C).
 Store enoxaparin sodium injection prefilled syringes in the original carton or packaging until ready to use. Keep enoxaparin sodium injection prefilled syringes and all medicines out of the reach of children.

Parts of enoxaparin sodium injection prefilled syringe

Preparing to inject enoxaparin sodium injection prefilled syringe: Step 1: Gather the following supplies for your injection (see Figure A):

enoxaparin sodium injection prefilled syringe 1 alcohol wipe'

1 cotton ball or gauze*
a small adhesive bandage, if needed a sharps disposal container* (see Step 13) *Items not included

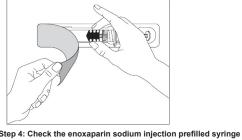


Step 2: Wash your hands well with soap and water Step 3: Preparing a dose of enoxaparin sodium injection

Take the prefilled syringe out of the package. Open the packaging by peeling the lid at the arrow as directed.

Take the prefilled syringe out of the plastic container by holding the middle of the syringe body (see Figure B).

Do not remove the prefilled syringe by pulling on the plunger rod or the needle cap as this may damage the syringe.
Do not pull off the needle cap until you are ready to inject.
Do not use the enoxaparin sodium injection prefilled syringe if it has been dropped on a hard surface or damaged.

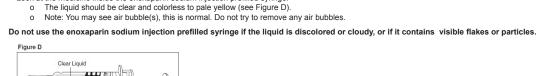


· Look at the medicine inside the enoxaparin sodium injection prefilled syringe

the expiration date on the prefilled syringe has not passed (see Figure C).

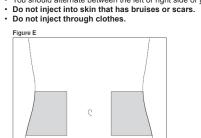
When you receive your enoxaparin sodium injection syringes, always check to see that

Do not use the enoxaparin sodium injection prefilled syringe if the expiration date has passed.



you have the correct medicine and dose.

Step 6: Choose your injection site You can inject into either the right or left side of your stomach area (abdomen), at least 2 inches away from your belly button and out towards your side (see Figure E). You should alternate between the left or right side of your stomach each time you give yourself an injection.

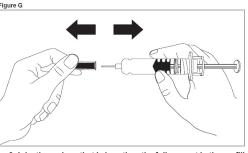


Step 7: Clean the injection site Clean the injection site with an alcohol wipe (see Figure F). Let your skin dry before injecting.



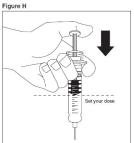
Step 8: Remove the needle cap
Hold the prefilled syringe in the middle of the body with the needle pointing away from you. Remove the needle cap by pulling it straight off the syringe (see Figure G).

 Do not twist the needle cap to avoid bending the needle. Do not put the needle cap back on. · Do not touch the needle

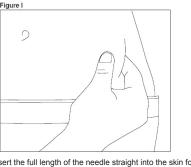


Step 9: Injecting a dose that is less than the full amount in the prefilled syringe. If your prescribed dose is the same as the amount in the prefilled syringe, go to Step 10. If your dose is based on your bodyweight, your healthcare provider may prescribe less than the full amount in the syringe. You will have to get rid of (discard) some of the medicine from the prefilled syringe before you inject enoxaparin sodium injection.

To measure your prescribed dose, hold the prefilled syringe with the needle pointing down. Carefully watch the numbers on the syringe as you push the plunger down until the amount left in the syringe is the same as your prescribed dose. The tip of the plunger should line up with the number for your prescribed dose (see Figure H)



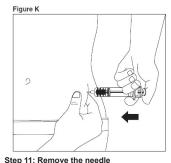
Step 10: Injecting enoxaparin sodium injection prefilled syringe
Hold the prefilled syringe like a pencil in your hand with the needle pointing down. With your other hand, pinch the cleaned stomach (abdomen) area between your forefinger and thumb to make a fold in the skin (see Figure I). Make sure you hold the skin fold during the entire injection.



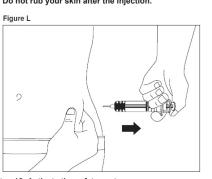
Insert the full length of the needle straight into the skin fold at about a 90° angle (see Figure J).



Push the plunger rod down slowly and steadily with your thumb until the enoxaparin sodium injection prefilled syringe is empty (see Figure K).



Step 11: Remove the needle Remove the needle from the injection site by pulling it straight out while keeping your fingers on the plunger rod (see Figure L). · Do not put the needle cap back on. Do not rub your skin after the injection.



Step 12: Activate the safety system Point the needle away from yourself and other people, and firmly push the plunger rod again to activate the safety system. The protective sleeve will automatically come down and You will feel some resistance. This is normal. Keep pushing until you hear the "click."

Activate the system while facing the syringe away from yourself and other people CLICK

Step 13: Dispose of used enoxaparin sodium injection prefilled syringes and needle caps
Put the used enoxaparin sodium injection prefilled syringe and needle cap in an FDA-cleared sharps disposal container right away after use (see Figure N). Do not dispose of enoxaparin sodium injection prefilled syringes or needle caps in your household trash.

o made of a heavy-duty plastic, o can be closed with a tight-fitting, puncture-resistant lid, without sharps being able to come out, o upright and stable during use,

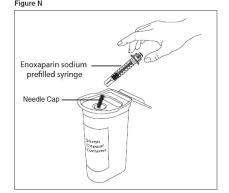
If you do not have an FDA-cleared sharps disposal container, you may use a household container that is:

The safety system can only be activated after the syringe has been emptied.
Only activate the safety system after you have removed the needle from your skin.

Activation of the safety system may cause a small amount of liquid to leak out of the syringe

o properly labeled to warn of hazardous waste inside the container.

When your sharps disposal container is almost full, you will need to follow your community guidelines for the right way to dispose of your sharps disposal container. There may be state or local laws about how you should throw away used needles, syringes, and prefilled syringes. For more information about safe sharps disposal, and for specific information about sharps disposal in the state that you live in, go to the FDA's website at http://www.fda.gov/safesharpsdisposal.



This Instructions for Use has been approved by the U.S. Food and Drug Administration Issued: January 2025