

Fig. 1. Chest X-ray showing a mass shadow in the left hilar region and bilateral diffuse interstitial opacities (A). Computed tomography revealing a mass in the left upper pulmonary lobe (B), pulmonary carcinomatosis lymphangitis, and minor bilateral pleural effusions (C). Chest X-ray showing marked improvement after four weeks of treatment (D).

depending on the general condition of the patient, such as respiratory failure, which is often a time-limited situation in such patients. Liquid biopsy is useful when tissue sampling is difficult or inadequate.² Although liquid biopsy can only currently be used in clinical practice for EGFR sensitizing and resistance mutations, various NSCLC driver genes have been detected successfully with the help of several plasma-based tests including molecular tests (such as detection of ROS-1) even before proceeding to tissue-based tests.^{3–6}

In the present case, the time gap between the plasma EGFR mutation test and result report was only three days. This case indicates that liquid biopsy for detecting other driver gene mutations in addition to the EGFR mutation are warranted, since a therapeutic effect can be expected in the patient with driver gene mutation-positive NSCLC; this is true even in cases where tissue tests are difficult, such as in patients with lymphangitic carcinomatosis, which generally have poor prognosis.

In summary, we reported a case of EGFR mutation positive lung adenocarcinoma with acute severe respiratory failure caused by pulmonary lymphangitic carcinomatosis. Although tissue collection was difficult, the patient could receive appropriate therapy by using liquid biopsy. Liquid biopsy could help in addressing the problem of poor prognosis in lymphangitic carcinomatosis patients.

References

1. Klimek M. Pulmonary lymphangitis carcinomatosis: systematic review and meta-analysis of case reports, 1970–2018. Postgrad Med. 2019;131:309–18.
2. Rolfo C, Mack PC, Scagliotti GV, Bass P, Barlesi F, Bivona TG, et al. Liquid biopsy for advanced non-small cell lung cancer (NSCLC): a statement paper from the IASLC. J Thorac Oncol. 2018;13:1248–68.

Rib Osteomyelitis Mimicking a Pleural Empyema in a Patient with Invasive Pneumococcal Disease*

Osteomielitis costal que simula un empiema pleural en una enfermedad neumocócica invasiva

Dear Editor,

Invasive pneumococcal disease (IPD) is a serious form of *Streptococcus pneumoniae* infection that can produce a wide range of



clinical manifestations including septicemia, meningitis, arthritis, osteomyelitis, cellulitis, endocarditis,¹ etc., which confers significant morbidity and mortality (especially in children under 2 years of age and in adults at risk due to medical conditions and/or advanced age). Incidence is variable, although in developed countries it is estimated to affect about 10 cases per 100 000 individuals/year in the general population.² We report the case of a patient with IPD who, after a non-consolidating acute respiratory infection, developed several thoracic and extrathoracic infectious complications.

Our patient was a 67-year-old woman with no personal clinical history of interest, except smoking (pack-year index of 30) until 7 years previously. She consulted with a clinical picture of respiratory infection (without radiographic consolidation [Fig. 1A]). *S. pneumoniae* susceptible to multiple drugs was isolated from a blood

* Please cite this article as: Gorospe Sarasúa L, Martín-Pinacho JJ, Ayala-Carbonero AM, Gambí-Pisonero E, Martín-Dávila P, Mirambeaux-Villalona RM, et al. Osteomielitis costal que simula un empiema pleural en una enfermedad neumocócica invasiva. Arch Bronconeumol. 2020;56:535–537.

Akihito Okazaki ^{a,c,*}, Kazuhiko Shibata ^b,
Yasuhiro Matsuda ^a, Yasutaka Shiba ^a, Keiichi Iwasa ^b,
Kazuo Kasahara ^c

^a Department of Respiratory Medicine, Koseiren Takaoka Hospital, 5-10, Eiraku-machi, Takaoka 933-8555, Japan
^b Department of Medical Oncology, Koseiren Takaoka Hospital, 5-10, Eiraku-machi, Takaoka 933-8555, Japan
^c Department of Respiratory Medicine, Kanazawa University Faculty of Medicine, Institute of Medical, Pharmaceutical and Health Sciences, 13-1, Takara-machi, Kanazawa 920-8641, Japan

* Corresponding author.

E-mail address: akihitookazaki1017@gmail.com (A. Okazaki).

<https://doi.org/10.1016/j.arbres.2020.03.032>
0300-2896/ © 2020 SEPAR. Published by Elsevier España, S.L.U. All rights reserved.

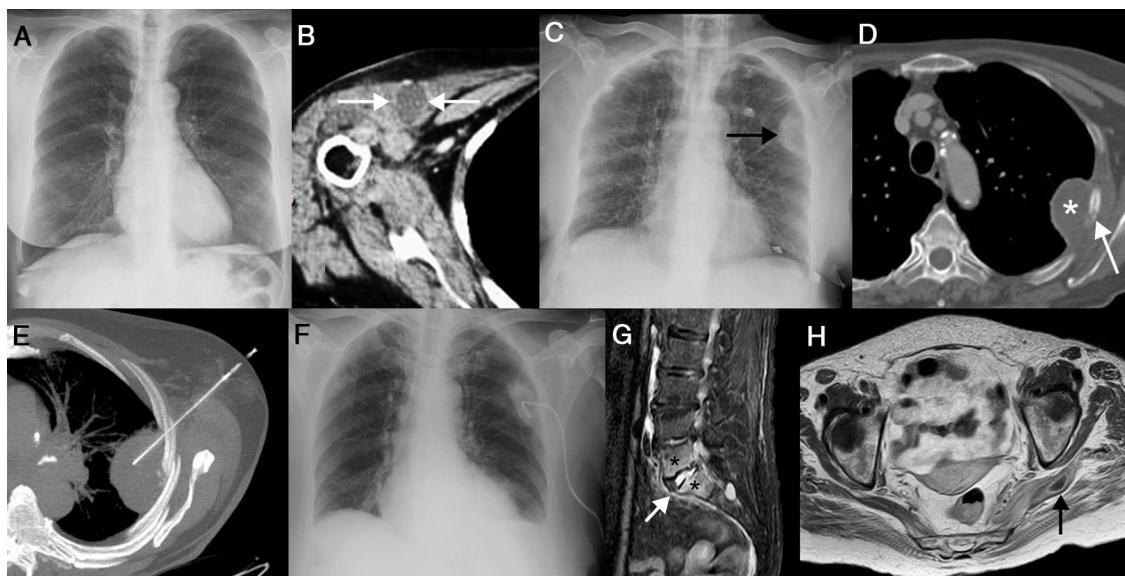


Fig. 1. (A) Posteroanterior chest X-ray showing no pneumonic consolidation. (B) Chest CT axial image showing a collection in the thickness of the right pectoralis major muscle (arrows). (C) Anteroposterior chest X-ray identifying a new extraparenchymal lesion. (D) Chest CT axial image showing a focal lytic lesion in the left fourth costal arch (arrow) surrounded by an extrapleural fluid collection (asterisk). (E) Chest CT axial image (maximum intensity projection) showing the percutaneous drainage procedure of the left hemithorax collection. (F) Follow-up anteroposterior chest X-ray showing radiological improvement of the collection. (G) MRI sagittal image (inversion-recovery sequence with fat suppression) identifying typical signs of acute spondylodiscitis: increased signal intensity in the L5-S1 intervertebral disc (arrow) and altered signal of the vertebral endplates adjacent to the disc (asterisks). (H) MRI axial image (T1-weighted sequence after intravenous gadolinium administration) showing a ring-enhancing collection in the thickness of the left pyramidal muscle (arrow).

culture kit. The patient, who was not vaccinated against pneumococcus, was initially treated with azithromycin, and showed rapid improvement in her respiratory symptoms. Within days of completing treatment, the patient developed low-grade fever and mixed mechanical and inflammatory pain in the right shoulder, which caused severe functional loss; physical examination showed right pre-pectoral erythema, increased ipsilateral shoulder volume, and intense pain in active and passive mobilization. Although an X-ray of the right shoulder showed no bone erosions, ultrasound and a computed tomography (CT) scan of the right shoulder confirmed joint effusion and the existence of periarticular collections (located in the pectoral musculature, Fig. 1B). Ultrasound-guided arthrocentesis demonstrated the presence of pus and encapsulated gram-positive cocci in the right glenohumeral joint (pneumococci were subsequently isolated in synovial fluid culture). Arthrotomy of the right shoulder and surgical cleaning of the periarticular collections were performed. A combined treatment of ceftriaxone and vancomycin was started, resulting in progressive improvement. Five days after joint drainage, the patient reported pleuritic-type pain in the left hemitorax. Chest X-ray showed the appearance of an extraparenchymal collection in the left hemitorax, initially interpreted as a pleural collection (Fig. 1C). A chest CT showed focal lysis of the left fourth costal arch surrounded by a collection. No parenchymal consolidations were identified (Fig. 1D). In view of these findings, interpreted as left costal osteomyelitis complicated by an extrapleural abscess, percutaneous drainage of this collection was performed with radiological control (Fig. 1E), obtaining purulent material (pneumococci confirmed on culture). Drainage continued for several days and several washings were performed with physiological serum, and a clinical and radiological improvement of this infectious focus was observed (Fig. 1F). Ten days after drainage, the patient experienced progressive and increased pain in the lumbar region radiating to the left gluteal and ipsilateral lower limb. A CT scan and magnetic resonance imaging (MRI) of the lumbar spine confirmed signs of spondylodiscitis of the L5-S1

intervertebral space (Fig. 1G) and an abscess in the left pyramidal muscle (Fig. 1H), which were treated conservatively (continued parenteral antibiotic treatment). The patient was discharged a few days later and did not develop any infectious complications again, after maintaining antibiotic treatment for an additional 8 weeks.

Humans are the only known natural reservoirs of *S. pneumoniae* (pneumococcus). More than 30% of children and 5%–10% of adults are asymptomatic carriers of pneumococcus in their nasopharynx; however, this bacteria has a remarkable ability to spread to the airway (upper and lower) and invade the bloodstream or lymph vessels, causing a wide variety of infectious diseases.³ IPD represents the most serious form of pneumococcal infection, and can produce a wide range of clinical manifestations (from meningitis to endocarditis). This is a severe condition with significant morbidity and mortality (especially in children with hemoglobinopathies and immunosuppressed adults) and a variable incidence (10 cases per 100 000 individuals/year in developed countries).^{1,2}

Spontaneous osteomyelitis and septic arthritis (not occurring in patients surgically operated on a bone or joint) due to pneumococcus are very rare and particularly affect immunocompromised children, adolescents, or adults (especially patients with human immunodeficiency virus infection and with asplenia/hypoplasia).^{4,5} Our patient did not have any predisposing factors that increased the risk of pneumococcal osteomyelitis and arthritis. Generally, the bone manifestations of pneumococcal osteomyelitis usually occur in patients with upper or lower respiratory tract infections (more or less clinically or radiologically evident), as in our patient. Symptoms are generally non-specific and include fever, chills, local pain, and increased soft tissue in the affected bone/joint.⁴ In our case, the patient developed several sequential joint and bone complications, despite the initial treatment of a non-consolidating respiratory infection, with low-grade fever and pain in different sites. Pneumococcal

glenohumeral arthritis and spondylodiscitis are described in the literature, as is *S. pneumoniae* pyomyositis.^{6–8} Costal osteomyelitis is extremely rare (less than 1% of hematogenous osteomyelitis), and may be caused by: (1) penetrating trauma; (2) regional spread from a pneumonic empyema or focus; or (3) hematogeneous transmission. The most common causative microorganisms are *Staphylococcus aureus* and mycobacteria.⁹ To our knowledge, no similar cases of pneumococcal osteomyelitis in adults have been described in the literature. The only documented case of spontaneous pneumococcal costal osteomyelitis occurred in a 4-month-old infant, and eventually required partial resection of the rib after failed percutaneous fine-needle aspiration.¹⁰ The authors recommend surgical resection for the proper management of costal osteomyelitis. In our case, attention is drawn to the initial radiological presentation of this osteomyelitis in the form of an extraparenchymal collection, which suggested pleural empyema. A chest CT scan showed a focus of costal osteolysis surrounded by an extrapleural collection, differentiating it from pleural empyema. Percutaneous drainage was performed using a pigtail catheter with radiological control, achieving clinical and radiological improvement of this complication. Although the serotype was not investigated in this case, it is important to remember the importance of routinely investigating the different serotypes of *S. pneumoniae* isolates in invasive diseases for possible prevention.

We believe that early use of imaging tests for the diagnosis of infectious complications of IPD, as well as early surgical drainage of treatable infectious foci, are necessary, in addition to systemic antibiotic therapy, for optimal diagnostic and therapeutic management of these patients. We found no previously reported cases in the literature of adults with extrapleural collection associated with costal osteomyelitis in the context of an IPD, and we believe that early percutaneous drainage of this collection may contribute to the favorable control of the infection.

References

- Fitzgerald D, Waterer GW. Invasive pneumococcal and meningococcal disease. *Infect Dis Clin North Am*. 2019;33:1125–41.
 - Rueda AM, Serpa JA, Matloobi M, Mushtaq M, Musher DM. The spectrum of invasive pneumococcal disease at an adult tertiary care hospital in the early 21st century. *Medicine (Baltimore)*. 2010;89:331–6.
 - Yildirim I, Shea M, Pelton SI. Pneumococcal disease in the era of pneumococcal conjugate vaccine. *Infect Dis Clin North Am*. 2015;29:679–97.
 - Pääkkönen M, Peltola H. Bone and joint infections. *Pediatr Clin North Am*. 2013;60:425–36.
 - Baraboutis I, Skourelis A. *Streptococcus pneumoniae* septic arthritis in adults. *Clin Microbiol Infect*. 2004;10:1037–9.
 - Lotz H, Strahm C, Zdravkovic V, Jost B, Albrich WC. Septic arthritis due to streptococci and enterococci in native joints: a 13 year retrospective study. *Infection*. 2019;47:761–70.
 - Forestier E, Sordet C, Cohen-Solal J, Remy V, Javier RM, Kuntz JL, et al. Bone and joint infection due to *Streptococcus pneumoniae* in two immunocompetent adults. *Joint Bone Spine*. 2006;73:325–8.
 - Wong SL, Anthony EY, Shetty AK. Pyomyositis due to *Streptococcus pneumoniae*. *Am J Emerg Med*. 2009;27:633.e1–3.
 - Nibley PD, Kraus CK. Rib osteomyelitis in a pediatric patient: case report and review of the literature. *Clin Pract Cases Emerg Med*. 2018;2:294–6.
 - Kalouche I, Ghanem I, Kharrat K, Dagher F. Osteomyelitis of the rib due to *Streptococcus pneumoniae*: a very rare condition in children. *J Pediatr Orthop B*. 2005;14:55–60.
- Luis Gorospe Sarasúa^{a,*}, Jesús Javier Martín-Pinacho^a, Ana María Ayala-Carbonero^a, Esther Gambí-Pisonero^a, Pilar Martín-Dávila^b, Rosa Mariela Mirambeaux-Villalona^c, Gemma María Muñoz-Molina^d
- ^a Servicio de Radiodiagnóstico, Hospital Universitario Ramón y Cajal, Madrid, Spain
^b Servicio de Enfermedades Infecciosas, Hospital Universitario Ramón y Cajal, Madrid, Spain
^c Servicio de Neumología, Hospital Universitario Ramón y Cajal, Madrid, Spain
^d Servicio de Cirugía Torácica, Hospital Universitario Ramón y Cajal, Madrid, Spain
- Corresponding author.
E-mail address: luisgorospe@yahoo.com (L. Gorospe Sarasúa).
- <https://doi.org/10.1016/j.arbr.2020.04.004>
1579-2129/ © 2020 SEPAR. Published by Elsevier España, S.L.U. All rights reserved.

Thrombus in Transit[☆]



Trombo en tránsito

Dear Editor,

Thrombi in right cardiac cavities, described in less than 4% of patients with pulmonary embolism, are rare and in most cases confer a poor prognosis.^{1–3} Nevertheless, there is no consensus on the treatment of choice in these patients.

We report the case of a 72-year-old man, former smoker, diagnosed with severe COPD, Parkinson's disease and Pick's disease, who consulted the emergency department for a sudden onset of loss of strength and sensitivity in the right side of the face and right hand, along with dysarthria. The stroke code was activated, and a perfusion computed tomography (CT) was per-

formed which revealed early signs of acute infarction in the left middle cerebral artery. Due to the low initial deficit and the progressive improvement of the clinical picture, fibrinolysis was rejected.

During his stay in the emergency department, the patient developed epigastric pain. An electrocardiogram was performed showing negative T waves in DI to DII, in aVF and V1–V6. Serial troponin T determinations rose from 38 to 570 ng/L.

Acute coronary syndrome was suspected, so a transthoracic echocardiogram was performed, which revealed an image consistent with very long, mobile thrombi in both the left atrium and the right atrium each measuring approximately 4 cm, that appeared to cross continuously through a patent foramen ovale, suggestive of thrombus in transit (Fig. 1). Findings consistent with severe pulmonary hypertension and severe right ventricular dilatation and dysfunction were also observed.

To confirm the presence of thrombus in transit, the study was completed with a transesophageal echocardiogram, and a chest CT was requested in which acute pulmonary thromboembolism (PTE)

[☆] Please cite this article as: Chasco L, Ballaz A, Oribe MA, Artaraz A, Aramburu A, Velasco S, et al. Trombo en tránsito. *Arch Bronconeumol*. 2020;56:537–539.