Multidetector Computed Tomography Findings of Persistent Left Superior Vena Cava: A Report of Five Cases

Persistan Sol Vena Kava Süperiorun Çok Kesitli Bilgisayarlı Tomografi Bulguları: Beş Olgunun Sunumu

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Abstract

Persistent left superior vena cava is the most common variation of the anomalous venous return to the heart. It may coexist with the right superior vena cava but may also appear on its own. In addition, persistent left superior vena cava is usually asymptomatic and is discovered incidentally. When present, this condition could complicate intravascular procedures or cardiac surgery. Herein, is a report of the multidetector computed tomography findings of five cases with persistent left superior vena cava.

Key words: Multidetector computed tomography, persistent left superior vena cava, superior vena cava.

ÖZET


Anahtar Sözcükler: Çok kesitli bilgisayarlı tomografi, persistan sol vena kava süperior, vena kava süperior.

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Persistent left superior vena cava (PLSVC) is the most common variation of the anomalous venous return to the heart affecting about 0.3-0.5% of the general population (1,2). It can also be associated with 5-10% of congenital heart disease (3) and usually coexists with the right superior vena cava (RSVC), but this is not always the case (2,4,5). Persistent LSVC is usually asymptomatic and is discovered incidentally. In addition, this condition can present technical difficulties during intravascular procedures or cardiac surgery (1,5). Herein, we present the multidetector computed tomography (MDCT) findings of five cases with PLSVC.

CASE
Case 1: A 57-year-old male was admitted to the chest disease department of our facility with obstructive sleep apnea disorder. Physical examinations and laboratory results revealed no abnormalities, but posteroanterior chest radiography detected an enlarged mediastinum. Therefore, the patient underwent chest MDCT in the supine position, during maximum inspiration, using an eight-channel MDCT system (GE Healthcare, Milwaukee, WI, USA). Contiguous axial slices with contrast-enhanced CT were undertaken at 5 mm intervals with 5 mm collimation at automatically modulated amperage of 120 Kvp (120 to 225 mA). All images were obtained at window levels appropriate for the lung parenchyma (window width, 1.500-1.700 HU; window level, -600 or -700 HU) and mediastinum (window width, 250-400 HU; window level, 40-50 HU). The images were then reconstructed via a high-resolution algorithm, and the multiplanar reformatted (MPR) images were interpreted in various planes. The MDCT revealed cardiomegaly and aneurysmatic dilatation of the ascending aorta. Furthermore, PLSVC was observed at the left side of the mediastinum and the lateral side of the aortic arch, pulmonary arteries, and left atrium. Furthermore, they were draining into the right atrium via a dilated coronary sinus (Figure 1a and b). It was draining left jugular and subclavian veins and joining the left brachiocephalic vein (Figure 1c). Chronic fibrotic changes with traction bronchiectasis and peribronchial thickening were also observed.

Case 2: A 63-year-old male had been suffering from abdominal pain for two months prior to being admitted to our radiology department for abdominal ultrasonography (US). A clinical examination revealed no abnormalities, but laboratory tests showed elevated carbohydrate antigen 19-9 (CA19-9) (635.8 (0-35) U/mL) and carcinoembryonic antigen (CEA) (9.6 (0-5) ng/mL) levels. A pancreatic mass measuring about 2x2 cm in diameter and multiple metastases at the liver of varying diameters
were detected on the abdominal US. Diagnostic and staging procedures were then performed. The patient underwent chest MDCT with the same protocol used in Case I, and multiple metastatic nodules of various diameters at the lungs were detected. Moreover, the MDCT showed PLSVC at the left side of the mediastinum and on the lateral side of the aortic arch, pulmonary arteries, and left atrium, with drainage into the right atrium via the coronary sinus (Figure 2a and b) as well as the left jugular and subclavian veins. However, the researchers found no left brachiocephalic vein.

**Case 3:** A 64-year-old female with chronic renal insufficiency was admitted to our emergency department with dyspnea. The patient’s breathing and heart rate were rapid, and laboratory tests revealed a creatinine level of 8.4 (0.5-0.9) mg/dl, blood urea nitrogen level of 68.7 (6-23) mg/dl, and C reactive protein (CRP) level of 13.7 (0-5) mg/L. The patient’s shortness of breath was attributed to a probable pulmonary embolism, and the patient underwent MDCT angiography of the pulmonary artery in the supine position, during maximum inspiration, using an eight-channel MDCT system (GE Healthcare, Milwaukee, WI, USA). Contiguous axial slices with contrast-enhanced CT were obtained at 2.5 mm intervals, 0.875 mm slice thickness, and 105 Kvp, 305 mA. All images were obtained at window levels appropriate for the mediastinum (window width, 250-400 HU; window level, 40-50 HU). The images were reconstructed with high-resolution algorithm, and MPR images were interpreted in various planes. The MDCT angiography revealed PLSVC at the left side of the mediastinum and the lateral side of the aortic arch, pulmonary arteries, and left atrium, with drainage into the right atrium via a dilated coronary sinus. No right-sided superior vena cava (SVC) was detected, and we found no left brachiocephalic vein (Figure 3a and b). Additionally, no embolus was observed at the pulmonary arteries. However, the researchers did detect cardiomegaly, pleural effusion with compression atelectasis at both hemithoraces, and a mosaic perfusion pattern at the lungs.

**Figure 2a, b:** An axial MDCT image a) obtained at the level of the aortopulmonary window and a coronal MPR MDCT image b) reveal a left-sided SVC (✱) along with a right-sided SVC (white arrow).

**Figure 3a, b:** An axial MDCT image a) obtained at the level of the aortic arch and an oblique coronal MPR MDCT image b) show a left-sided SVC (✱) and a dilated coronary sinus. There was no right-sided SVC.
Case 4: A 77-year-old male was admitted to our emergency department with a three-day history of constipation and abdominal pain. The patient's medical history revealed chronic obstructive lung disease (COLD) and cardiac insufficiency. A physical examination was normal, and laboratory tests revealed an erythrocyte sedimentation rate (ESR) of 18 mm/h, and a CRP level of 50 mg/dL. For the detection of a probable malignancy, the patient underwent chest MDCT using the aforementioned protocol, which revealed PLSVC at the left side of the mediastinum and the lateral side of the aortic arch, pulmonary arteries, and left atrium, with drainage into the right atrium via a dilated coronary sinus. The researchers observed no left brachiocephalic vein (Figure 4a and b). Cardiomegaly, dilated pulmonary arteries, and pleural effusion at both hemithoraces were found.

Case 5: A 55-year-old female patient was admitted to our oncology department with abdominal pain and weight loss. A cholecystectomy was performed, and a pathological examination revealed poorly differentiated adenocarcinoma of the gall bladder with serosa and colonic invasion. Radiation therapy and systemic chemotherapy were administered; however, after completing the treatment, liver metastases developed. For re-staging, the patient underwent chest MDCT using the same protocol as the other cases, and PLSVC was discovered at the left side of the mediastinum and the lateral side of the aortic arch, pulmonary arteries, and left atrium, with drainage into the right atrium via a dilated coronary sinus. However, no left brachiocephalic vein was detected (Figure 5a and b). Furthermore, the MDCT revealed a dilated ascending aorta, a few nodules with ground glass opacity at the lungs, and dilated biliary tracts.

**DISCUSSION**

The anterior cardinal vein (ACV) and the posterior cardinal vein drain the cranial and caudal parts of the embryo, respectively. In addition, the left brachiocephalic vein develops and connects the cranial portions of the two ACVs in the first eight gestational weeks. The caudal part of the right ACV then becomes the normal RSVC, and the
left ACV caudal to the left brachiocephalic vein regresses and forms the oblique ligament and the vein of Marshall. If this portion fails to involute, it becomes a PLSVC (1). John Marshall published the first report of the great anterior veins of the thoracic region in humans and mammals in 1850, which including a description of PLSVC (6).

Persistent LSVC can occur in several anatomic variations. If bilateral SVC is present, left brachiocephalic veins may be completely absent in up to approximately 65% of the patients (6). In the current cases, no brachiocephalic veins were detected, with the exception of Case I. Persistent LSVC coexists with RSVC in up to 80-90% of cases, but isolated PLSVC is possible, as was seen in Case III. However, this is very rare (7).

In most of the cases with PLSVC, drainage into the right atrium via the coronary sinus occurs, resulting in no hemodynamic consequence. However, in approximately 10-20% of cases, drainage takes place via the left atrium, either through an unroofed coronary sinus or in a straight-line fashion into the roof of the left atrium or through the left superior pulmonary vein (6). In all of the current patients, drainage occurred in the right atrium via the coronary sinus.

Patients with PLSVC can have a variety of cardiac anomalies, such as atrial septal defect, a bicuspid aortic valve, coarctation of the aorta, coronary sinus ostial atresia, and cor triatriatum, and these issues are more commonly seen with the concomitant absence of the RSVC (8). The most frequently associated extracardiac anomaly is esophageal atresia (6), but in the present cases, no cardiac or extracardiac anomalies were detected.

Patients with PLSVC can present technical difficulties during intravascular procedures, for example Swan–Ganz catheterization, the insertion of pacing systems, cardiac catheterization, cardiac dialysis, or the cardiac surgery itself (4,5). Serious complications have also been reported when pacemaker leads or catheters have been inserted via the PLSVC (8). In addition, this condition can cause problems during central venous catheterization since accessing the coronary sinus can cause hypotension, angina, perforation of the heart, tamponade, and arrest. Furthermore, pacemaker implantation can also be difficult due to the circuitous path taken by the electrode, which can lead to the inability to obtain a stable electrode position and sustained capture. Moreover, in cardiopulmonary bypass, isolated PLSVC impairs the use of retrograde cardioplegia (7). Another problematic situation can occur during heart transplantation when the coronary sinus must be dissected carefully to permit re-anastomosis of the PLSVC to the right atrium (8). In the absence of an RSVC, central venous access should be made via the femoral vein in patients with PLSVC. During right-sided open-heart surgical procedures, drainage should be done by inserting a separate cannula. If the PLSVC drains into the left atrium and creates a large right-to-left shunt, surgical correction should be undertaken by central venous access via the femoral vein (7). In addition, PLSVCs may predispose the heart to arrhythmia owing to the close proximity of the dilated coronary sinus to the final position of the left-sided primitive pacemaking tissue (5).

When PLSVCs are present, electrocardiography often shows an abnormal P-wave axis and a normal or shortened PR interval. In addition, on chest X-ray, a crescent-shaped shadow of the PLSVC can be seen at the aortic knob or left upper mediastinum. After the insertion of a pulmonary artery catheter into the left subclavian or jugular vein, a control chest X-ray can give the false appearance that the catheter has passed through the vessel (7). However, transthoracic echocardiography can be used to reveal the dilated coronary sinus and the diagnosis can be confirmed with the use of a saline contrast (“bubble study”) echocardiography. Single or multiplane transesophageal echocardiography and radionuclide angiocardiography have also been used to establish a diagnosis. Multidetector CT or magnetic resonance venography can also be employed to diagnose this condition and are useful for ruling out variations in the typical anomalous venous course (8). In addition to the axial images, MPR images are also important for the detection of PLSVCs and the evaluation of the presence of an RSVC, left brachiocephalic vein, and other vascular and cardiac anomalies. Furthermore, MPR allows for the evaluation of the chest in more than one plane to show the drainage of the PLSVC into the heart.

In conclusion, radiologists and clinicians should be aware of PLSVC and its variations in order to avoid possible complications and should use MDCT to aid in the diagnosis of this condition. Therefore, it is beneficial to be aware of the presence of PLSVC via MDCT before invasive procedures are performed.

CONFLICTS OF INTEREST
None declared.

AUTHOR CONTRIBUTIONS

YAZAR KATKILARI

REFERENCES