## Deep vein thrombosis at King Abdul Aziz University Hospital

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## **ABSTRACT**

**Objective:** To study the incidence of deep vein thrombosis and pulmonary embolism at King Abdul Aziz University Hospital. To determine the risk factors, use of different diagnostic modalities, treatment given and to compare our findings with those reported in the literature.

**Methods:** Retrospective study which included all cases of deep vein thrombosis and pulmonary embolism admitted to the medical ward of King Abdul Aziz University Hospital during the period between January 1994 till March 1999 were analyzed.

**Results:** Total of 75 patients were diagnosed to have deep vein thrombosis with mean age of 44.16 +/- 14.5 years and male:female ratio of 1:2. Doppler ultrasound was used for the diagnosis in 56 of 75 patients (75%). Pulmonary embolism as a complication of deep vein

thrombosis developed in 24 of 75 patients (32%). Prolonged immobilization was found to be the most common risk factor 17 of 75 (23%). All the patients were treated with conventional heparin followed by warfarin.

**Conclusion:** As discussed, our results are comparable with those reported in the literature. Post operative patients who are anticipated to have prolonged immobilization should receive prophylactic anti coagulation with subcutaneous heparin. Thrombophillia screeing should be reserved for those with recurrent deep vein thrombosis or patients with positive family history.

**Keywords:** Deep vein thrombosis, pulmonary embolism, risk factors.

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Venous thrombo embolism is a serious disorder and frequently heralds fatal outcome. The various modalities of treatment of deep vein thrombosis (DVT) and pulmonary embolism (PE) have changed considerably over the last 10 - 20 years and will continue to do so over the next coming years. The non specific nature of the clinical presentations of both DVT and PE may lead to substantial delay in diagnosis and initiation of therapy, in turn accounting for significant morbidity and mortality. Objective testing for DVT is crucial as clinical assessment alone is unreliable<sup>2,3</sup> because undiagnosed DVT can cause fatal PE<sup>4,5</sup> and because treatment of DVT is effective. More than half of the cases are never diagnosed and all too frequently PE is first determined at autopsy.<sup>6,8</sup> Patients with acute

lower extremity DVT often do not exhibit erythema, warmth, swelling, pain or tenderness and the sensitivity and specificity of the clinical examination are too low to be relied upon.<sup>9</sup> The objective of our study is to cause awareness of the risk factors for DVT in our hospital to enhance clinical suspicion and encourage prophylaxis to high risk group.

**Methods.** The study included all cases of DVT that were admitted to the medical ward of King Abdul Aziz University Hospital (KAUH), Jeddah, Saudi Arabia during the period between January 1994 till March 1999. The medical records of all patients were reviewed. Patients age, sex, and nationality were recorded as well as the site of DVT,

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whether right, left or bilateral and its extent, whether above knee, below knee or extensive (above and below knee together). Patients admitted to our hospital with suspected clinical diagnosis of DVT first had doppler ultrasound as the initial investigation to prove the diagnosis. If the results were inconclusive and there was a strong clinical suspicion of DVT, they underwent contrast venography. Patients with suspected PE had ventilation perfusion scan and if this was negative and there was a strong clinical suspicion, they would undergo pulmonary angiography. History of previous DVT was reported. Different risk factors such as prolonged immobilization, pregnancy, post operative DVT and the type of surgery, use of contraceptive pills, heart failure, bone fracture, underlying malignancy were all recorded. Measurement of protein C, S and anti thrombin III were carried out in patients with recurrent DVT, those with family history of thrombosis and those with unusual thrombosis site. Type of treatment given to the patient was recorded whether heparin followed by warfarin or thrombolytic therapy. Statistical analysis was carried out using the statistical package for social sciences (SPSS 7.5). T test and chi-square were used appropriately. Results were considered significant if the P value is less than 0.05.

**Results.** A total of 75 patients were analyzed with mean age of  $45.16 \pm 14.5$  years and male:female ration of 1:2. Most of the admissions were non-Saudi (67%). Left sided DVT was the most frequent type of presentation 44 of 75 (59%) left sided while 26 of 75 (35%) right sided and 5 of 75 (7%) had bilateral DVT. Extensive DVT was noticed in 57 of 75 (77%) while below knee was reported in 11 of 75 (25%) and 6 of 75 (8%) above knee. Doppler ultra sound was carried out for 56 of 75 (75%) of the patients to confirm the diagnosis while venography was required in 28 of 75 (37%) of the patients. Pulmonary embolism developed in 24 of 75 (32%) who were diagnosed by ventilation perfusion scan, none of the patients required pulmonary angiography. History of recurrent DVT was documented in 19 of 75 (26%) of patients, 7 of them (37%) had antiphospholipid syndrome, one (5%) had Bechet's disease, while in the remaining 11 (58%) no cause could be found. Seventeen of 19 patients who had recurrent DVT had extensive DVT involving both above and below knee (90%), whereas 3 of the nineteen patients (16%) had more than 2 recurrences. Risk factors as shown in Table 1 showed that prolonged immobilization was the most common risk factors 17 of 75 (23%). Twelve of 17 (70%) of patients who had history of prolonged immobilization developed extensive DVT, versus 5 of 17 (29%) who developed below knee DVT (p0.07). Three of 5 (60%) pregnant ladies and 3 of 5

Table 1 - Frequency of different risk factors for deep vein thrombosis.

| Risk Factors        | Number of patients (%) |
|---------------------|------------------------|
| Immobilization      | 17 (23%)               |
| Post operative      | 12 (16%)               |
| Post delivery       | 7 (9%)                 |
| Travel              | 6 (9%)                 |
| Contraceptive pills | 5 (7%)                 |
| Pregnancy           | 5 (7%)                 |
| Heart Failure       | 5 (7%)                 |
| Fracture            | 3 (4%)                 |
| Malignancy          | 3 (4%)                 |
| Drug addiction      | 3 (4%)                 |

(60%) of those who were using contraceptive pills developed recurrent DVT versus 2 of 5 (40%) and 2 of 5 (40%) who did not (both p0.06). Investigations, such as measurement of protein C and S and antithrombin III were carried out for 6 of 19 (84%) patients with recurrent DVT, none of them had any abnormality. All patients were treated with heparin followed by warfarin, only one patient received strepto kinase for extensive pulmonary embolism.

Discussion. Deep vein thrombosis is more common in the 4th decade of life. Risk is increased by advancing age and patients over 40 may qualify for prophylaxis. 10 Despite the different diagnostic tools available for the diagnosis of DVT, contrast venography (CV) remains the gold standard<sup>11</sup> but it is not an ideal one due to its invasive nature, risks associated with contrast media and its technical Of the non invasive methods for demand. diagnosing DVT, impedence plethysmography, venous ultra sonography have been extensively evaluated and widely used. In our study doppler ultrasound was used more than contrast venography. None of the patients had impedence plethysmography as it is not available in our institute. In general impedence plethysmography is less accurate than venous ultrasonography. 12-15 Left sided DVT is the most common type of presentation as reported by Lindblad et al.6 Extensive DVT was found to be more common in immobilized patients, this has been reported by others, thus emphasizing the use of prophylactic subcutaneous heparin which decreases the risk of DVT.1,18-23

Pregnancy and hormone replacement therapy doubles the risk of venous thrombo embolism.<sup>24</sup> The risk is higher near the start of therapy than after long term use.<sup>25</sup> In our study post delivery, contraceptive

pills and pregnancy were risk factors in that order of frequency. Malignancy as a cause of DVT was seen in only 4%, although it has been reported that previously unsuspected cancer is identified in patients with newly diagnosed DVT.26,27 predisposes patients to DVT and PE as late as one month post operatively.28 Thrombophillia screening like measurement of protein C, S and anti thrombin recommended for recurrent DVT.<sup>29</sup> Thrombophillia screening was negative in 84% of our patients who had recurrent DVT. Low molecular weight heparin was not used in any of our patients as it is not available in our hospital, although a number of prospective trials have demonstrated the efficacy of low molecular weight heparin.<sup>30,31</sup> D-Dimer test was not used in any of our patients as a diagnostic tool. The reason being the substantial variation in the assay performed for D-Dimer. At present, specific recommendations regarding the appropriate use of this test cannot be made.32,33

## References

- 1. Anderson FA, Wheeler HB, Goldberg RJ, Hosmer DW, Forcier A. The prevalance of risk factors for venous thrombo embolism among hospital patients. Arch Internal Med 1992; 152: 1660-1664.
- 2. Sandler DA, Martin JF, Duncan JS, Blake GM, Ward P, Ramsay LE et al. Diagnosis of DVT Comparison of Clinical Evaluation, Ultrasound, Impedance plethysnography with xray venography. Lancet 1984; 29: 716-719.
- Wells PS, Hirsh J, Anderson DR, Lensing AW, Foster G, Keoran C, et al. Accuracy of clinical assessment of DVT Lancet 1995; 345: 1326-1330.
- 4. Bell WR, Simon TL. Current status of pulmonary embolic Pathophysiology, diagnosis, prevention and disease. treatment. Am Heart J 1982; 103: 239-262.
- Anderson FA, Wheeler HB, Goldberg RJ, Hosmer DW, Palwardhan NA, Jovanovic B. A population based perspective of the hospital incidence and case fatality rates of DVT and PE. Arch Intern Med 1991; 151: 933-938.
- 6. Lindblad B, Erikson A, Bergqvist D. Autopsy verified PE in surgical department analysis of the period from 1995 - 1998. Br J Surg 1991; 78: 849-852.
- 7. Goldhaber SZ, Hennekens CH, Evans DA, Newton EC, Godleski JJ. Factors associated with correct antemortem diagnosis of major PE. Am J Med 1982; 73: 822-826.
- 8. Bergqvist D, Lindblad B. A 30 year survey of PE verified at autopsy. An analysis of 1274 surgical patients. Br J Surgical 1985; 72: 105-108.
- 9. Lectere Jr, Illescas F, Jarzen P. Diagnosis of DVT. In: Lectere Jr, editor. Venous thrombo embolic disorders. Philladelphia (USA): Lea and Febiger; 1991. p. 176-228.
- 10. Hirsh J. Modern management of pulmonary embolism. Proceedings of the Royal College of Physicians of Edinburgh 1994; 24: 548-553
- 11. Redman HC. DVT, is Contrast Venography still the 'Gold
- Standard'. Radiology 1988; 168: 277-278. Kearon C, Julian JA, Newman TE, Ginsberg JS. Non invasive diagnosis of DVT. Ann Intern Med 1998; 128: 663-
- 13. Ginsberg JS. Management of venous thrombo embolism. N Eng J Med 1996; 335: 1816-1828.

- 14. Comerota AJ, Katz ML, Greenwald LL, Leefmans E, Czeredarczuk M, White JV et al. Venous duplex imaging should it replace haemodynamic tests for DVT. J Vasc Surg 1990; 11: 53-59
- 15. Heijboer H, Buller HR, Lensing AW, Jongbloets LM. A comparison of real time compression. Ultrasonography for diagnostic management of patients with recurrent venous thrombosis. Acta Radiol 1992; 33: 297-300.
- 16. Bucur IJ. Venous thrombo embolism in high altitude. Saudi Medical Journal 1999; 20: 56-62.
- 17. Toglia MR, Weg JG. Venous thromboembolism during pregnancy. N Eng J Med 1996; 335: 108-114.
- Monreal M, Lofez E, Navarro A, Granew X, Cajo V, Salvador R. A prospective double blind trial of low molecular weight heparin with conventional low dose heparin 3 time daily to prevent pulmonary embolism and DVT in patients with hip fracture. J Trauma 1989; 6: 873-
- 19. Leyvraz PF, Van-melle G, Tregvard JM, Livio JJ, Candardjis G, Richard J et al. Adjusted versus fixed close subcutaneous heparin in the prevention of DVT after total hip replacement. N Eng J Med 1983; 309: 954-958.
- 20. Weitz JI. Low molecular weight heparin. N Eng J Med 1997; 337: 688-698.
- 21. Anderson DR, O'Brian BJ, Levine MN, Roberts R, Wells PS, Hirsh J. Efficacy and cost of low molecular weight heparin for prevention of DVT after total hip arthroplasty. Ann Intern Med 1993; 119: 1105-1112.
- 22. Hirish J , Joak J. Venous thrombo embolism and pulmonary embolish - a statement of Health Care Professionals. Circulation 1996; 93: 2212-2245.
- 23. Collins R, Scrimgeour A, Yousuf S, Peto R. Reduction in fatal PE and venous thrombosis by pre operative administration of subcutanous heparin. Overview of the result of randomized trials in general, orthopaedic and urologic surgery. N Eng J Med 1988; 318: 1162-1173.
- 24. McColl MD, Ramsay JE, Tait RC, Walker ID, Conkie JA, Carty MJ. Risk factors for pregnancy associated venous thrombo embolism. Thromb Haemost 1997; 78: 1183-1188.
- 25. Vandenbroucke JP, Helmerhorst FM. Risk of venous thrombo embolism with hormone replacement therapy. Lancet 1996; 348: 972.
- 26. Nodstrom M, Lindblad B, Anderson H, Bergqvist D, Kjellstrom T. DVT and occult malignancy, an Kjellstrom T. DVT and occult malignancy, an epidemological study. BMJ 1994; 308: 891-894.
- 27. Prandoni P, Lensing AW, Buller HR, Cogo A, Prins MH, Cattelan AM, et al. DVT and the incident of subsequent cancer. N Eng J Med 1992; 327: 1128-1133.
- 28. Lindblad B, Eriksson A, Bergqvist D. Autopsy verified pulmonary embolism in a surgical department. Analysis of the period from 1951-1988. Br J Surg 1991; 78: 849-852.
- 29. Heijboer H, Brandjes DP, Buller HR, Sturk A, ten Cate JW. Deficiencies of coaggulation inhibiting and fibrinolytic proteins in out patients with DVT. N Eng J Med 1990; 323: 1512-1516.
- 30. Koopman MM, Prandoni P, Piovell F, Ockelford PA, Brandjes DP, Gallus AS. Low molecular weight heparin and proximal DVT. N Eng J Med 1996; 334: 682-687.
- 31. Sirgusa S, Cosmi B, Piovell F, Hirsh J, Ginsberg JS. Low molecular weight heparin and unfractionated heparin in the treatment of patients with acute venous thrombo embolism. Results of meta analysis. Am J Med 1996; 100: 269-277.
- 32. Philbrick JT, Bachhuber TL, Humphries JE, Becker DM. D-Dimer testing and acute venous thrombo embolism - a short cut to accurate diagnosis. Arch Inter med 1996; 156: 939-946.
- Diagnosing PE, D-Dimer needs vigorous 33. Moser KM. evaluation. BMJ 1994; 309: 1525-1526.