Making Cancer History®

MD Anderson Cancer Center History Making Cancer History Making Can

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MANAGEMENT INITIAL EVALUATION VTE RISK LEVEL • Initiate mechanical prophylaxis prior to anesthesia in the OR and continue postoperatively until patient is fully ambulatory if no contraindications exist⁴ • Encourage ambulation • Mitigate risk factors • Reassess for pharmacologic contraindications⁵ daily Yes **HIGH RISK** Does patient Expected altered have contraindications $mobility^2 > 48 hours$ • Initiate mechanical prophylaxis prior to anesthesia in the OR and continue to pharmacologic and > 2 VTE risk postoperatively until patient is fully ambulatory if no contraindications exist⁴ prophylaxis³? No • Discuss pharmacological prophylaxis with pediatric hematology^{5,6}. Notify factors anesthesia team as this may affect anesthetic choice. • Encourage ambulation • Mitigate risk factors Patients 10-17 years old¹ Assess for • If neuraxial catheter or spinal procedure planned or in place, see Appendix E who are expected to VTE risk have a surgical factors, see procedure lasting Appendix A > 60 minutes **MODERATE RISK** • Expected altered mobility² > 48 hours • Initiate mechanical prophylaxis prior to anesthesia in the OR and continue and either 0 or 1 VTE risk factor postoperatively until patient is fully ambulatory if no contraindications exist⁴ • Encourage ambulation • Mitigate risk factors LOW RISK • Expected altered mobility² < 48 hours

OR = operating room

regardless of VTE risk factors

¹ Patients < 10 years old do not need VTE prophylaxis perioperatively unless there is known inherited thrombophilia or previous history of deep vein thrombosis (DVT); consult Pediatric Hematology in such cases

² Altered mobility is defined as a permanent or temporary state in which the child has a limitation in independent, purposeful physical movement of the body or of one or more extremities

³ See Appendix B for contraindications to pharmacological options for VTE prophylaxis

⁴ See Appendix C for mechanical VTE prophylaxis

⁵ See Appendix D dosing for VTE pharmacologic prophylaxis in pediatric patients

⁶ Obtain pediatric hematology consult when weighing risk versus benefit in patients at risk of bleeding



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APPENDIX A: VTE Risk Factors

- Active cancer (or suspicion of cancer)
- Blood stream infection
- Central venous catheter (including non-tunneled, tunneled and peripherally inserted central catheters)
- Treatment factors including chemotherapy (especially asparaginase, bevacizumab, thalidomide/lenalidomide plus high-dose dexamethasone), protein kinase inhibitors, immunotherapy, and/or antiangiogenic agents
- Exogenous estrogen compounds (contraceptives, hormone replacement, tamoxifen/raloxifene, diethylstilbestrol) within past two months
- History of venous thrombosis
- Hyperosmolar state (serum osmolality > 320 mOsm/kg)
- History of inflammatory diseases (e.g., inflammatory bowel disease, systemic lupus erythematosus)
- Obesity (BMI > 95th percentile for age)
- Orthopedic procedures: hip or knee reconstruction
- History of nephrotic syndrome
- History of familial and/or acquired hypercoagulability
- Major trauma: more than 1 lower extremity long bone fracture, complex pelvic fractures, spinal cord injury
- Major surgery: abdominal, pelvic, orthopedic
- Erythropoietin stimulating agents in patients undergoing orthopedic surgery
- Altered mobility¹
- History of antiphospholipid antibodies
- History of polycythemia
- History of congenital heart disease (non-biologic reconstruction)

APPENDIX B: Contraindications to Pharmacological Options for VTE Prophylaxis

Absolute Contraindications

- Evidence of or high risk of bleeding: cerebral, gastrointestinal or genitourinary
- Uncorrected coagulopathy
- Bleeding disorder (known or suspected)
- Severe thrombocytopenia: platelets < 30 K/microliter
- Hypersensitivity to enoxaparin, heparin, pork products, or any component of the formulation
- Epidural or paraspinal hematoma

Relative Contraindications

- Moderate thrombocytopenia: platelets 30-50 K/microliter
- For patients undergoing spinal procedures and/or epidural placement/removal, see Appendix E
- Intracranial or spinal lesion at high risk of bleeding
- Recent major surgery at high risk of bleeding (e.g., neurosurgical)
- Pelvic fracture within past 48 hours
- Uncontrolled hypertension
- Renal failure

Altered mobility is defined as a permanent or temporary state in which the child has a limitation in independent, purposeful physical movement of the body or of one or more extremities



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APPENDIX C: Mechanical VTE Prophylaxis

Sequential compression devices (SCDs) should be used for pediatric patients requiring mechanical VTE prophylaxis. The goal is to use SCDs for 18 hours per day.

Contraindications:

- DVT, suspected or existing (can use graduated compression stockings)
- Extremity to be used has acute fracture
- Extremity to be used has peripheral IV access
- Skin conditions affecting extremity (e.g., dermatitis, burn)
- Unable to achieve correct fit due to patient's size

APPENDIX D: Dosing for VTE Pharmacologic Prophylaxis in Pediatric Patients

Enoxaparin:

Weight < 50 kg: 0.5 mg/kg subcutaneously twice daily

Weight > 50 kg: 40 mg subcutaneously once daily

Aspirin (may be used in orthopedic patients, not recommended in other populations): 81 mg PO

APPENDIX E: Spinal Procedure and/or Neuraxial Catheter Management

Hold times prior to Lumbar Puncture (LP) or neuraxial catheter removal or placement:

• Enoxaparin: 12 hours

Hold time <u>after LP</u> or neuraxial catheter <u>placement:</u>

• Enoxaparin: 8 hours (if blood in CSF sample: 24 hours)

Hold time after neuraxial catheter removal:

• Enoxaparin: 8 hours

CSF = cerebrospinal fluid



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SUGGESTED READINGS

- Cincinnati Children's Hospital Medical Center, Multidisciplinary VTE Prophylaxis BESt Team (2014). Best evidence statement venous thromboembolism (VTE) prophylaxis in children and adolescents (Hospital Medicine/Prophylaxis/Venous Thromboembolism/BESt 181). Retrieved from https://www.cincinnatichildrens.org/-/media/cincinnati%20childrens/home/service/j/ anderson-center/evidence-based-care/recommendations/type/venous%20thromboembolism%20best%20181
- Jinks, S., & Arana, A. (2019). Venous thromboembolism in paediatrics. British Journal of Anaesthesia, 19(9), 305. doi:10.1016/j.bjae.2019.05.003
- 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: American College of Chest Physicians evidence-based clinical practice guidelines. CHEST Journal, 141(2 Suppl), e737S–e801S. doi:10.1378/chest.11-2308
- Morgan, J., Checketts, M., Arana, A., Chalmers, E., Maclean, J., Powis, M., ... Association of Paediatric Anaesthetists of Great Britain and Ireland guidelines working group on thromboprophylaxis in children. (2018). Prevention of perioperative venous thromboembolism in pediatric patients: Guidelines from the Association of Paediatric Anaesthetists of Great Britain and Ireland (APAGBI). Pediatric Anesthesia, 28(5), 382-391. doi:10.1111/pan.13355
- Punzalan, R. C., Hillery, C. A., Montgomery, R. R., Scott, J. P., & Gill, J. C. (2000). Low-molecular-weight heparin in thrombotic disease in children and adolescents. *Journal of Pediatric* Hematology/Oncology, 22(2), 137-142. Retrieved from https://journals.lww.com/jpho-online/fulltext/2000/03000/Low Molecular Weight Heparin in Thrombotic Disease.11.aspx



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DEVELOPMENT CREDITS

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