Role of multidetector spiral CT scanning for pulmonary embolism confirmation in a child with pulmonary hypertension: a case report

Heda Melinda Nataprawira¹, Sri Endah Rahayuningsih¹, Nono Sumarna Afandi¹, Armijn Firman¹, Tan Siauw Koan²

Pulmonary embolism (PE) is associated with considerable morbidity and mortality. Early diagnosis and prompt treatment is essential,¹,² however PE is rarely clinically diagnosed or treated in children. Most clinically significant PE is not recognized antemortem.³ While its diagnosis remains a challenge as the signs and symptoms can often be non-specific, an accurate diagnosis is essential for the management of this disease. It is known that a number of non-invasive diagnostic tools are available for its detection nowadays.¹²,⁴ Even though multi-detector spiral, also called helical, CT scanning is promising and has been proven to be useful in diagnosing this condition with high sensitivity and specificity,⁵ it is unavailable even in referral hospitals in Indonesia. The gold standard, pulmonary angiography, is considered as the procedure of choice to diagnose PE, but unfortunately it is invasive. Failure to diagnose PE accurately and promptly can result in excess morbidity and death due to pulmonary hypertension (PH) and recurrent venous thromboembolic events. Conversely, unnecessary anticoagulation therapy poses a risk without any benefit.²

Pulmonary embolism should be considered in the evaluation of unexplained PH, respiratory insufficiency, and disseminated intravascular coagulation in children.³ A diagnosis of PH can be made when the mean pulmonary artery pressure is greater than 25 mmHg in a resting individual at sea level.⁶ Pulmonary hypertension refers to a group of conditions with multiple causes rather than a single condition. PH is caused by increased pulmonary blood flow as seen in congenital heart defects with large left-to-right shunts (hyperkinetic pulmonary hypertension), alveolar hypoxia, increased pulmonary venous pressure, primary pulmonary vascular disease and chronic thromboembolic disease.⁶,⁷ An extensive evaluation should be performed in children with severe PH, as the most successful strategy for managing PH involves treatment of any underlying disorders. A specific phosphodiesterase inhibitor, sildenafil, is known as a potent and selective pulmonary vasodilator that has been proven to be superior in decreasing the mean pulmonary artery pressure.
pulmonary artery pressure. It is also equally effective and selective in reducing pulmonary vascular resistance compared to inhaled nitric oxide (iNO) treatment.6,8

The Case

An 11-year-old boy was referred to Hasan Sadikin Hospital with fatigue as the chief complaint. Eight months before admission he felt fatigue that was sometimes accompanied by shortness of breath. During the last two months before admission, these complaints had become worse, accompanied by sudden onset left chest pain. He also suffered from two episodes of syncope and had infrequent coughing symptoms. There were no symptoms of chronic pulmonary problems or disease. He was previously described as healthy and similar histories were not found in family members. Echocardiography examination revealed PH (pulmonary artery pressure of 140 mmHg) with severe tricuspid insufficiency but no cardiac defect was found (Figure 1).

He was referred to Cardiac Center at Harapan Kita Hospital where a lung perfusion scintigraphy examination was performed. This procedure revealed primary pulmonary hypertension (PPH) with suspected PE (Figure 2). He was then given oral sildenafil and the anticoagulant warfarin, to maintain the international normalized ratio (INR) within the range of 1.5–2.0. Afterwards, he received follow-up care at Hasan Sadikin Hospital and at a private clinic.

On physical examination, he was fully alert; the pulse rate was 96 beats/minute and was regular. The respiratory rate was 24 times/minute, and blood pressure was 120/80 mmHg. No cyanosis or clubbing of the fingers was noted. On auscultation of the heart, the first sound was normal with a loud second heart sound. A 3/6 systolic murmur in the third and fourth intercostal spaces was noted. No liver enlargement was found. Laboratories findings were within normal limit including an INR of 1.5–2.0. Chest X-rays showed cardiomegaly, while an electrocardiogram revealed patterns characteristic of right heart strain. The spiral CT scan of the thorax was performed at Borromeus Hospital and this supported the diagnosis of PH with right ventricular and atrium dilatation.

Figure 1. Echocardiogram revealed pulmonary hypertension with severe tricuspid insufficiency.

Figure 2. Lung perfusion scintigraphy showed primary pulmonary hypertension with observed pulmonary embolism.
tricuspid regurgitation, and contrast regurgitation from the right atrium to the inferior cava vein and the hepatic vein. There was no sign of PE i.e. no filling defect in the subsegmental branches of the pulmonary arteries on both sides (Figure 3). Right and left lung examination revealed no disorder or abnormality explaining the cause of PH. Therefore, the warfarin was discontinued.

One month after these examinations he was brought to the hospital again with a complaint of shortness of breath that was accompanied by low grade fever and cough for one day. On examination he was alert but tachypneic (respiratory rate of 43/minute), with blood pressure of 110/70 mmHg, pulse rate of 107/minute with an irregular pulse (10 extrasystoles/minute), and a temperature of 37.9°C. The sclera of the eyes was slightly icteric and crackles in the lung base were detected. On auscultation, the first heart sound was normal, a loud second heart sound was found in the pulmonary area, and a systolic murmur of tricuspid regurgitation was heard. The liver was enlarged and a slight peripheral edema was noted. He was then diagnosed as having right heart failure (RHF) with PH, tricuspid insufficiency, cardiac arrhythmia, bronchopneumonia and cardiac liver cirrhosis. He was given oral sildenafil, captopril, clarithromycin, and furosemide. Repeated echocardiogram revealed PH with decreasing pulmonary artery pressure (pulmonary artery pressure 40 mmHg). A liver ultrasonogram showed cardiac liver cirrhosis.

Sildenafil and captopril were taken regularly. One month later, the patient still had jaundice and the echocardiography showed decreasing pulmonary artery pressure (pulmonary artery pressure 40 mmHg). So forth, he never turned-up in the clinic or hospital.

Discussion

Pulmonary embolism (PE) remains a common cause of mortality and its diagnosis is often missed. Although prompt diagnosis and appropriate therapy have been shown to reduce the mortality from 30% to less than 10%, the diagnosis of PE is often not suspected in children. Even though most of the signs and symptoms, including chest pain (70%), tachypnea (70%), cough (40%), tachycardia (33%), shortness of breath (25%), signs of deep venous thrombosis (10%) and syncope (5%), are likely to be seen in PE patients, diagnosis based on the clinical manifestations is not reliable. It has been noted that in adults clinical diagnosis has a sensitivity of 85% but a specificity of 38%, reflecting the vast differential diagnosis found in both adults and children. These clinical symptoms (chest pain, dyspnea, shortness of breath, syncope) also occur in PH.

The conventional chest x-ray is the commonest
investigation performed in a cardiac or respiratory emergency. In acute pulmonary thromboembolism it may be normal or non-specific.9 Chest radiographs and electrocardiograms are often normal in young patients with PE.7 The use of several imaging modalities such as ventilation-perfusion (V/Q) lung scanning, spiral volumetric computed tomography (CT) scanning, magnetic resonance imaging (MRI), CT angiography and digital subtraction angiography suggests that one single technique is not reliable to confirm or exclude the diagnosis in a patient with clinically suspected PE.2,4 V/Q scanning is the most frequently ordered diagnostic test in patients with clinically suspected PE.7 The obvious advantages of V/Q scanning are its low cost, ease of performance and its non-invasive nature. However, this type of scan can be problematic because it rarely provides a definitive “high probability” or “normal” diagnosis.4 In addition, this diagnostic test is not available in this hospital. Even though it was established in the PIOPED study (Prospective Investigation of Pulmonary Embolism Diagnosis) that a normal or low probability scan has a high negative predictive value and a high probability scan has a high positive predictive value when the results are interpreted along with the clinical assessment of the likelihood of PE, one study has shown that in approximately 70% of patients, the results of V/Q scanning were non-diagnostic.2 In patients with a non-diagnostic V/Q scan, pulmonary angiography is the next recommended imaging modality to establish the diagnosis of PE. However, pulmonary angiography is not often requested by clinicians because of its invasive nature.2,10 Pulmonary angiography is the historical gold standard for diagnosing PE, with which all other imaging tests have been compared.4,7,11,12 In our case study, lung perfusion scintigraphy performed on this patient revealed idiopathic or PPH with an observation of PE which was then treated with sildenafil and warfarin. As a confirmation of PE was needed to justify prescription of warfarin, a multidetector helical/spiral CT scan was performed. This technique has been proven to be useful to diagnose PE, and is available in another hospital (Borromeus Hospital) in the Bandung area. The choice between V/Q scanning and helical CT scanning tends to dominate most current discussions about non-invasive PE diagnosis.10 Recent studies have reported the usefulness of spiral CT scanning for the detection of central and segmental PE. It is non-invasive, easy to perform, quick and has been reported to have good specificity and sensitivity.2 Compared to V/Q scintigraphy or pulmonary angiography, the sensitivity of spiral CT has ranged from 53% to 100% and its specificity from 78% to 100%.7

Based on those findings, even though anticoagulation may be required in cases of PH that are associated with low cardiac output, leading to sluggish blood flow through the pulmonary artery which may predispose the patient to the development of pulmonary thrombi, we decided to stop warfarin treatment as PE was not diagnosed by multidetector spiral CT scanning. In the treatment of PH, oral sildenafil is given to promote an increase of cGMP levels and thus cause pulmonary vasodilatation.6 Although we had some limitations in performing all tests for the diagnosis of PH as proposed by the WHO, this patient likely had an idiopathic PH/PPH, as we could not determine other possible causes.6 Although PPH is a rare disease in childhood, and is associated with poor outcome and long term survival, the natural history of disease is heterogeneous with some patients dying within months of diagnosis and others living for decades.13 In present days, the treatment of PH is changing.3 Most cases of PH are difficult to treat and are irreversible unless the cause can be eliminated.5 Previously, the main treatment was aimed only to control congestive heart failure (CHF).3 The prognosis of PH is variable and represents a significant improvement compared to several years ago. Children who respond to short-term drug testing (responders) have a five year survival rate of 90%, while those who are nonresponders have a five year survival rate of only 33%. Many of the children in the later group may be candidates for lung transplants.14,15

During follow up, this patient developed jaundice, and decreased liver function, and ultrasonography examination suggested cardiac liver cirrhosis, leading to a diagnosis of decompensated RHF. Cardiac cirrhosis, also called congestive hepatopathy or chronic passive liver congestion, includes a spectrum of hepatic derangements that occur in the setting of RHF. The RHF causes transmission of elevated central venous pressures directly to the liver via the inferior cava and hepatic veins. At a cellular level, venous congestion impedes efficient drainage of sinusoidal...

Paediatr Indones, Vol. 49, No. 1, January 2009 • 57
blood flow into terminal hepatic venules. Sinusoidal stasis results in accumulation of deoxygenated blood, parenchymal atrophy, necrosis, perisinusoidal collagen deposition, and ultimately, fibrosis.\(^\text{16}\)

In conclusion, although the precise role of various non-invasive imaging techniques in the diagnosis of acute PE remains to be clarified, the result from using multidetector spiral CT as a diagnostic tool in the detection of PE is promising. It has high sensitivity and specificity, gives quick results and can help to provide alternative diagnosis, despite the fact that PE in this patient was unlikely to be confirmed based on multidetector spiral CT-scanning results.

References