



## Scientific Letters

**Synchronous Cardiopulmonary Consequences of the Hypercoagulable State Associated With Cancer<sup>☆</sup>**

**Consecuencias cardiopulmonares sincrónicas del estado de hipercoagulabilidad asociado al cáncer**

To the Editor,

Venous thromboembolic disease (VTD) is a health problem of the first order, with an incidence of deep vein thrombosis (DVT) of 145 per 100,000 inhabitants and pulmonary thromboembolism (PTE) of 65.8 per 100,000 inhabitants.<sup>1,2</sup> Three factors are responsible for venous thrombosis: blood stasis, vascular damage, and a hypercoagulable state, that can be congenital or acquired.<sup>3</sup> The latter may be due to malignant, autoimmune, or infectious diseases. The use of complementary examinations for the detection of occult cancer in patients with VTD is controversial, so tools are being developed to improve the identification of these patients.<sup>4</sup> The manifestations of hypercoagulability associated with cancer include the common, conventional conditions, such as pulmonary thromboembolism (PTE), deep vein thrombosis (DVT), and migratory thrombophlebitis, more uncommon entities, such as arterial thrombosis, and the rare, difficult-to-diagnose entities, such as non-bacterial thrombotic endocarditis (NBTE).<sup>5</sup>

We retrieved very few references from the literature on the subject (Medline and Pubmed search engines, keywords “endocarditis, non-infective” and “pulmonary embolism”). Only 3 cases have been reported in which both entities coexist, and all of them were associated with underlying cancers of the lung,<sup>6</sup> ovaries,<sup>7</sup> and pancreas.<sup>8</sup>

We report the case of a 64-year-old man, former smoker of 20 pack-years who gave up 5 years previously, with a personal history of vitiligo, receiving treatment with omeprazole 20 mg. He consulted due to sudden onset of dyspnea, pleuritic-type chest pain, and syncope. Physical exploration revealed O<sub>2</sub> saturation 87% and tachycardia 134 beats per minute. Most significant findings included a systolic murmur I/VI in the mitral valve, non-tender hepatomegaly measuring 3 cm on the costal margin, and a solid mass in the right lower limb. Complete blood count (CBC) showed microcytic anemia (hemoglobin 9.9 g/dl), with ferritin 1728 ng/ml (normal values [NV] 30–400 ng/ml). Biochemistry was significant

for lactate dehydrogenase 814 U/l (NV: 125–250 U/l), glutamic pyruvic transaminase 71 U/l (NV: 2–33 U/l), C-reactive protein 193 mg/l (VN 0–5 mg/l), and D-dimer >20,000 ng/ml. Computed tomography angiography (CTA) showed filling defects in the arteries of the lower right and left lobes compatible with PTE (Fig. 1a). Lower limb venous Doppler showed a femoro-popliteal DVT in the right lower limb. Based on these findings, the patient was admitted and began treatment with enoxaparin adjusted for weight every 12 h.

The patient reported a weight loss of 6 kg in 2 months, asthenia, and fever. Given this constitutional syndrome, an abdominal CT scan was performed, which revealed a mass measuring 38 × 37 mm in the body of the pancreas and multiple liver masses consistent with pancreatic cancer with liver metastasis. Fine-needle aspiration with endoscopic ultrasonography confirmed the presence of pancreatic adenocarcinoma.

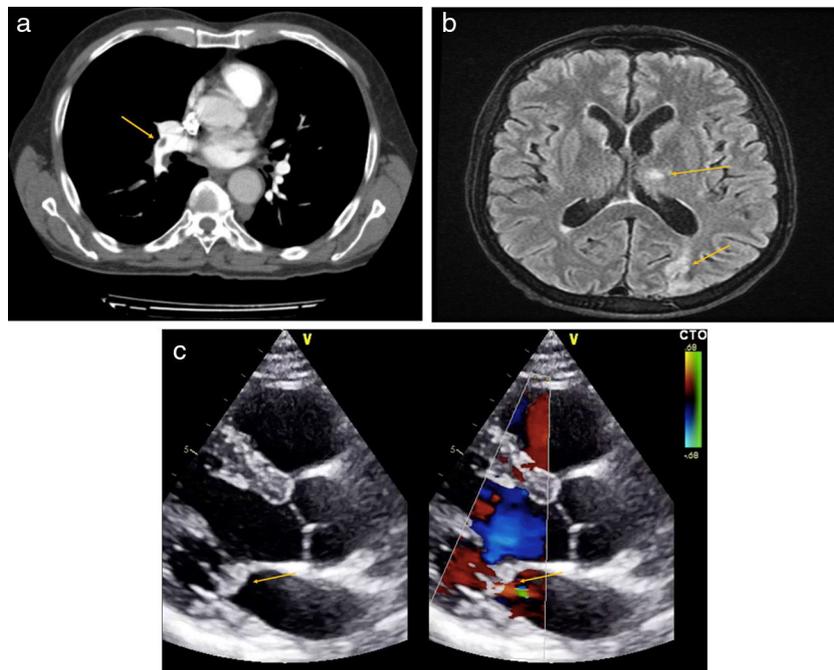
Seventy-two hours later, the patient presented loss of strength in the right upper limb and reduced level of consciousness. The neurological examination revealed a loss of strength in the proximal right arm, accompanied by an increased base of support and positive Romberg test. CT of the brain revealed hypodense lesions without contrast uptake in both cerebellar hemispheres and the right occipital lobe. Brain magnetic resonance imaging (MRI) confirmed the presence of ischemic lesions (Fig. 1b). In view of suspected embolic syndrome, we performed a transthoracic echocardiogram that showed severe mitral insufficiency and endocardial masses on the mitral valve, while the presence of patent foramen ovale was ruled out (Fig. 1c). Blood cultures were repeatedly negative, as were serologies for *Coxiella* spp., *Brucella* spp., *Bartonella* spp., and *Legionella* spp.

With a diagnosis of VTD, NBTE, and stroke of embolic origin caused by a hypercoagulable state due to stage IV pancreatic adenocarcinoma (T2N1M1), the treating physicians decided to initiate treatment with tinzaparin 14,000 IU every 24 h and palliative chemotherapy.

NBTE, formerly known as marantic endocarditis, is a rare clinical entity that has devastating consequences.<sup>5</sup> It may appear concurrently with VTD as an expression of a procoagulant state or be the primary source of the embolic disease.

Ziegler<sup>9</sup> first described NBTE in 1888 using the term “thromboendocarditis”, but it was not until 1954 that Angrist and Marquiss highlighted the strong association of systemic emboli with this entity.<sup>10</sup> It is characterized by the presence of fibrin vegetations on the heart valves in the absence of bacterial infection.<sup>5,11</sup> The valves typically affected are the mitral and aortic valves, although the involvement of right heart valves has also been less commonly reported.

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**Fig. 1.** (a) CT angiography of pulmonary arteries. Filling defect in arteries of the right and left lower lobes consistent with PTE, arrow. (b) Brain MRI in T2. Ischemic impacts in cerebellar hemispheres and in the occipital region, arrows. (c) Echocardiogram in the long parasternal axis. Severe mitral insufficiency and endocardial masses on mitral valve, indicated with arrows.

The incidence on autopsy ranges between 0.3% and 9.3%.<sup>11</sup> This contrasts with the low incidence of diagnosis in clinical practice,<sup>8,12</sup> and demonstrates the need for a high clinical suspicion. Like VTD, it is associated with a wide variety of procoagulant states, such as autoimmune diseases, infections, and cancer, especially of the pancreas and lung.<sup>13</sup>

The pathogenesis of NBTE involves the interaction between macrophages and malignant cells, which releases cytokines that damage the endothelium and promote the aggregation and deposition of platelets and thrombus formation. Overactivation of the coagulation cascade provokes a hypercoagulable state, which fosters the development of thrombi. This leads to the growth of sterile vegetations composed of fibrin and blood platelets.<sup>14</sup>

Clinically, it manifests with systemic embolisms and, more rarely, valvular dysfunction. Heart murmurs are uncommon, and tend to be systolic if present, complicating the suspected diagnosis. Diagnosis is clinical, accompanied by the demonstration of vegetations on echocardiography.

The main sites of embolization are the spleen, kidney, and limbs, while embolic events in the coronary arteries and the central nervous system produce greater morbidity and mortality.<sup>9,10</sup> Although it tends to occur in the left valves, pulmonary circulation is frequently affected by the embolic phenomenon.<sup>15</sup>

Treatment is based on control of the underlying disease and anticoagulation with heparin or low molecular weight heparin; vitamin K antagonists should be avoided. Valvular surgery should be considered an option in selected patients.<sup>8</sup>

We believe that this case of synchronous presentation of NBTE and PTE illustrates the need for clinical suspicion in patients with VTD and a clinical picture of systemic embolism, after other more common entities such as patent foramen ovale have been ruled out. NBTE is a rare entity that is difficult to diagnose, but one that has great prognostic relevance for the patient.

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Javier Miguel Martín Guerra,\* Miguel Martín Asenjo,  
Carlos Jesús Dueñas Gutiérrez, Inmaculada Gil González

Servicio de Medicina Interna, Hospital Clínico Universitario de  
Valladolid, Valladolid, Spain

\* Corresponding author.

E-mail address: [javi6vega@hotmail.com](mailto:javi6vega@hotmail.com) (J.M. Martín Guerra).

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## Extramedullary Multiple Myeloma With Pleural Involvement: A Rare Clinical Entity<sup>☆</sup>

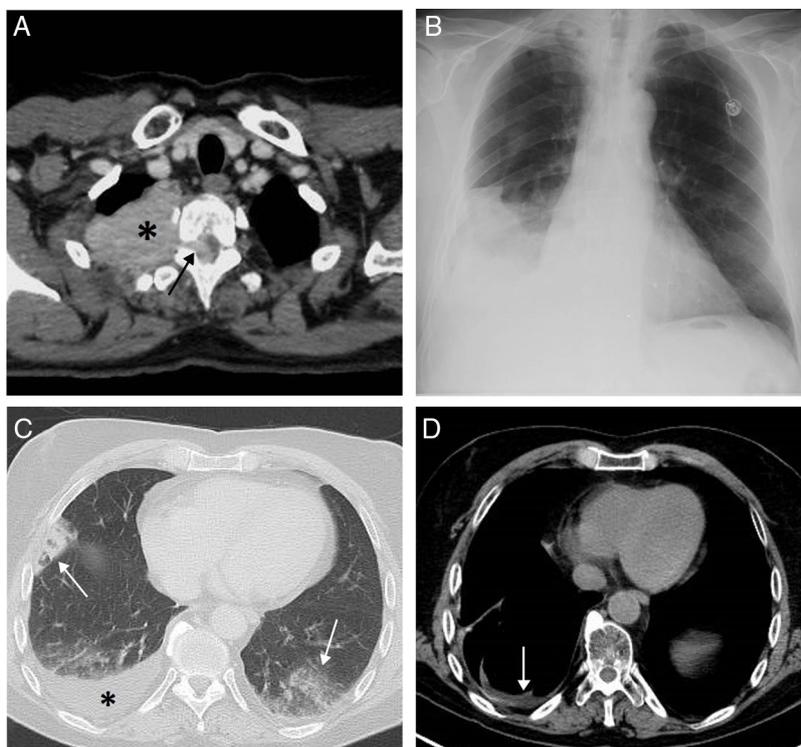


### Mieloma múltiple extramedular con afectación pleural: una rara entidad clínica

To the Editor,

Thoracic involvement has been described in the course of multiple myeloma (MM), in the form of bone lesions, extraosseous plasmacytomas, pulmonary infiltration and, exceptionally, pleural effusion (PE). Myelomatous pleural effusion (MPE) occurs in only 1% of cases of PE in patients with MM, and is associated with a poor prognosis (median survival of 1.5–3 months after appearance). We report a patient with MM with extramedullary thoracic involvement who developed secondary MPE.<sup>1,2</sup>

This was a 67-year-old man with a history of quiescent MM IgA kappa diagnosed in 1998 who attended the emergency room of our hospital in November 2017 due to fever, general malaise, asthenia, and cough. The patient had presented several plasmacytomas in the right chest wall (Fig. 1A) and the spinal canal in 2012 and 2014, but without PE, and was treated with different therapeutic options (chemotherapy, autologous hematopoietic stem cell transplantation, local radiation therapy), resulting in complete resolution of the lesions. In 2015, he presented a new extramedullary relapse in the form of a paravertebral mass in the right hemithorax, which was treated with chemotherapy. Given the patient's clinical picture and radiographic findings (not present in previous studies) of right PE observed in the emergency department (Fig. 1B), we decided to admit him and perform a positron emission tomography/computed tomography (PET/CT).



**Fig. 1.** (A) Axial CT scan of the chest (mediastinum window) performed in 2012 that identifies a solid mass in the upper right hemithorax (asterisk) infiltrating the chest wall and penetrating the spinal canal through the right T2–T3 intervertebral foramen (arrow). (B) Posteroanterior chest X-ray performed in November 2017 showing right pleural effusion for the first time in this patient. (C) Chest axial image of a PET/CT scan performed in November 2017 which identifies bilateral opacities of pneumonic aspect (arrows) and right pleural effusion (asterisk). (D) Thoracic axial image of another PET/CT scan performed in March 2018 which shows the disappearance of pneumonic opacities and the presence of minimal pleural effusion.

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