

The Importance of Deep Vein Thrombosis Prophylaxis in Orthopaedic Patients

For more than 30 years multiple national and international evidence based guidelines and consensus papers have highlighted how patients undergoing major orthopaedic surgery are at very high risk of venous thromboembolism (VTE)¹⁻¹⁰. Without prophylaxis, patients undergoing hip and knee arthroplasty have been found to have a deep vein thrombosis (DVT) prevalence of up to 70%^{1,2,9-11}.

VTE is the most commonly seen post-operative complication following joint replacement in the lower extremity^{6,11}. A study conducted in 220 acute US hospitals, identified that following major orthopaedic surgery, those patients who developed VTE had significantly longer hospital stays and in-patient costs were twice as much as standard care¹².

Why does both hip and knee arthroplasty surgery place the patient at risk of VTE development?

Hip arthroplasty

To enable femoral preparation and insertion of the prosthesis, the lower leg is disarticulated. This results in obstruction and kinking of the femoral vein, producing an endothelial injury which then can potentiate thrombus formation and propagation^{4,13}. Heat activated cement may also be used, which has the potential to cause vascular damage¹³.

Knee arthroplasty

During knee replacement surgery, a tourniquet is used to reduce blood loss and also to maintain a clear surgical field. This causes damage to the endothelial lining of the vein, increasing risk of thrombus formation¹³. Without prophylaxis, the total DVT rate is greater in knee replacement surgery compared to hip replacements¹⁰.

Venous stasis

Both during and after surgery, patients who have undergone lower limb orthopaedic surgery have a marked decrease in pulsatile blood flow, due to loss of normal physiological muscle contraction in the lower limbs¹⁴. Patients are not able to perform active dorsal to plantar flexion and are also immobile for long periods of time during recovery.

Alteration in fibrinolytic activity

Vessel damage and bone trauma during the operative procedure leads to the enhanced release of circulating procoagulants. It has been established for many years that major surgery and trauma are accompanied by a recognised reduction in the spontaneous fibrinolytic activity of the blood, a process called 'fibrinolytic shutdown'²³⁻²⁷. This phenomenon is reported to commence during or soon after the surgical procedure and last for at least 3 days^{23,26}.

Mechanical and biochemical effects of FLOWTRON® DVT Prophylaxis Systems

Prevention of venous stasis

FLOWTRON Systems actively augment venous blood flow¹⁴⁻²¹. This reduces stasis, flushes valve pockets where many thrombi originate, decreases venous hypertension and decreases interstitial oedema²².

Increased fibrinolytic activity

Use of *FLOWTRON* Systems results in an increase in the fibrinolytic activity of the blood^{28,29}, a suppression of procoagulant factors³⁰ and may assist in the reversal or prevention of fibrinolytic shutdown.

Clinical studies using the FLOWTRON DVT Prophylaxis Systems

Clinical studies undertaken in elective orthopaedic surgical patients utilising *FLOWTRON* Systems have

established high levels of efficacy combined with excellent patient concordance and freedom from adverse effects^{11,13,14,31-36}.

An additional benefit found that when *FLOWTRON* Systems were used as the method of prophylaxis compared to low molecular weight heparin (LMWH), the operative field was easier to work in and drier^{31,32,36}.

Significant cost savings have also been reported in a randomised controlled trial of 442 patients comparing use of *FLOWTRON* DVT Prophylaxis Systems with Enoxaparin³⁷. Over a 23 month period, similar incidence rates of VTE were recorded in each cohort and a cost saving of US \$67,300 was made in the *FLOWTRON* Systems group. The paper highlights how the use of *FLOWTRON* Systems for the prevention of DVT is a safe and cost effective alternative to LMWH.

Duration of prophylaxis

Thromboprophylaxis using IPC should be commenced intra-operatively as DVT often develops as a result of the surgical procedure⁴. Evidence based guidelines and consensus papers^{3,8,10} highlight the requirement for prolonged prophylaxis of up to 4-6 weeks post-operatively. Use of IPC can be continued for as long as tolerated by the patient and if necessary switched to chemical prophylaxis for the duration of the post-operative at risk period³.

References

1. National Institutes of Health (1986). Prevention of venous thrombosis and pulmonary embolism. *National Institutes of Health Consensus Development Conference Statement*; 6: 2.
2. European Consensus Statement (1992). Prevention of venous thromboembolism. *International Angiology*; 11(3): 151-159.
3. Nicolaidis AN, Fareed J, Kakkar AK et al (2006). Prevention and treatment of venous thromboembolism - International Consensus Statement. *International Angiology*; 25(2): 101-147.
4. Salvati E, Pellegrini V, Sharrock N et al (1998). Recent advances in venous thromboembolic prophylaxis during and after total hip replacement – presented at the Annual Meeting of the American Orthopaedic Association, Asheville, North Carolina. *The Journal of Bone and Joint Surgery*; 82-A(2): 252-270.
5. Second Thromboembolic Risk Factors (THRIFT II) Consensus Group (1998). Risk of and prophylaxis for venous thromboembolism in hospital patients. *Phlebology*; 13: 87-97.
6. Freedman K, Brookenthal K, Fitzgerald R et al (2000). A meta-analysis of thromboembolic prophylaxis following elective total hip arthroplasty. *The Journal of Bone and Joint Surgery*; 82-A(7): 929-939.
7. Silleran-Chassany J, Safran D (2000). Prophylaxis of perioperative venous thrombosis: Role of venous compression. *Phlebology*; 15: 138-142.



FLOWTRON Excel DVT Prophylaxis System

8. The Australia and New Zealand Working Party on the



FLOWTRON Universal DVT Prophylaxis System

management and prevention of venous thromboembolism (2007). Best practice guidelines for Australia and New Zealand 4th Edition. *HEMI Publishing*.

9. Scottish Intercollegiate Guidelines Network (SIGN) (2002). Prophylaxis of venous thromboembolism: A national clinical guideline recommended for use in Scotland; SIGN: Edinburgh.
10. Geerts WH, Bergqvist D, Pineo GF et al (2008). Prevention of venous thromboembolism: American College of Chest Physicians Evidence-Based Clinical Practice Guidelines (8th Edition). *Chest*; 133(6Suppl): 381S-453S.
11. Pidala J, Duane L, Donovan M et al (1992). A prospective study on intermittent pneumatic compression in the prevention of deep vein thrombosis in patients undergoing total hip or total knee replacement. *Surgery*; 175: 47-51.
12. Ollendorf DA, Vera-Llonch V, Oster G (2002). Cost of venous thromboembolism following major orthopaedic surgery in hospitalised patients. *American Journal of Health Syst Pharm*; 59(18): 1750-1754.

13. Capper C (1998). External pneumatic compression therapy for DVT prophylaxis. *British Journal of Nursing*; 7(14): 851-854.
14. Westrich G, Specht LM, Sharrock NE et al (1998). Venous hemodynamics after total knee arthroplasty: Evaluation of active dorsal to plantar flexion and several mechanical compression devices. *The Journal of Bone & Joint Surgery*; 80-B(6): 1057-1066.
15. Flam E, Berry M, Coyle A et al (1993). DVT Prophylaxis: Comparison of two thigh-high intermittent pneumatic compression systems. Presented at the meeting of the American College of Surgeons, San Francisco October.
16. Flam E, Berry S, Coyle V et al (1996). Blood-flow augmentation of intermittent pneumatic compression systems used for the prevention of deep vein thrombosis prior to surgery. *The American Journal of Surgery*; 171(3): 312-315.
17. Flam E, Nackman G, Tarantino D and Raab, L (2000). Intermittent pneumatic compression devices of the foot: A comparison of various systems on femoral vein blood flow velocity augmentation in the supine and dependent, non-weight bearing positions. *ArjoHuntleigh Clinical Report*.
18. Procter MC, Zajkowski PJ, Wakefield TW et al (2001). Venous hemodynamic effects of pneumatic compression devices. *The Journal of Vascular Technology*; 25(3): 141-145.
19. Woodcock J and Morris R (2002). The effect of the Kendall SCD® and Huntleigh Flowtron DVT30 garments on femoral and popliteal vein blood flow measurements. *ArjoHuntleigh Clinical Report*.
20. Morris RJ, Giddings JC, Ralis HM et al (2003). The hematologic and hemodynamic effects of the Aircast Venaflo calf-length and the Huntleigh Flowtron Calf-length intermittent pneumatic compression for deep vein thrombosis prophylaxis. *ArjoHuntleigh Clinical Report*.
21. Morris RJ, Giddings JC, Jennings GM et al (2003). The hematological and hemodynamic comparison of the Kendall AV Impulse™ and the Huntleigh FP5000 Intermittent Pneumatic Foot Compression System. *ArjoHuntleigh Clinical Report*.
22. Kumar S and Walker M (2002). The effects of intermittent pneumatic compression on the arterial and venous system of the lower limb: a review. *Journal of Tissue Viability*; 12(2): 58-65.
23. Griffiths NJ (1979). Factors affecting the fibrinolytic response to surgery. *Annals of the Royal College of Surgeons of England*; 61(1): 12-16.
24. D'Angelo A, Kluff C, Verheijen JH et al (1985). Fibrinolytic shut-down after surgery: impairment of the balance between tissue type plasminogen activator and its specific inhibitor. *European Journal of Clinical Investigation*; 15: 308-312.
25. Kassis J, Hirsh J, Podor TJ (1992). Evidence that postoperative fibrinolytic shutdown is mediated plasma factors that stimulate endothelial cell type 1 plasminogen activator inhibitor biosynthesis. *Blood*; 80(7): 1758-1764.
26. Dahl OE, Pedersen T, Kierulf P et al (1993). Sequential intrapulmonary and systemic activation of coagulation and fibrinolysis during and after total hip replacement surgery. *Thrombosis Research*; 70: 451-458.
27. Cahan MA, Hanna DJ, Wiley LA et al (2000). External pneumatic compression and fibrinolysis in abdominal surgery. *Journal of Vascular Surgery*; 32(3): 537-543.
28. Giddings JC, Ralis H, Davis D et al (2004). Systematic haemostasis after intermittent pneumatic compression. Clues for the investigation of DVT prophylaxis and traveller thrombosis. *Clin. Lab. Haematol*; 26(4): 269-273.
29. Morris RJ, Giddings JC, Ralis HM et al (2006). The influence of inflation rate on the haematologic and haemodynamic effects of intermittent pneumatic compression for deep vein thrombosis prophylaxis. *Journal of Vascular Surgery*; 44(5): 1039-1045.
30. Giddings JC, Ralis H, Davies D et al (2001). Suppression of the tissue factor pathway combined with enhanced tissue plasminogen activator activity (tPA) and urokinase plasminogen activator activity (scuPA) after intermittent pneumatic compression. *Thrombosis and Haemostasis*; 86: S2240.
31. Stone M, Limb D, Campbell P et al (1996). A comparison of intermittent calf compression and enoxaparin for thromboprophylaxis in total hip replacement. *International Orthopaedics*; 20: 367-369.
32. Richards S, Espahbodi S, McCarthy I et al (2001). Intermittent pneumatic foot compression for prophylaxis against thromboembolic disease in total hip replacement. *ArjoHuntleigh Clinical Report*.
33. Cameron J (2001). Avoiding haemorrhage and wound infection in total arthroplasty whilst maintaining deep venous thrombosis prophylaxis. Presented at RCN Orthopaedic Nursing Conference, Belfast, September.
34. Pagella P, Cipolle M, Sacco E et al (2007). A randomised trial to evaluate compliance in terms of patient comfort and satisfaction of two pneumatic compression devices. *Orthopedic Nursing*; 26(3): 169-174
35. Brooks PJ, Keramati M, Wickline A (2007). Thromboembolism in patients undergoing total knee arthroplasty with epidural analgesia. *The Journal of Arthroplasty*; 22(5): 641-3
36. Eskander M, Limb D, Stone M et al (1997). Sequential mechanical and pharmacological thromboprophylaxis in the surgery of hip fractures. *International Orthopaedics*; 21: 259-261.
37. Ginzburg E, Cohn S, Lopez J et al (2003). Randomised clinical trial of intermittent pneumatic compression and low molecular weight heparin in trauma. *British Journal of Surgery*; 90: 1338-1344.
38. Sullivan SD, Kahn SR, Davidson BL et al (2003). Measuring the outcomes and pharmacoeconomic consequences of venous thromboembolism prophylaxis in major orthopaedic surgery. *Pharmacoeconomics*; 21(7): 477-496.