



**Inclusion criteria:**

- Therapeutic indication for LMWH management

**Exclusion criteria:**

- Renal failure
- Diagnosis of HIT

Patient with indication for low molecular weight heparin (LMWH) management

**Indications for management:**

- Treatment or prophylaxis of VTE
- Treatment or prophylaxis of arterial thrombosis

LMWH  
Considerations for use

**Adverse effects:**

- Most common event is bleeding; refer to [LMWH antidote](#) for management
- Osteoporosis
  - Rare but may occur with prolonged use of LMWH
  - Refer to [monitoring](#) section for additional recommendations
- Thrombocytopenia due to HIT
  - Risk in children is low overall
  - May be asymptomatic
  - Risk increases after 5 days of therapy
  - May be associated with life threatening arterial or venous thrombosis
  - Refer to [monitoring](#) section for additional recommendations

**Drug Interactions:**

- Increased potential for hemorrhage:
  - Anticoagulants: heparin, vitamin K antagonists, direct thrombin inhibitors
  - Thrombolytic agents: alteplase, streptokinase, urokinase
- Drugs affecting platelet function: aspirin, NSAIDs, dipyridamole, clopidogrel, ticlodipine, cilostazol
- Complementary/alternative medications known to have the potential to increase bleeding risk: garlic, ginger, ginkgo biloba, fenugreek, St. John's Wort

**Other considerations:**

- [Fast Facts](#)
- For assistance transitioning between anticoagulants, contact Hematology
- For analgesia, consider alternative agents (e.g. acetaminophen), as clinically appropriate
- Avoid IM injections and arterial punctures
- Encourage mobilization as tolerated

**Guidance for holding prior to procedures:**

- Hold LMWH doses for 24 hours prior to **immunizations or invasive procedures** such as LP or surgery, unless clinical situation requires emergent intervention.
- Restart 12 hours after the procedure, surgery or immunization.

**Initiation and Maintenance**

- [Indications for Hematology Consult](#)
- Obtain baseline CBC, PT, aPTT
- Refer to [Initiation and Maintenance](#) for dosing and titration guidelines
- [Patient Education](#)

**Monitoring**

- Monitor bone density if therapy exceeds 3 months; consider bone densitometry studies on day 1 and approximately every 12 months to assess for osteoporosis
- Monitor platelet counts weekly until stable on LMWH
  - If platelet count decreases below 150,000/μL or drops by ≥ 50%, determine if decrease in platelet count is related to underlying disorder or LMWH therapy
  - If likely due to LMWH, suspect HIT and discontinue use; initiate an alternative therapy and consult Hematology
- The **optimal sample** for LMWH levels is a fresh venipuncture site
  - Alternate sites may be considered but present limitations with interpretation of levels
  - Capillary samples are not appropriate
  - Ensure sample is not contaminated by heparin (e.g. from arterial line) by drawing adequate waste volume to clear line before obtaining sample

**Abbreviations:**

DVT = deep venous thrombosis  
 HIT = heparin induced thrombocytopenia  
 IM = intramuscular  
 PE = pulmonary embolism  
 VTE = venous thromboembolism

Duration of Therapy		
<b>Provoked thrombus</b>	DVT	Minimum 3 months of therapy suggested; consider conversion to <b>warfarin*</b> based on clinical situation.
	Extensive DVT or large PE	Administer LMWH for 7-14 days prior to considering conversion to <b>warfarin*</b> . Minimum 6 months of therapy with prolongation and/or transition to warfarin based on clinical situation.
<b>Unprovoked thrombus</b>		Administer LMWH for 7-14 days prior to considering conversion to <b>warfarin*</b> . Minimum 6 months of therapy with prolongation and/or transition to warfarin based on clinical situation.
*Warfarin (not typically used in infants) is generally added on day 1 for a DVT and day 5 for an extensive DVT or large PE		



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## References:

David, M., et al. (2007, Jan). Heparin and LMWH in Children. Thrombosis Interest Group of Canada. Retrieved Nov 15, 2018, from <http://www.tigc.org/eguidelines/heparinchild07.htm>.

Lexicomp Online, Pediatric and Neonatal Lexi-Drugs. *Enoxaparin*. Retrieved Nov 2018, from <https://online.lexi.com>.

Monagle, P., Chan, A. K. C., Goldenberg, N. A., Ichord, R. N., Journeycake, J. M., Nowak-Göttl, U., & Vesely, S. K. (2012). Antithrombotic therapy in neonates and children: Antithrombotic Therapy and Prevention of Thrombosis, 9th ed: American College of Chest Physicians Evidence-Based Clinical Practice Guidelines. *Chest*, *141*(2 Suppl), e737S-e801S. <https://doi.org/10.1378/chest.11-2308>

Monagle, P., Cuello, C. A., Augustine, C., Bonduel, M., Brandão, L. R., Capman, T., Chan, A. K. C., Hanson, S., Male, C., Meerpohl, J., Newall, F., O'Brien, S. H., Raffini, L., van Ommen, H., Wiernikowski, J., Williams, S., Bhatt, M., Riva, J. J., Roldan, Y., . . . Vesely, S. K. (2018). American Society of Hematology 2018 Guidelines for management of venous thromboembolism: treatment of pediatric venous thromboembolism. *Blood Adv*, *2*(22), 3292-3316. <https://doi.org/10.1182/bloodadvances.2018024786>

Roach, E. S., Golomb, M. R., Adams, R., Biller, J., Daniels, S., Deveber, G., Ferriero, D., Jones, B. V., Kirkham, F. J., Scott, R. M., & Smith, E. R. (2008). Management of stroke in infants and children: a scientific statement from a Special Writing Group of the American Heart Association Stroke Council and the Council on Cardiovascular Disease in the Young. *Stroke*, *39*(9), 2644-2691. <https://doi.org/10.1161/strokeaha.108.189696>