



Clinical Image

Pulmonary thromboembolism in an anticoagulated oncological patient

Diana Alegre-González^{a,*}, Ramón Baeza-Trinidad^a , Leticia de Ávila-Lizarraga^b, Laura Fernández-Prado^b

^a Internal Medicine Department, Hospital San Pedro, Logroño, Spain

^b Medical Oncology Department, Hospital San Pedro, Logroño, Spain

ARTICLE INFO

Article history:

Received 15 January 2023

Received in revised form 11 February 2023

Accepted 11 August 2023

Keywords:

Thromboembolic disease

Chemotherapy

Cisplatin

© 2023 The Authors. Published by Iberoamerican Journal of Medicine. This is an open access article under the CC BY license (<http://creativecommons.org/licenses/by/4.0/>).

Tromboembolismo pulmonar en un paciente oncológico anticoagulado

INFO. ARTÍCULO

Historia del artículo:

Recibido 15 Enero 2023

Recibido en forma revisada 11 Febrero 2023

Aceptado 11 Agosto 2023

Palabras clave:

Enfermedad tromboembólica

Quimioterapia

Cisplatino

© 2023 Los Autores. Publicado por Iberoamerican Journal of Medicine. Éste es un artículo en acceso abierto bajo licencia CC BY (<http://creativecommons.org/licenses/by/4.0/>).

HOW TO CITE THIS ARTICLE: Alegre-González D, Baeza-Trinidad R, de Ávila-Lizarraga L, Fernández-Prado L. Pulmonary thromboembolism in an anticoagulated oncological patient. Iberoam J Med. 2023;5(4):186-187. doi: 10.53986/ibjm.2023.0026.

A 59 year-old woman recently diagnosed of lung adenocarcinoma and treated with Cisplatin-Pemetrexed was admitted to our hospital for one-week dyspnea. She was treated with Enoxaparine (60mg/kg/12 hours) due to ischemic stroke secondary to marantic endocarditis of the

aortic valve 3 months ago. On physical examination, she presented oxygen saturation less than 90%, tachypnea, left lung crackles and increased right lower extremity diameter. Laboratory data revealed D-Dimer elevation [952 ug/L (normal range less than 250 ug/L)] with pO₂ 69 mmHg

* Corresponding author.

E-mail address: dalegre@riojasalud.es

ISSN: 2695-5075 / © 2023 The Authors. Published by Iberoamerican Journal of Medicine. This is an open access article under the CC BY license (<http://creativecommons.org/licenses/by/4.0/>).

<https://doi.org/10.53986/ibjm.2023.0026>

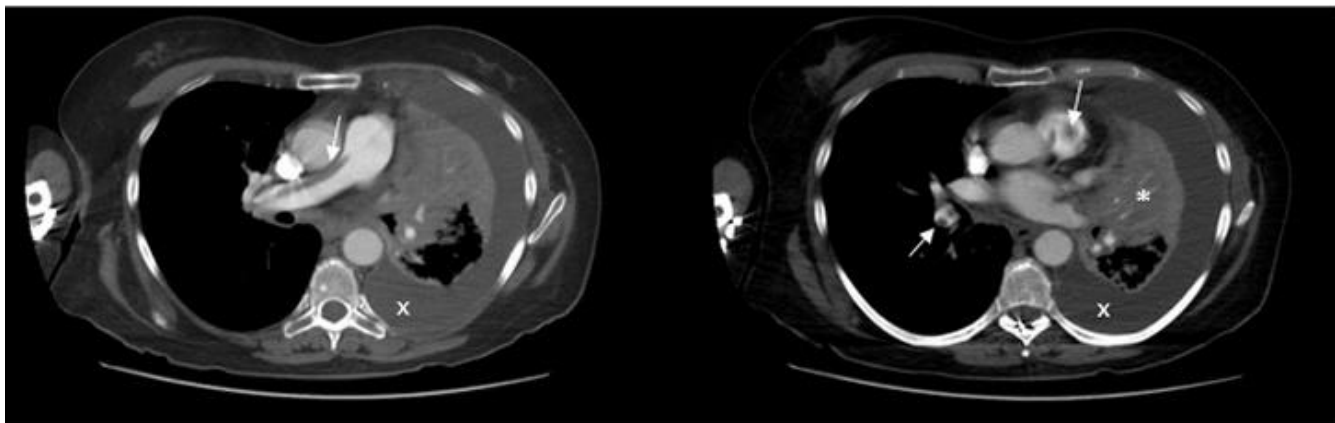


Figure 1: Chest computed axial tomography showing mediastinal- hilar mass (*), a moderate pleural effusion (x) and an embolism in the pulmonary artery (arrow).

(normal range between 80-100 mmHg) and normal pCO₂. A moderate pleural effusion with a mediastinal-hilar mass and an extensive pulmonary embolism from lower right lobe segmental branches to right ventricle was observed on pulmonary computed tomography (Figure 1). Moreover, a right deep venous thrombosis in popliteal vein extending to saphenous vein was showed at Doppler ultrasound.

Oncological patients have a higher risk of venous thromboembolism (VTE) due to a pro-inflammatory condition (activated tissue factor, inflammatory cytokines...), tumour type and treatment. Patients with advanced cancer have a higher risk of VTE during the first year of starting chemotherapy, being higher during the first month of starting chemotherapy [1, 2]. Certain chemotherapy treatment regimens (such as those based on Cisplatin and antiangiogenic drugs) [1, 3, 4] are associated with an increased risk of thromboembolic disease. Seng S et al showed VTE rate of 1.92% in patients treated with cisplatin-based and 0.79% for non-cisplatin-based chemotherapy regimens [3]. Some authors think about endothelial damage, elevation of Von Willebrand factor levels and drug-induced reduction of left ventricular function as the cause of increased risk of thrombosis related to cisplatin regimens [5]. In patients with lung adenocarcinoma and high thrombotic risk, we should exclude therapies with antiangiogenic agents and assess the use of Carboplatin treatment [3, 4]. If these alternatives were

not possible, thromboprophylaxis could be considered in those patients without bleeding risk [1, 3].

1. CONFLICT OF INTERESTS

The authors have no conflict of interest to declare. The authors declared that this study has received no financial support.

2. REFERENCES

1. Nasser NJ, Fox J, Agbarya A. Potential Mechanisms of Cancer-Related Hypercoagulability. *Cancers (Basel)*. 2020;12(3):566. doi: 10.3390/cancers12030566.
2. Khorana AA, Dalal M, Lin J, Connolly GC. Incidence and predictors of venous thromboembolism (VTE) among ambulatory high-risk cancer patients undergoing chemotherapy in the United States. *Cancer*. 2013;119(3):648-55. doi: 10.1002/ncr.27772.
3. Seng S, Liu Z, Chiu SK, Proverbs-Singh T, Sonpavde G, Choueiri TK, et al. Risk of venous thromboembolism in patients with cancer treated with Cisplatin: a systematic review and meta-analysis. *J Clin Oncol*. 2012;30(35):4416-26. doi: 10.1200/JCO.2012.42.4358.
4. Nadeau Nguyen MC, Woronow D, Binks B, Nayernama A, Jones SC. Cisplatin-Associated Aortic Thrombosis: A Review of Cases Reported to the FDA Adverse Event Reporting System. *JACC CardioOncol*. 2021;3(1):165. doi: 10.1016/j.jaccao.2020.11.017.
5. Sato N, Mishima T, Okubo Y, Okamoto T, Shiraiishi S, Tsuchida M. Acute aortic thrombosis in the ascending aorta after cisplatin-based chemotherapy for esophageal cancer: a case report. *Surg Case Rep*. 2022;8(1):75. doi: 10.1186/s40792-022-01431-8.