A female child with visceral heterotaxy (presumed left isomerism), interrupted inferior vena cava with azygos continuation to the superior vena cava, and double-outlet right ventricle underwent a total cavopulmonary anastomosis (end-to-side anastomosis of the superior vena cava to the right pulmonary artery) in infancy. Completion of Fontan circulation was undertaken at 3 years of age with construction of a tunnel from the hepatic veins to the pulmonary artery confluence.

By 6 years of age, the child had become increasingly cyanotic, and cardiac catheterization was performed to define the mechanism of the cyanosis. Figure 1 is a pulmonary artery angiogram that reveals moderate hypoplasia of the pulmonary artery confluence. Figure 2 is a selective angiogram performed in the hepatic vein tunnel, with hepatic venous flow directed predominantly toward the left lung. The left pulmonary artery angiogram was normal. A right pulmonary artery angiogram (Figure 3) was markedly abnormal, showing a diffusely granular appearance of the distal vasculature and early appearance of contrast in the right pulmonary veins, characteristic of the presence of pulmonary arteriovenous malformations. The development of arteriovenous malformations in the right lung allowed for intrapulmonary right-to-left shunting, accounting for the patient’s progressive cyanosis. Figure 4 is an azygos vein angiogram, performed after the

**Figure 1.** Pulmonary artery angiogram, anteroposterior view. Catheter course is from azygos vein to superior vena cava to right pulmonary artery (RPA) via cavopulmonary anastomosis. White arrowhead denotes hypoplastic segment between right pulmonary artery and left pulmonary artery (LPA).

**Figure 2.** Selective angiogram in hepatic vein tunnel (HVT) outlining hepatic veins (HV). Note distribution of hepatic venous flow toward left pulmonary artery (LPA). No evidence of pulmonary arteriovenous malformations was found in the left lung.
implantation of an intravascular stent into the hypoplastic pulmonary artery confluence. Note the presence of a duplicated inferior vena cava (the left inferior vena cava received the left renal vein).

This case illustrates the importance of hepatic venous return circulating through the pulmonary vasculature in inhibiting pulmonary angiogenesis. The “antiangiogenic” properties of hepatic venous blood are attributed to an as-yet uncharacterized hepatic factor. The left lung, which received hepatic venous return through the surgically created hepatic vein tunnel, was normal. However, the right lung, which was underperfused with hepatic venous effluent both by virtue of the geometry of the hepatic vein to pulmonary artery tunnel and the narrowing of the pulmonary artery confluence, developed arteriovenous malformations.
Development of Unilateral Pulmonary Arteriovenous Malformations due to Unequal Distribution of Hepatic Venous Flow
Henri Justino, Lee N. Benson and Robert M. Freedom

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