Septic Pulmonary Embolism in a Child

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Abstract

Septic pulmonary embolism (PE) is a diagnosis seldom considered in children. A 15-year-old girl presented with fever, extremity swelling and pain for 5 days. Chest radiograph revealed a large, round density in the right lung and consolidation areas in peripheral portions of both lungs. Computed tomography showed bilateral multiple round and cavitary nodules in peripheral portions of both lungs, pleural effusion in the right lung and hypodense round lesion (8×4 cm) in the superior segment of right lower lobe, characteristic of PE. Cefotaxime and clindamycin were administered. By the 5th hour, the patient passed away. Staphylococcus aureus was isolated from blood, pleural fluid and tracheal aspirate cultures.

Key words: Children, Septic pulmonary embolism, Staphylococcus aureus.

Introduction

Septic pulmonary embolism (PE) is an uncommon condition in children(1). Numerous pulmonary infarcts resulting from small emboli may be associated with right sided bacterial endocarditis, septic thrombophlebitis and osteomyelitis(2). The

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clinical picture is variable, often suggests pneumonia and the diagnosis may not be established until autopsy(3). Septic PE usually presents as fever, hemoptysis and patchy infiltrates on chest radiographs(4). We hereby present a case of septic PE in a child.

CASE REPORT

A previously healthy 15-year-old girl with a 5-day history of sprain in the right ankle was admitted to emergency ward. She presented with fever, pain, erythema, and swelling on the left leg, left arm and right ankle. Physical examination showed fever of 38.5°C; wide-spread rales on right lung fields, swelling, erythema and tenderness at the left leg, left arm and right ankle. Laboratory analysis showed white blood cell count of 7800/mm³ (72% neutrophils, 22% lymphocytes, 4% monocytes, 2% eosinophils), hemoglobin 12.4 g/dL, platelet 172000/mm³, erythrocyte sedimentation rate 30 mm/h and C-reactive protein 7.5 mg/dL. Chest radiograph revealed large sized round density in the right lung and consolidation areas in peripheral portions of both lungs (Fig 1). There was soft tissue



Fig. 1. Chest radiograph revealed large, round density in the right lung and consolidation areas in peripheral portions of both lungs.

swelling on extremity radiographs. At the begining cefotaxime and clindamycin were initiated for soft tissue infection, however clindamycin was later replaced with vancomycin, because of suspicion of methicillin resistant *Staphylococcus aureus* (MRSA) infection. By the second hour, she had developed progressive respiratory distress and fever. Computed tomography (CT) showed bilateral multiple round and cavitary nodules in peripheral portions of both lungs and, pleural effusion in the right lung (*Fig* 2). Other chest CT section revealed a thin walled, hypodense round lesion (8×4 cm) in the right lower lobe superior segment (*Fig* 3). Albendazole therapy was also added because we could not rule out the possibility of hydatid cysts. By the 3rd hour of



Fig. 2. Computed tomography showed bilateral multiple round (black arrow) and cavitary nodules (white arrow) in peripheral portions of both lungs and, pleural effusion (plus sign) in the right lung.

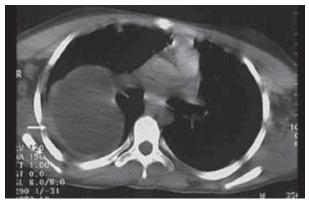


Fig. 3. Computed tomography revealed a thin walled, hypodense round lesion (8x4 cm) (arrow) in the superior segment of right lower lobe.

admission, progressive deterioration resulted in respiratory failure, requiring mechanical ventilation. The patient developed septic shock. Septic shock treatment was administered according to our clinic protocol (intravenous fluid, dopamin and adrenalin perfusion). Echocardiography proved normal. By the 5th hour, the patient passed away. Methicillin sensitive *Staphylococcus aureus* (MSSA) was isolated from blood, pleural fluid and tracheal aspirate fluid cultures. Serum *Echinococcus granulosis* IgE and its indirect hemagglutination were both negative. Lymphocyte subsets, immunoglobulin level, and tetanus antibody response were normal.

DISCUSSION

Septic PE is an uncommon disorder with an insidious onset and is difficult to diagnose(5). Septic PE can progress to abscesses, empyema and bronchopleural fistula(6). Our patient presented with fever as well as signs and symptoms of soft tissue infections. To rule out the possibility of lung abscesses, patient underwent a CT, which showed bilateral multiple round and cavitary nodules in peripheral portions of both lungs and a hypodense lesion (8×4 cm) in the right lower lobe superior segment. Septic PE diagnosis was confirmed by CT in our patient. Typical radiographic features of septic PE include patchy air space lesions simulating nonspecific bronchopneumonia; multiple round or wedge shaped densities of varying sizes from 0.5 to 3.5 cm located peripherally on chest CT scans (7,8). It has been reported that empyema, bronchopleural fistula, pneumothorax, cavity and abscess formations are common in septic PE(7-9). While most frequent underlying cause is a medical device, soft tissue infections may also constitute a focus for septic PE. Wong, et al.(10) reported that the commonest causes of septic PE were soft tissue and bone infections. Primary staphylococcal pneumonia is usually unilateral. In our patient, pulmonary features, include bilateral, peripherally located multiple round and cavitary nodules, rapid progression of cavities and empyema, are compatible with septic embolisation. The cause of septic PE in our patient was staphylococcal bacteremia secondary to soft tissue infection. Staphylococci are the most commonly isolated pathogens in patients developing

septic PE(5,10), Anticoagulation is not used in cases of septic embolisation because of higher chances of bleeding in the area of infected embolus and lack of benefit of anticoagulation therapy(6). Eradication of infection is the cornerstone in the management of septic PE(6). Septic embolisation caused by MRSA is best treated with a lactam antibiotic. Our patient was treated with intravenous vancomycin plus cefotaxime, however, she failed to respond to antibiotic therapy. Lee, et al.(11) reported that radiological or surgical interventions performed on the sites where the emboli originated in addition to antimicrobial therapy. In that study, six patients underwent drainage of the extrapulmonary infection site. Pigtail catheter drainage was performed for liver abscess and for thigh cellulitis(11). In our patient, drainage of the soft tissue infection was not performed due to the rapid deterioration of patient's condition.

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REFERENCES

- 1. Stern RC. Pulmonary Embolism and Infarction. In: Nelson Textbook of Pediatrics, Behrman R, Kliegman RM, Jenson HB (eds), 17th ed. Philadelphia: WB Saunders; 2004. p. 958-972.
- MacMillan JC, Milstein SH, Samson PC. Clinical spectrum of septic pulmonary embolism and infarc-tion. J Thorac Cardiovasc Surg 1978; 75: 670-679.

- 3. Evans DA, Wilmott RW. Pulmonary embolism in children. Pediatr Clin North Am 1994; 41: 569
- 4. Connolly J, Tarver RD, Meyer C, Winer-Muram H. Fever and patchy infiltrates: pulmonary septic emboli. Semin Respir Infect 2002; 17: 85-88.
- 5. Cook RJ, Ashton RW, Aughenbaugh GL, Ryu JH. Septic pulmonary embolism: presenting features and clinical course of 14 patients. Chest 2005; 128: 162-166.
- Aslam AF, Ahmad KA, Thakur CT, Vasavada BC, Khan IA. Staphylococcus aureus infective endocarditis and septic pulmonary embolism after septic abortion. Intern J Cardiol 2005; 105:233-235.
- Huang RM, Naidich DP, Lubat E, Schinella R, Garay SM, McCauley DI. Septic pulmonary emboli-CT radiographic correlation. AJR 1989; 153: 41-45.
- 8. Rossi S, Goodman PC, Franquet T. Nonthrombotic pulmonary emboli. AJR 2000; 174: 1499-1508.
- 9. Kuhlman JE, Fishman EK, Teigen C. Pulmonary septic emboli: diagnosis with CT. Radiology 1990; 174: 211-223.
- Wong KS, Lin TY, Huang YC, Hsia SH, Yang PH, Chu SM. Clinical and radiographic spectrum of septic pulmonary embolism. Arch Dis Child 2002; 87: 312-315.
- 11. Lee SJ, Cha SI, Kim CH, Park JY, Jung TH, Jeon KN, *et al.* Septic pulmonary embolism in Korea: Microbiolgy, clinicoradiologic features, and treatment outcome. J Infect 2007; 54: 230-234.