Assessment of additional risk factors for deep vein thrombosis in patients with various malignancies

¹Ştefan C. Vesa, ²Antonia E. Macarie, ³Sonia I. Vlaicu, ⁴Bogdan Micu, ¹Anca D. Buzoianu, ³Sorin Crişan

¹ Departement of Pharmacology, Toxicology and Clinical Pharmacology, "Iuliu Hatieganu" University of Medicine and Pharmacy, Cluj-Napoca, Romania; ² Department of Geriatrics and Gerontology, "Iuliu Hatieganu" University of Medicine and Pharmacy, Cluj-Napoca, Romania; ³ Department of Internal Medicine, "Iuliu Hatieganu" University of Medicine and Pharmacy, Cluj-Napoca, Romania; ⁴ Vth Department of Surgery, "Iuliu Hatieganu" University of Medicine and Pharmacy, Cluj-Napoca, Romania.

Abstract. Abstract: Aim: to analyze the additional risk factors of deep vein thrombosis (DVT) in patients with a positive oncological history. Material and methods: The included 62 patients (29 (46.8%) women and 33 (53.2%) men) diagnosed with various cancers. The median age was 73.5 (68.75; 81) years. Patients were divided in two groups, the first group comprised 40 patients with DVT and the second group comprised 22 patients without DVT. The following risk factors for DVT were recorded for each patient: age, gender, varicose veins, history of bed rest longer than 3 days, bilateral lower limb edema, infections, chemotherapy, history of DVT, and other comorbidities such as: heart failure and diabetes mellitus. Results: History of DVT and the bed rest for more than 3 days were associated in univariate analysis with an increased risk of DVT in patients with cancer, and the prior enhanced the risk of DVT in multivariate analysis. Conclusion: History of DVT was an additional risk factor for DVT in patients with cancer.

Key Words: deep vein thrombosis, cancer, risk factors.

Copyright: This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Corresponding Author: A. E. Macarie, email: macarieantonia@yahoo.com

Introduction

Venous thromboembolism (VTE) is an entity that encompasses two diseases: deep vein thrombosis (DVT), usually involving the lower limbs, and pulmonary embolism (PE). VTE is an important health and economic issue, affecting over 10 million people every ear (Jha et al 2013). The mortality related to VTE events is high with over 300000 annual deaths in USA, and over 500000 in UE (Cohen et al 2008; Tapson 2008).

VTE due to deep vein thrombosis (DVT) is a common complication in patients with malignancies, for whom the risk of such an event is four- to six-fold higher than for age and sex matched controls (Bloom et al 2005). Since VTE is the second leading cause of death worldwide in oncological patients, the risk factors for DVT have been the object of many research groups looking to reduce the morbidity and mortality rate in these patients (Heit et al 2000). VTE is symptomatic in only 4-20% of cancer patients, but it is discovered in over 50% of autopsies (Khorana et al 2007).

There are several additional risk factors for DVT described in literature in oncologic patients. Some factors are related to the patient: advanced age, race (higher risk in Afro-Americans and lower in Asians) and comorbidities such as obesity, chronic obstructive pulmonary disease, previous DVT, thrombophilia. Other factors are linked to the cancer itself: primary location of the tumor, the first 3-5 months from diagnosis, presence of

metastases, chemotherapy, surgery related to the cancer, etc. (Lee&Levine 2003; Rosendaal 2005; Lyman et al 2015). We aimed to analyze the additional risk factors of DVT in patients with a positive oncological history.

Materials and methods

This retrospective, analytical, observational, transversal, cohort study included 62 patients (29 (46.8%) women and 33 (53.2%) men) diagnosed with various tumor types (lung, digestive system, genitourinary tract and hematological malignancies). The median age was 73.5 (68.75; 81) years. The patients were recruited from those admitted to the internal medicine ward of the Municipal Clinical Hospital of Cluj-Napoca between September 2007 and September 2009. The patients were divided in two groups, the first group comprised 40 patients with DVT and the second group comprised 22 patients without DVT. The diagnosis of acute DVT was established using duplex ultrasonography by means of an Aloka SSD-4000 device with a linear transducer (7-10 MHz), according to the criteria in force. Patients having signs and symptoms that were evocative for acute DVT underwent ultrasound examination.

All patients completed a signed consent prior to the enrolment in the study. The study protocol was approved by the Ethics Committee of "Iuliu Hatieganu" University of Medicine and Pharmacy, Cluj-Napoca.

Patients receiving oral or intravenous anticoagulant therapy at the time of enrolment or did not signed the informed consent were not included in the study.

The following data were recorded for each patient: age, gender, BMI, varicose veins, history of bed rest longer than 3 days, bilateral lower limb edema, infections, chemotherapy, history of DVT, and other comorbidities such as: heart failure, diabetes mellitus and COPD.

Statistical analysis was performed using MedCalc Statistical Software version 15.5 (MedCalc Software bvba, Ostend, Belgium; https://www.medcalc.org; 2015). Data were classified as qualitative or quantitative. Nominal variables were described using frequency and percentage. The Kolmogorov-Smirnov test was used to determine if the continuous variables were normal distributed. Quantitative variables were described using the median and the 25th and 75th percentile. The chi-square test or Fisher's exact test were used to analyse the difference in frequency between two groups of nominal variables. Multivariate analysis was performed using binary logistic regression. The level of statistical significance was set at p<0.05.

Results

Data recorded for the patients can be seen in table 1.

Table 1. Characteristics of patients with and without DVT

Table 1. Characteristics of patients with and without DV1					
Variables		Patients with DVT	Patients without DVT	P	
Age (years)		74 (69; 82)	73 (65.76; 77.25)	0.1	
Gender	Female	21 (52.5%)	8 (36.4%)	0.3	
Gender	Male	19 (47.5%)	14 (63.6%)	0.3	
Environment	Rural	19 (47.5%)	13 (59.1%)	0.5	
	Urban	21 (52.5%)	9 (40.9%)	0.5	
Heart failure		11 (27.5%)	6 (27.3%)	1	
Diabetes mellitus		4 (10%)	2 (9.1%)	1	
COPD		7 (17.5%)	2 (9.1%)	0.4	
Obesity		12 (30%)	3 (13.6%)	0.2	
History of DVT		14 (35%)	1 (4.5%)	0.01	
Bed rest for more than 3 days		8 (20%)	-	0.04	
Major surgery		1 (2.5%)	-	1	
Bilateral edema		10 (25%)	1 94.55)	0.07	
Varicose veins		13 (32.5%)	2 (9.1%)	0.08	
Chemotherapy		4 (10%)	-	0.2	

The different types of cancer are described in table 2. The location of the cancer was not related to the risk of DVT (p=0.5). A multiple logistic binary regression was used to determine the independent association between variables and DVT. The diagnosis of DVT was the dependent variable, while the following were introduced as independent variables: history of DVT, varicose veins and lower limb edema. The variable bed rest longer than 3 days was not included due to statistical concerns. These variables statistically significantly predicted the occurrence of

DVT, R2=0.19. As seen in table 3, the factor with the most important contribution to the occurence of DVT was the history of DVT (OR, 7.91; CI 0.91-68.73; p=0.029).

Table 2. Types of cancer

Type of cancer	Patients without DVT	Patients with DVT	Total
Brest/uterus	6	8	14
Drest/uterus	27.3%	20.0%	22.6%
D:	7	12	19
Digestive system	31.8%	30.0%	30.6%
O4h 4	3	2	5
Other types	13.6%	5.0%	8.1%
Pulmonary	2	7	9
rumonary	9.1%	17.5%	14.5%
Genitourinary	4	11	15
Genitourmary	18.2%	27.5%	24.2%

Table 3. The multiple logistic binary regression for DVT

Variables	В	P	OR	95% CI for OR	
		1	OK	Lower	Upper
History of DVT	2.069	0.05	7.915	0.912	68.735
Bilateral edema	1.303	0.258	3.68	0.384	35.24
Varicose veins	1.252	0.145	3.498	0.649	18.854

Discussion

Cancer-associated thrombosis is a common problem in cancer patients. Patients with cancer which are not hospitalized do not require routine anticoagulation. Only patients with additional comorbidities or conditions that place them in the high risk percentile for DVT should receive thromboprophylaxis (Lyman et al 2015). Thus it is extremely important to identify which of the risk factors are associated with an increase chance of DVT onset. The occurrence of DVT in patients with cancer is due to concomitant action of several factors: inflammation caused by necrosis or release of acute phase proteins, stasis caused by external tumoral compression or vascular invasion. The tumors have the capacity of secreting specific substances (tissue factor, cytokines) which can inhibit the release of anticoagulation factors or can upregulate the procoagulant factors, and thus resulting a hypercoagulable state (Falanga et al; 1993; Falanga&Rickles 1999; Blow et al 2004).

In our study, the history of DVT was the strongest risk factor for onset of DVT (OR, 7.9; p=0.05). The risk of another DVT is lower in patients with transient risk factors like temporary immobility, estrogen therapy, and higher in those with permanent conditions (cancer, obesity, paralysis; thrombophillia) (Heit et al 2000a). Despite anticoagulation after the thrombotic episode, there is still a 5 to 7% chance of thrombosis recurrence within a year for cancer patients (Ginsberg 1996).

The bed rest for more than 3 days was the other factor associated with the onset of DVT. Due to the fact that there were no controls with bed rest, this parameter could not be included in the multivariate analysis, but this does not mean that it's not a strong risk factor. The long period of bed rest increase the chance of DVT, due to venous stasis (Weill-Engerer et al 2004). Limitations of the study include the small number of cases and controls, the absence of data regarding thrombophilia and the limited data regarding metastases.

Conclusion

The history of DVT and bed rest for more than 3 days increase the risk of an acute DVT in cancer patients.

References

- Blom JW, Osanto S, Rosendaal FR. The risk of a venous thrombotic event in lung cancer patients: higher risk for adenocarcinoma than squamous cell carcinoma. J Thromb Haemost 2004;2:1760–1765.
- Bloom JW, Doggen CJM, Osanto S, Rosendaal FR. Malignancies, prothrombotic mutations and the risk of venous thrombosis. JAMA 2005;293(6):715-22.
- Cohen AT, Agnelli G, Anderson FA, et al. Venous thromboembolism (VTE) in Europe. Thromb Haemost 2007;98:756-764.
- Falanga A, Rickles FR. Pathophysiology of the thrombophilic state in the cancer patient. Semin Thromb Hemost 1999;25:173-182.
- Falanga A, Ofosu FA, Cortelazzo S, et al. Preliminary study to identify cancer patients at high risk of venous thrombosis following major surgery. Br J Haematol 1993;85:745-750.
- Ginsberg JS. Management of venous thromboembolism. N Engl J Med 1996;335:1816-1828.
- Heit JA, Silverstein MD, Mohr DN, et al. Risk factors for deep vein thrombosis and pulmonary embolism: A population-based casecontrol study. Arch Intern Med 2000:160:809–815.
- Heit JA, Mohr DN, Silverstein MD, et al. Predictors of recurrence after deep vein thrombosis and pulmonary embolism: a population-based cohort study. Arch Intern Med 2000a;160:761–768.
- Jha AK, Larizgoitia I, Audera-Lopez C, Prasopa-Plaisier N, Waters H, Bates DW. The global burden of unsafe medical care: analytic modeling of observational studies. BMJ Qual Saf 2013;22;809-15.
- Khorana AA, Francis CW, Culakova E, Kuderer NM, Lyman GH. Thromboembolism is a leading cause of death in cancer patients receiving outpatient chemotherapy. J Thromb Haemost 2007;5(3):632–634.

- Lee AYY, Levine M. Venous Thromboembolism and Cancer: Risks and Outcomes. Circulation 2003;107:1-17.
- Lyman GH, Bohlke K, Khorana AA, Kuderer NM, Lee AY, Arcelus JI, et al. Venous thromboembolism prophylaxis and treatment in patients with cancer: american society of clinical oncology clinical practice guideline update 2014. J Clin Oncol 2015;33(6):654-6.
- Rosendaal FR. Venous Thrombosis: The Role of Genes, Environment, and Behavior. Hematology 2005:1-12.
- Tapson VF. Acute pulmonary embolism. N Engl J Med 2008;358(10):1037-52.
- Weill-Engerer S, Meaume S, Lahlou A, Piette F, Saint-Jean O, Sachet A, et al. Risk Factors for Deep Vein Thrombosis in Inpatients Aged 65 and Older: A Case-Control Multicenter Study. J Am Geriatr Soc 2004;52(8):1299–1304.

Authors

- •Ștefan C. Vesa, Department of Pharmacology, Toxicology and Clinical Pharmacology, "Iuliu Hatieganu" University of Medicine and Pharmacy, 23 Gheorghe Marinescu Street, 400337, Cluj-Napoca, Cluj, Romania, EU, email: stefanvesa@gmail.com
- •Antonia E. Macarie, Department of Geriatrics and Gerontology, "Iuliu Haţieganu" University of Medicine and Pharmacy, Geriatric Unit, Municipal Clinic Hospital, 11 Tăbăcarilor Street, 400139, Cluj-Napoca, Cluj, Romania, EU, email: macarieantonia@yahoo.com
- •Sonia I. Vlaicu, Department of Internal Medicine, 1st Medical Clinic, University of Medicine and Pharmacy "Iuliu Hațieganu", Cluj-Napoca, 3-5th Clinicilor Street, 400139, Cluj-Napoca, Romania, EU, e-mail: vlaicus@yahoo.com
- •Bogdan Micu, Vth Departament of Surgery, "Iuliu Hatieganu" University of Medicine and Pharmacy, Muncipal Clinical Hospital, 11 Tabacarilor Street, 400139, Cluj-Napoca, Cluj, Romania, EU, email: micubogdan@yahoo.com
- •Anca D. Buzoianu, Department of Pharmacology, Toxicology and Clinical Pharmacology, "Iuliu Hatieganu" University of Medicine and Pharmacy, 23 Gheorghe Marinescu Street, 400337, Cluj-Napoca, Cluj, Romania, EU, email: abuzoianu@umfcluj.ro
- •Sorin Crişan, Department of Internal Medicine, "Iuliu Hatieganu" University of Medicine and Pharmacy, Municipal Clinic Hospital, 11 Tăbăcarilor Street, 400139, Cluj-Napoca, Cluj, Romania, EU, email: crisan.sorin@gmail.com

Citation	Vesa ŞC, Macarie AE, Vlaicu SI, Micu B, Buzoianu AD, Crişan S. Assessment of additional risk factors for deep vein thrombosis in patients with various malignancies. HVM Bioflux 2016;8(1):17-19.
Editor	Ioan V. Petrescu-Mag
Received	30 November 2015
Accepted	28 December 2015
Published Online	29 December 2015
Funding	None reported
Conflicts/ Competing Interests	None reported