Pulmonary Atresia or Persistent Truncus Arteriosus
Is It Important to Make the Distinction and How Do We Do It?
Margaret L. Kirby

The congenital cardiac anomaly known as tetralogy of Fallot (TOF) is characterized by right ventricular outflow tract obstruction caused by subpulmonary stenosis, dextroposition (overriding) of the aorta, a ventricular septal defect, and right ventricular hypertrophy. The right ventricular hypertrophy is secondary to the presence of right ventricular outflow obstruction (pulmonic valvar or subvalvar stenosis or in the most severe case, atresia). Cyanosis in these patients is attributable to the passage of systemic venous blood into the aorta, bypassing the lungs, with the degree of cyanosis dependent on the severity of the outflow tract obstruction. The malformations that are classified as persistent truncus arteriosus (PTA) are characterized by a single multicuspid semilunar valve with a single vessel, the truncus, arising from the ventricles and giving rise to systemic, pulmonary, and coronary circulations. Thus, both PTA and TOF with pulmonary atresia are characterized by a single vessel emanating from the heart. In PTA, the septation that would divide the common arterial trunk into an aorta and pulmonary trunk is missing, whereas in TOF with pulmonary atresia, it is unclear whether this septation is missing or misplaced. Diagnosis of TOF with pulmonary atresia relies on the presence of a pulmonary valve remnant by clinical imaging. This criterion permits differentiation of TOF with pulmonary atresia and PTA. However, if the pulmonary atresia develops embryonically before a valve is formed, then this criterion would not distinguish PTA from TOF with pulmonary atresia. New findings in experimental mouse models by Théveniau-Ruiss et al reported recently in Circulation Research shed unexpected light on the embryogenesis of these defects and may allow differentiation of the 2 defects in ways that have not been used previously.1

The myocardium and smooth muscle at the arterial pole, that is, the subaortic and subpulmonary myocardium and the smooth muscle at the base of the aortic and pulmonary arterial trunks, is added from a specific region of the ventral pharyngeal mesoderm caudal to the attachment point of the outflow tract with the pharynx that has been called secondary heart field.2,3 The secondary heart field contains the progenitors of both the myocardium and smooth muscle of the arterial pole and ablation of this region has been shown to result in pulmonary stenosis/atresia with overriding aorta.4 This implicates secondary heart field as the defining cell population associated with pulmonary stenosis/atresia. Other experimental causes of pulmonary stenosis/atresia are associated with decreased proliferation in this field of progenitors. Decreased proliferation is experimentally associated with decreased fibroblast growth factor signaling.5 Tbx1-null mice also have decreased proliferation in the secondary heart field, and this is associated with TOF.5,6

Reports from Margaret Buckingham and colleagues in the last 3 years have shown that myocardium originating from this region has unique aortic and pulmonary identity well before septation of the outflow tract. This group has identified 2 separate loci that specifically express a lacZ reporter in outflow myocardium in what will become the subaortic and subpulmonary myocardium. The first report used the y96-Myf5-nlacZ-16 (96-16) transgene that marks the pulmonary outflow myocardium in a pattern complementary to 96-16 and continues development as the subaortic myocardium. The 2 myocardial populations represented by the 96-16 (subpulmonary) and T55 (subaortic) appear from a clonal analysis to arise from distinct precursor populations.8

The article by Théveniau-Ruiss et al identifies the gene that is represented by the 96-16 transgenic as Semaphorin3C (Sema3C).1 Sema3C, a secreted growth factor associated with axonal path-finding and angiogenesis, has been reported previously to be expressed in the outflow myocardium, although not specifically identified as subpulmonary myocardium. Sema3C mutant mice have cardiovascular defects that include conotruncal malformations, although these are, unfortunately, not well described.9 Plexins are the receptors for semaphorins, and PlexinA2 is expressed by neural crest cells. In Sema3C-null embryos, distribution of the plexinA2-expressing neural crest cells in the outflow tract is altered but the cells are present in the outflow tract.9 Furthermore, cardiac neural crest form the outflow septation complex...
valves with biventricular origin of the arterial trunk is intriguing. In virtually all the animal models of PTA, the arterial trunk arises from the right ventricle.

To summarize these comments, the diagnosis of PTA and pulmonary atresia in patients has used criteria that may not have recognized variants of pulmonary atresia beyond an atretic pulmonary valve. If the subpulmonary myocardium never grows into the heart, as in the case shown by Theveniau-Ruissy et al, the outflow tract and right ventricle would lack this tissue from very early and not have a chance to form a pulmonary outlet. Because septation of the arterial pole requires an entirely different developmental process involving neural crest cells which appear to be guided by the junction of the subpulmonary and subaortic myocardium, it is possible that septation is present, albeit in the wrong place because of the reduction in the subpulmonary myocardium. This implies an altogether different embryogenesis of PTA (a septation problem) and pulmonary atresia (a problem with development of the subpulmonary myocardium). Even though the clinical treatment of the 2 types of patients is the same, to ultimately understand the genetic and epigenetic causes of the 2 defects requires that the criteria for diagnosis be reevaluated.

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