Radiology of Anomalies of Pulmonary Veins

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Keywords

► congenital diseases
► inferior vena cava
► vertical vein

Abstract

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Settings and Design

Selection of cases—the adolescents, infants, and children, of both sexes, from day 1 to 15 years and who had clinical signs of CVS and positive 2D echo findings were selected. The dose of contrast and radiation dose were kept as per the recommendations. MDCT was performed on Dual Source CT (Siemens Co Ltd.) which has a high isotropic resolution with 0.6 mm resulting into high quality orthogonal images.1,2

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Technique of MDCT Angiography

1. Age group: day 1 to 15 years.
2. Inclusion criteria:
   Patients presenting to the hospital with congenital heart disease referred for cardiac MDCT were included.
3. Exclusion criteria:
   a. Patients coming for postoperative follow-up or complications.
   b. Patients coming with recurrent disease.
   c. Contrast enhanced CT was performed using (SIEMENS SOMATOM 64 dual source) MDCT scanner, under light sedation if required.
d. Sedation—chloral hydrate or IV sedation or general anesthesia (GA)—if SOS. Early cases were done without GA but in some cases, study had to be repeated. Hence GA was given in infants and co-operative children.
e. Injection of contrast—manual in neonates, 20 to 24 size needle, pressure injector in older children (1.5–4 mL/s). It was used to dilute contrast, so that a greater volume of 3 mL/kg could be used at a volume of 2 mL/kg.
f. Pitch of 1.2, 0.75, or 1-mm slice thickness, 64 × 0.625 mm collimation, gantry rotation 0.33 seconds.
g. Field of view (FOV) for cardiac anomalies—from the level of sternum notch up to diaphragm.
h. If major aortopulmonary collateral arteries (MAPCA) are suspected—from neck to inferior vena cava (IVC).
i. Low radiation—80 to 120 mA, 80 to 120 KV.
j. Scanning: (a) at the end for right heart and (b) bi-phasic for Lt heart: with bolus tracking.
k. Multiplanar reconstruction (MPR), maximum intensity projection (MIP) (with varying slice thickness), axial, coronal, sagittal, or oblique recons, rotate images, volume rendering technique (VRT) images were used.
l. In some cases, conventional pulmonary angiography was performed.

Discussion
The anomalies of pulmonary veins can be grouped as follows:

Type I: Supracardiac
Anomalous pulmonary veins terminate at the supracardiac level. Pulmonary veins converge to form a left vertical vein, which drains to either brachiocephalic vein, superior vena cava (SVC), or azygos vein. The supracardiac variant can classically depict a snowman appearance (figure of eight heart or cottage loaf heart) on a frontal chest radiograph. In this child MDCT showed the vertical vein even when Chest X-ray was normal (► Figs. 1 and 2). This is a left to right shunt. This can be TAPVR (total anomalous pulmonary venous return) or PAPVR (partial anomalous pulmonary venous return).

Type II: Cardiac
Pulmonary veins drain into the coronary sinus and then the right atrium or SVC (► Figs. 3, 4, and 5). A specific variant is a meandering pulmonary vein (► Fig. 6). It is an anomalous pulmonary vein, taking a circuitous route through the lung to enter into (in contrast to scimitar syndrome) the left atrium, rather than the IVC. Meandering pulmonary veins can occur on the left side or both sides and coincide with features of scimitar syndrome. This is a left to left shunt.

Fig. 1 A female neonate, presented with respiratory distress and cyanosis soon after birth. MPR image of MDCT shows supracardiac type of total anomalous return of pulmonary veins (white arrow). MDCT, multidetector computed tomography.

Fig. 2 In this child CXR was normal. CT showed supra-cardiac type of TAPVR with all pulmonary veins on right draining into left SVC (vertical vein, marked by a white arrow), then into right brachiocephalic vein and normal SVC. TAPVR, total anomalous pulmonary venous return.

Fig. 3 A 7-year-old female with ASD. MDCT shows a small tributary of RIPV opening into SVC—cardiac type (marked by arrows). MDCT, multidetector computed tomography.
Type III: Infracardiac
TAPVR, in which all four pulmonary veins drain abnormally to the right atrium instead of the left atrium. This becomes a left to left shunt.

PAPVR is a congenital malformation in which pulmonary vein enters the systemic veins, e.g., in Scimitar syndrome. This is a left to right shunt. All pulmonary veins may drain into hepatic veins, portal vein or IVC. An inferior pulmonary vein may have a meandering course before ending in right atrium, IVC, or even portal vein (pulmonary veno-arterial or systemic venous shunt).

Rarely the pulmonary veins may join behind the left atrium to form a common vertical descending vein, coursing anterior to the esophagus and passing through the diaphragm at the esophageal hiatus and then joining the portal system.

There have been a few studies in the literature describing the technique of CT of heart in pediatric patients (Figs. 5, 7).

Type IV—miscellaneous anomalies—pulmonary AVM and pulmonary varix (Figs. 6, 8, and 9).

Summary
A study of 43 suspected congenital diseases of heart supported by 2D echo was performed in children and adults. Only anomalies of pulmonary veins are
presented. These were better accepted by clinicians. In most cases it tallied with 2D echo but in some cases it did not, as there can be difference in interpretation between the two cardiologists.

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None.

Conflict of Interest
None declared.

Acknowledgment
None.

References

Fig. 8  CXR in this adolescent showed a round density to the right and behind right hilum (arrow). A nodal mass or a vascular lesion was suspected. CT showed this to be a pulmonary venous angioma (arrowhead). CT, computed tomography.

Fig. 9  PA view of CXR shows bilateral anomalous curvilinear vessels in the lower pulmonary regions. MDCT with axial and coronal MIPs demonstrates bilateral pulmonary veins with anomalous routes in the lower pulmonary regions but draining normally into the left atrium. Coronal volume-rendering 3D reconstruction shows the anomalous veins and normal pulmonary arteries. PA, posteroanterior; 3D, three dimensional; MDCT, multidetector computed tomography.