In the first part of our review of cardiac development, we discussed the initial changes involved in transformation of the heart forming regions of the embryo into the great veins, the atrial and ventricular chambers, and the arterial trunks. We showed that this first part of cardiac development could be divided into phases of formation of the primary myocardial tube, looping of the tube, during which additional parts are added that give the future compartments their definitive topography, and the assembly of these components into the cardiac chambers and arterial trunks. In this second review, we discuss the processes that complete the separation of the two sides of the definitive heart, for the most part involving septation of the parts of the primary tube not themselves directly involved in formation of the chamber-specific compartments of the atriums and ventricles. In this respect, when concluding our first review, we explained how the arterial trunks were also formed by septation within the solitary outflow tract of the primary heart tube. We also showed, however, that subsequent to formation of the two arterial trunks, there was disappearance of the cushions that initially divided them. Thus, in the definitive heart, the proximal parts of the aorta and pulmonary trunk, along with the sinuses of the arterial roots and the subpulmonary infundibulum, possess their own discrete walls, separated by extra-cardiac space. Although septation by fusion of endocardial cushions will be a topic included in this second review, septation and separation of the outflow tracts is sufficiently complicated to require special treatment. Because of this, we will devote a third review specifically to the mechanisms underscoring the remodelling of the outflow tracts. In this review, therefore, we will confine our considerations to the formation of the atrial, atrioventricular, and ventricular septal structures.

MORPHOLOGICAL CONSIDERATIONS

As will become evident, separation of the four cardiac chambers involves more than simple formation of partitions between them. There is justification for arguing that understanding the complex mechanics of development itself will provide the information required properly to interpret the postnatal morphology. Clinicians, however, are more likely to be acquainted with the structure of the heart rather than its development. We will preface our account of the developmental events, therefore, with a brief review of the nature of the tissues that separate the cavities on the right and left sides in the definitive heart. Appreciation of the precise structure of these separating walls underpins the understanding of the developmental steps involved in their formation.

Thus, careful examination of the heart subsequent to separation of the chambers shows that several of the structures usually described as “septums” in classical accounts are, in reality, no more than folds, with extracardiac adipose tissue incorporated between the folded layers of myocardium. It can reasonably be argued, of course, that any partition interposed between the cavities of adjacent cardiac chambers is justifiably described as a septum, irrespective of its precise structure. Our own understanding was helped, nonetheless, when we came to appreciate the anatomic difference between shelves within the chambers, as opposed to those folded myocardial walls that incorporate between them wedges of extracardiac fibro-adipose tissue. This difference is best seen at the oval fossa (“fossa ovalis”). The flap valve, forming the floor of the fossa, and being free to open and close while the fossa is patent, is the exemplar of the shelf-like partition that we define as a true septal structure. The antero-inferior muscular buttress that anchors the flap valve to the fibrous atrioventricular septal structures is also a true septum, since it, too, can be removed without creating a communication with the extracardiac space. In contrast, the so-called “septum secundum”, seen supero-posteriorly, is a deep infolding between the walls of the superior caval and right pulmonary veins at their junctions with their respective atriums (fig 1). Incisions in this area, initially extending into the subpapillary fibro-adipose tissue, can potentially extend to the pericardial cavity.

Yet another arrangement of overlapping tissues is to be found in the area of the triangle of Koch. In this area, which separates the off-set attachments of the tricuspid and mitral valvar leaflets, a layer of fibro-adipose tissue is again incorporated between two muscular layers, but in this example, the muscular layers—one atrial and the other ventricular—are both attached antero-superiorly.

See end of article for authors’ affiliations

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to the central fibrous body. The fibro-adipose layer between the two muscular walls is itself continuous with the inferior atrioventricular groove (fig 2). Although previously we nominated this area as a “muscular atrioventricular septum”, septation itself cannot produce this arrangement. Rather, as we will describe, the folding occurs concomitant with caudal expansion of the right atrioventricular junction over the base of the ventricular mass.

The significance of morphologic distinctions of this type should become evident as we discuss the development of the various structures that, eventually, separate the atrial and ventricular chambers within the definitive heart. These are the atrial septum, the atrioventricular septum, and the ventricular septum. We will start, however, with a brief consideration of another area often described as a septum, again erroneously in our view, namely the tissues that separate the two horns of the systemic venous sinus (“sinus venosus”).

FORMATION OF THE “SEPTUM” OF THE SYSTEMIC VENOUS SINUS
As described in our first review, the systemic venous tributaries exist from the outset of formation of the circulatory system, being continuous with the caudal, or atrial, end of the developing heart tube. These systemic venous tributaries are initially formed in symmetrical fashion in the right and left sides of the embryo. In the earliest human embryo that we have studied, nonetheless, at Carnegie stage 12, the veins are already joined to the primary atrial component of the heart tube in asymmetrical fashion (fig 3). This is because, already at this early stage of development in the human, the left sinus horn is becoming incorporated into the developing left atrioventricular groove, passing beneath the posterior mesocardium. The mesocardium itself provides the portal for the pulmonary vein, when it develops, to enter the left atrium.

An integral part of ongoing atrial development is reorientation of the openings of the systemic venous tributaries so that they empty only to the right side of the primary atrium, this being a necessary prelude to atrial septation. Almost always, in the human, there is an accompanying refiguration of the venous pathways themselves. Anastomoses are formed in the neck and abdomen, permitting a greater volume of blood to be returned to the heart through the right sided tributaries, which persist at their junction with the atrium as the orifices of the superior and inferior caval veins. There is concomitant diminution in size of the left sinus horn, which persists as the coronary sinus, developing so as to receive most of the venous return from the heart itself. The left sinus horn is also smaller than the right horn in the mouse heart (fig 4A). In the mouse, however, the left sinus horn remains patent in postnatal life, draining the persisting left superior caval vein. Such a
situation can also be found as a congenital malformation in the human, with the left caval vein maintaining both its patency and its connection to the right atrium. Reduction in size of the left horn, therefore, is not a pre-requisite for successful atrial septation. Be that as it may, it is the initial bifurcation of the walls of the right and left horns, at their entrances to the systemic venous sinus, that persists in the human and the mouse as the so-called “sinus septum” (fig 4B). This so-called septum, therefore, is no more than a muscular fold in the atrial walls of the right atrium, interposing between the orifices of the inferior caval vein and the coronary sinus.

FORMATION OF THE ATRIAL SEPTUM

As we have described, the systemic venous tributaries are re-oriented as a prelude to septation so as to enter exclusively the right side of the primary atrial component of the heart tube. And, concomitant with this reorientation of the systemic venous tributaries, the pulmonary vein develops as a new structure, using the persisting posterior mesocardium to gain its entrance to the left side of the primary atrium. Reduction in size of the left horn, therefore, is not a pre-requisite for successful atrial septation. Be that as it may, it is the initial bifurcation of the walls of the right and left horns, at their entrances to the systemic venous sinus, that persists in the human and the mouse as the so-called “sinus septum” (fig 4B). This so-called septum, therefore, is no more than a muscular fold in the medial wall of the right atrium, interposing between the orifices of the inferior caval vein and the coronary sinus.

Septation is accomplished in several steps. Two endocardial cushions are produced within the initially common atrioventricular junction, also part of the primary heart tube (figs 5 and 6), while the primary atrial septum grows as a muscular partition from the roof of the primary atrial component of the heart tube (figs 6 and 7). This muscular shelf grows into the atrial cavity between the systemic and pulmonary venous openings (fig 8), taking its origin from the part of the primary atrium that has molecularly left characteristics, as shown by its expression of the Pitx2 gene. The part of the primary atrium with these left sided molecular characteristics extends as far rightwards as the attachments of the left valve of the systemic venous sinus. As the primary muscular septum grows from the atrial roof, with its origin leftward relative to the developing left venous valve, it carries a mesenchymal cap on its leading edge (fig 9). With continuing growth of the muscular primary atrial septum, this mesenchymal cap approaches, and then fuses with, the superior endocardial cushion of the atrioventricular canal (fig 10). Concomitant with this development, a further mass of mesenchyme (fig 11) grows into the heart inferiorly from the posterior mediastinum. At this stage, the pulmonary vein remains a solitary channel, which opens to the left atrium inferiorly, adjacent to the atrioventricular junction. The mass of mesenchyme entering the heart from the mediastinum, called the “spina vestibuli” by His, grows through the rightward border of the new pulmonary venous orifice, enlarging the border in the process (fig 8). As it grows into the atrium, this vestibular spine also carries forward the inferior ends of the so-called venous valves, which by now guard the orifices of the systemic venous sinus, marking their junction with the developing right atrium (fig 9). The process of merging and fusion of the mesenchymal cap on the
primary septum with the mesenchymal tissues of the vestibular spine, combined with subsequent fusion with the superior and inferior atrioventricular cushions, themselves also fused by this stage, serves to fill in the gap initially seen beneath the leading edge of the primary septum and the atrioventricular endocardial cushions. In other words, it is the fusion of the various mesenchymal components that closes the primary atrial foramen, or “ostium primum”. At the same time, the process of fusion divides the common atrioventricular junction into its right and left components. Before the primary atrial foramen is closed, however, the upper margin of the muscular primary septum breaks down to form the secondary interatrial foramen, or the “ostium secundum” (figs 6 and 7). After fusion of the various mesenchymal structures, the primary septum itself is firmly adherent, via the tissues derived from the vestibular spine and the mesenchymal structures, to the crest of the ventricular septum. This permits the primary septum itself to function as the flap valve of the oval foramen. The mass of tissue derived from the vestibular spine now forms an extensive buttress at the right side of the base of the primary septum. This tissue then muscularises, except for the central part continuous posteriorly and inferiorly with the valves of the systemic venous sinus. This remains as the fibrous tendon of Todaro. The muscularised component forms a true secondary atrial septum at the base of the flap valve, the latter, of course, representing the persisting muscular primary atrial septum (fig 12).

When first formed, the secondary foramen has no upper rim, its border being the flat atrial roof (fig 10). Only much later, subsequent to separation of the right and left pulmonary veins, and incorporation of their orifices as separate openings to the roof of the left atrium, does the deep fold develop to form the structure usually described as the “septum secundum”. It is against this fold that the flap valve abuts in postnatal life to close the oval foramen (fig 12).

**FORMATION OF THE ATRIOVENTRICULAR SEPTUM**

As explained above, before growth of the primary atrial septum from the roof of the atrium, two masses of mesenchymal tissue had formed superiorly and inferiorly within the atrioventricular canal. These are the atrioventricular endocardial cushions (figs 5 and 6). The two cushions grow towards each other, fusing to divide the canal into its right and left components (fig 10). The cushions are also adherent to the crest of the muscular ventricular septum, this septum itself forming as the apical trabecular components of the developing left and right ventricles balloon from the primary heart tube (see below). A key feature in growth of this primary ventricular septum, however, is the increase in size of its posteroinferior component. This part of the muscular septum develops concomitantly with growth of the inlet of the right
ventricle, this self evidently occurring subsequent to division of the atrioventricular canal. Having divided, the atrioventricular canal has been dissected away from the atrial cavities, permitting the atrial cavities to be photographed from beneath. The valves of the systemic venous sinus have now formed, showing that the sinus itself (SVS) opens exclusively to the right side of the primary atrium. Note that the orifice of the pulmonary vein now opens inferiorly to the left side of the atrium. The primary atrial septum is growing as a shelf between the systemic and pulmonary venous orifices. Note also the bulge at the right side of the pulmonary venous orifice, between it and the opening of the systemic venous sinus. This is the "spina vestibuli" initially described by His, which we term the vestibular spine.

Figure 8  This scanning electron micrograph is from a mouse heart at a slightly earlier stage of development than the human embryo shown in fig 6. The atrioventricular canal has been dissected away from the atrial chambers, permitting the atrial cavities to be photographed from beneath. The valves of the systemic venous sinus have now formed, showing that the sinus itself (SVS) opens exclusively to the right side of the primary atrium. Note that the orifice of the pulmonary vein now opens inferiorly to the left side of the atrium. The primary atrial septum is growing as a shelf between the systemic and pulmonary venous orifices. Note also the bulge at the right side of the pulmonary venous orifice, between it and the opening of the systemic venous sinus. This is the "spina vestibuli" initially described by His, which we term the vestibular spine.

Figure 9  This higher power scanning electron micrograph shows the atrial roof, prepared in comparable fashion to fig 8, but at a slightly later stage of development. The upper margin of the primary atrial foramen has now broken down to form the secondary interatrial foramen (double headed arrow). Note the mesenchymal cap on the leading edge of the primary septum, which is growing towards the atrioventricular endocardial cushions, carrying forward the inferior end of the orifice of the systemic venous sinus (star). The upper end of the orifice becomes the septum spurium in the roof of the right atrial appendage.

Figure 10  This section, in four chamber plane, is from a human embryo at stage 16 of the Carnegie classification. It shows how the mesenchymal cap on the primary atrial septum has now fused with the superior cushion, with breakdown of the upper edge to form the secondary foramen. The upper margin of the foramen, however, is flat to the orifice of the right sinus horn (RSH, black line). Only subsequent to septation is there infolding of the superior margin (see fig 12).

Figure 11  This section, in four chamber plane, is from the same human embryo at stage 16 of the Carnegie classification as shown in fig 10. This section shows the superior cushion adherent to the ventricular septum, but with the larger part of its bulk within the left ventricle. It also shows the tissue growing into the heart through the vestibular spine, confining the pulmonary vein within the developing left atrium. Note the left sinus horn (LSH) with its own discrete walls.

ventricle, this self evidently occurring subsequent to division of the atrioventricular canal. Having divided, the atrioventricular canal expands rightward and, as it does so, the larger part of the inferior atrioventricular cushion becomes draped over the right side of the enlarging posteroinferior part of the muscular ventricular septum. The larger part of the superior cushion, in contrast, together with a smaller part of the inferior cushion at the site of fusion, remains within the cavity of the developing left ventricle. The line of fusion with the inferior cushion then forms the supero-posterior margin of the embryonic interventricular foramen. This part of the foramen has now become an arch-like structure that, eventually, will provide the vestibule of the subaortic part of the left ventricular outflow tract (fig 13).

The tricuspid orifice itself has increased in size concomitant with the expansion of the atrioventricular canal, and growth of the inlet part of the right ventricle. The mitral orifice also increases in size. It is only at a much later stage, after the completion of ventricular septation (see below), that the musculature of the initial atrioventricular canal that surrounded the developing valvar orifices, apart from the area destined to become the atrioventricular conduction axis, loses its continuity with the ventricular myocardium. This separation is produced by formation of rings of fibro-adipose tissue throughout the newly expanded atrioventricular junctions. This process of separation, which produces the essential plane of electrical atrioventricular insulation, occurs at a level that sequestrates the musculature of the initial atrioventricular...
Canal on the atrial side of the new junctions, thus forming the vestibules of the two atria.10

The atrioventricular cushions themselves occupy the newly formed septal component of the junctions. As explained above, the greater part of the superior cushion, along with a smaller part of the inferior cushion, are located within the left ventricle, where they form the basis of the atrioventricular septum. The ventricular aspect of this leaflet will eventually form the posterior part of the vestibule of the sub-aortic outflow tract (fig 13). The fused part of the cushions attached to the muscular ventricular septum itself, with the muscularising vestibular spine adherent to its atrial aspect, now forms the atrioventricular septal component of the so-called central fibrous body. The other part of this body is the rightward end of the arch of cushion tissue now occupying the roof of the left ventricle, which will become the right fibrous trigone. The central fibrous body, therefore, represents the fused remnants of the atrioventricular endocardial cushions. This fibrous tissue now occupies the space that, initially, was the central part of the atrioventricular canal. The musculature found immediately beneath and behind the fibrous tissue, therefore, is the musculature of the inferior quadrant of the atrioventricular canal, which is primary myocardium.11 This part of the canal musculature retains its continuity with both the atrial and ventricular muscular masses, becoming the atrioventricular node and the penetrating atrioventricular bundle.11

At the time that the cushions themselves initially fused to divide the right from the left atrioventricular junctions, this muscular pathway, being the wall of the atrioventricular canal, marked the postero-inferior extent of the cardiac cavities (fig 14). Subsequent to the sequestration of the right side of the initial musculature of the atrioventricular canal into the right atrium, the atrial end of the primary myocardium has become the atrioventricular node (fig 15), positioned in the definitive heart at the apex of the triangle of Koch. The atrial musculature forming the floor of the triangle of Koch has, by now, come to overlie the right ventricular septal musculature that was excavated concomitant with formation of the right ventricular inlet. It is the overlap of the muscular walls in this area that, initially, we described as representing a “muscular atrioventricular septum”.11 In reality, it is the process of expansion and separation of the atrial and ventricular muscle masses, rather than septation itself, that incorporates the wedge of extracardiac fibro-adipose tissue between the atrial and ventricular walls.

On the ventricular side, it is only subsequent to the closure of the interventricular foramen (see below) that delamination occurs within the myocardium to form the septal leaflet and inferior leaflets of the tricuspid valve, along with their supporting tension apparatus.11 Also only subsequent to delamination of these leaflets11 is it possible to recognise the ventricular component of the membranous septum (fig 16). This...
fibrous part of the septum is derived from the most proximal parts of the outflow cushions. It is the fusion of this ventricular component of the membranous septum with its atrioventricular component, the latter derived from the fused atrioventricular cushions, that closes the embryonic interven- 

tricular foramen, thus walling the aorta into the left ventricle (fig 17).

**FORMATION OF THE VENTRICULAR SEPTUM**

As explained in our first review, the muscular part of the ven-

tricular septum is produced concomitantly with the ballooning of the apical components of the right and left ventricles from the primary heart tube. As these “pouches” grow out from the heart tube, so the muscular septum is formed between them (fig 18). Until recently, there was disagreement concerning the developmental structures that contributed to the definitive muscular septum. For quite some time, it was believed that the inlet part of the septum formed separately from the primary apical muscular septum. Indeed, one of us (RHA), in early attempts to explain the development of the heart, illustrated the inlet part of the ventricular septum as a separate structure. Along these lines, others have argued that there is a muscular septum of the atrioventricular canal incorporated into the definitive ventricular septum. When adjudicating this process, it should be noted that there are pronounced differences between birds and mammals in the formation of the ventricular septum. In the chick, it is known that the primary apical muscular septum itself compacts by fusion of a series of layered trabeculations. Also in the chick, it is known that muscularisation of the atrioventricular cushions does, indeed, produce a muscular septum of the atrioventricular canal.

Thus far, no one has shown that similar mechanisms occur in the mammalian heart.

For the mammalian heart, it was the serendipitous discovery that an antibody to the nodose ganglion of the chick
marked the site of the developing atrioventricular conduction tissues in man that resolved the disputes concerning the structures contributing to the mammalian muscular ventricular septum. These studies showed unequivocally that the primary muscular ventricular septum produced by ballooning from the primary heart tube separated not only the apical components of the ventricles, but also interposed between the entirety of the inlet of the right ventricle and the cavity of the left ventricle.

Thus, as the apical parts of the ventricles balloon from the primary heart tube, the antibody to the nodose ganglion delimits a band of tissue that surrounds the primary interventricular foramen (fig 19). Inferiorly, the tissue lies within the musculature of the atrioventricular canal, marking the site of the subsequent atrioventricular conduction axis. Significantly, the stained tissue, which becomes the conducting tissues, is not only inter输给ventricular and atrioventricular. The section is taken between the unfused atrioventricular cushions, showing that most of the circumference of the atrioventricular canal is supported by the developing left ventricle (black bracket).

ventricular component of the ring, which forms the atrioventricular bundle, retains its position on the crest of the newly formed ventricular septum. As has already been shown (figs 14 and 15), at the site of its continuity with the atrioventricular node the penetrating bundle is sandwiched between the muscular ventricular septum and the atrioventricular component of the newly formed membranous septum, this now being part of the central fibrous body (fig 13). This anatomic arrangement would not be possible if a second muscular component was incorporated into the definitive ventricular septum, be it derived from a separate muscular “inlet septum” or a “septum of the atrioventricular canal”. As also explained, such a septum is found within the chick heart, produced by muscularisation of the atrioventricular endocardial cushions.

In the chick heart, nonetheless, the atrioventricular bundle is buried within the muscular ventricular septum, sandwiched between the primary ventricular septum and the part of the septum derived from the muscularised cushion. This arrangement is not seen in the human heart (fig 15). Subsequent to formation of the inlet of the right ventricle, therefore, the primary muscular ventricular septum separates the entirety of the cavities of the ventricles. The additional step then required to permit closure of the embryonic interventricular foramen, separating completely the newly formed right and left ventricles, is the “transfer” of the subaortic outlet to the left ventricle.

Expansion of the atrioventricular canal enlarged the initial continuity that existed between the cavities of the right atrium and right ventricle. Subsequent to this expansion of the canal, the newly formed right ventricle continues to support the entirety of the developing outflow tract. The subaortic outlet component itself has been formed by fusion of the proximal cushions, or ridges, found within the outflow segment of the primary heart tube (fig 15). This embryonic

Figure 18 This scanning electron micrograph, from a mouse embryo with 42 somites, shows the formation of the muscular ventricular septum. The specimen was prepared by transecting the heart through the atrioventricular canal, and the photograph is of the posterior segment. Note the inferior cushion (IC) occupying most of the posterior margin of the canal, but note also that the floor of the right atrium is in continuity with the roof of the developing right ventricle at the right margin of the atrioventricular canal, even though the atrioventricular groove interposes between the cavities of right atrium and right ventricle (yellow dashed line). Ballooning of the apical parts of the right and left ventricles from the ascending and descending parts of the ventricular component of the primary heart tube, respectively, has produced the primary muscular ventricular septum between them (star). The primary ventricular foramen (yellow bracket) provides the entrance at this stage to the developing right ventricle.

Figure 19 This section from a human embryo at Carnegie stage 14 is comparable to the picture of the mouse heart shown in fig 18. This section, however, has been stained with an antibody to the nodose ganglion of the chick, which recognises the developing atrioventricular conduction tissues, staining them brown. As can be seen, the conduction tissues at this stage form a ring around the primary interventricular foramen (double headed arrow), the inferior part of the ring marking the crest of the developing muscular ventricular septum, and the superior part occupying the right atrioventricular junction. Thus, at this stage the ring is both interventricular and atrioventricular. The section is taken between the unfused atrioventricular cushions, showing that most of the circumference of the atrioventricular canal is supported by the developing left ventricle (black bracket).
outlet septum, when first formed, is of necessity exclusively a right ventricular structure, since the right ventricle at this stage supports the entirety of the developing outflow tract. At this stage, the lumen of the developing aorta is in communication with the left ventricle through the embryonic interventricular foramen, but also remains in continuity with the cavity of the developing right ventricle. It is the fusion of the proximal ends of the outflow cushions with the crest of the muscular ventricular septum that walls the aorta into the left ventricle, this process incorporating the embryonic interventricular communication as the subaortic vestibule (figs 15 and 17). Subsequent to the fusion of the atriocentric and outflow cushions, the ring of tissue positive to the nodose ganglion of the chick can still be seen as a ring surrounding the subaortic vestibule, showing that not only the pulmonary trunk, but also the aorta, was originally supported by the embryonic interventricular foramen. This was confirmed by the development of the right ventricle (fig 20).

After closure of the embryonic interventricular foramen, the fused outflow cushions, which initially formed an embryonic outlet septum, become transformed into the free-standing muscular subpulmonary infundibulum and supraventricular crest of the right ventricle (fig 16). The proximal part of the fused outflow cushions, nonetheless, have fused with the atriocentric cushions to close the embryonic interventricular communication, a process well described by Odgers. It is the conversion of this part of the fused outflow cushions into fibrous tissue, occurring concomitant with delamination of the leaflets of the tricuspid valve, which produces the interventricular part of the membranous septum, by now an integral part of the subaortic vestibule. As has been explained, this interventricular part of the membranous septum is separate from, but contiguous with, the atrioventricular part of the membranous septum, the latter being derived from the fused atriocentric cushions (fig 16).

CONCLUSIONS

The definitive structures that separate the atrial and ventricular cavities are almost entirely muscular structures, albeit that parts are formed by folds or sandwiches between adjacent walls. The small part of the septum that is not muscular, the membranous septum, is formed in part from the atriocentric cushions, and in part from the proximal end of the fused outflow cushions. Although these outflow cushions at first formed an extensive embryonic outflow septum, this tissue, subsequent to its muscularisation, becomes incorporated into the right ventricle as the supraventricular crest and the free-standing subpulmonary infundibulum. The steps involved in this process, along with formation of the arterial valvar leaflets and sinuses, and development of the intrapericardial parts of the arterial trunks, will be the subjects of our final review of cardiac development.

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the heart tube. Only much later are the walls of the separate veins incorporated to form the dome of the left atrium.


In this earlier contribution to dev. Heart, we discussed in greater detail much of the information presented in this review concerning atrial septation.


This important study showed how it was possible to distinguish between the molecular right and left sides of the developing atriums according to the expression of various genes and proteins. The molecular arrangement did not correspond exactly with morphological boundaries.


The study in which we confirmed that far more developmental primordiums were involved in separation of the right and left sides of the atrioventricular canal than mere fusion of the atrioventricular endocardial cushions.


An exquisite study in which Wilhelm His the elder illustrated the mechanism of development of the heart, and pointed to the crucial role of the “spina vestibuli”, a structure subsequently ignored until very recently.


The study that showed how, subsequent to formation of the fibro-muscular atrioventricular junctions, the initial musculature of the embryonic atrioventricular canal was sequestered on the atrial sides of the new region of electrical insulation.


This study used graded series of human embryos, along with future development, to show the extensive moulding of the atrioventricular cushions occurring during human development as they form the aortic leaflet of the mitral valve. The study also re-emphasized the role of His’s “spina vestibuli”.


This study showed how the distribution of atrial natriuretic peptide distinguished between the areas of the cardiac chambers produced by “ballooning” as opposed to the parts derived from the initial primary heart tube.


The initial study that demonstrated how the axis of atrioventricular conduction tissue was derived from a ring of tissue encircling the primary interventricular canal, but a ring that also, from the outset, occupied part of the right atrioventricular junction.


This study charted the expansion of the right atrioventricular junction in the human heart, distinguishing those leaflets of the tricuspid valve involving the various cushions, and stressing the late delamination of parts of the leaflets from the ventricular myocardium.


This study, performed much earlier than the previous one, also stressed the role of late delamination of the leaflets of the tricuspid valve in formation of the membranous part of the septum.


A detailed study, difficult to understand, but showing that the endocardial cushions of both the atrioventricular canal and the outflow tract contributed to formation of the membranous part of the septum.


This exquisite study demonstrated convincingly that, in the chick, the ventricular septum was formed by the fusion of multiple layered trabeculations. There is no evidence thus far that the mammalian septum develops in comparable fashion.


In all three of these studies, it was presumed that the muscular part of the ventricular septum was derived from more than one embryonic primordium. As we have now shown, there is no convincing evidence that supports this notion for development of the mammalian heart.


This exquisite study demonstrated convincingly that, in the chick, the ventricular septum was formed by the fusion of multiple layered trabeculations. There is no evidence thus far that the mammalian septum develops in comparable fashion.


By using marking with Indian ink, these investigators showed that, in the chick, the crest of the muscular ventricular septum was derived from the atrioventricular endocardial cushions. Again, there is no evidence to show that the mammalian septum develops in comparable fashion.


This study, using molecular techniques, confirmed that, in the chick, there is muscularisation of the atrioventricular endocardial cushions, the newly formed muscular area becoming incorporated within the ventricular septum.


The study that amplified the findings initially described in reference 12, relating the findings to tricuspid atresia and double inlet left ventricle.


These studies showed how, in the bird heart, the atrioventricular bundle is sandwiched within the musculature of the ventricular septum, a finding consistent with the observation that the crest of the septum is formed by muscularisation of the atrioventricular endocardial cushions.
Development of the heart: (2) Septation of the atriums and ventricles

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