Common arterial trunk and ventricular non-compaction in Lrp2 knockout mice indicate a crucial role of LRP2 in cardiac development

Maria E. Baardman1, Mathijs V. Zwier2, Lambertus J. Wisse3, Adriana C. Gittenberger-de Groot3, Wilhelmina S. Kerstjens-Frederikse1, Robert M. W. Hofstra4,5, Angelika Jurdzinski2, Beerend P. Hierck3, Monique R. M. Jongbloed6, Rolf M. F. Berger7, Torsten Plösch8,9 and Marco C. DeRuiter3,*

ABSTRACT

Lipoprotein-related receptor protein 2 (LRP2) is important for development of the embryonic neural crest and brain in both mice and humans. Although a role in cardiovascular development can be expected, the hearts of Lrp2 knockout (KO) mice have not yet been investigated. We studied the cardiovascular development of Lrp2 KO mice between embryonic day 10.5 (E10.5) and E15.5, applying morphometry and immunohistochemistry, using antibodies against Tfpα2 (neural crest cells), Nkx2.5 (second heart field), WT1 (epicardium derived cells), tropomyosin (myocardium) and LRP2. The Lrp2 KO mice display a range of severe cardiovascular abnormalities, including aortic arch anomalies, common arterial trunk (persistent truncus arteriosus) with coronary artery anomalies, ventricular septal defects, overriding of the tricuspid valve and marked thinning of the ventricular myocardium. Both the neural crest cells and second heart field, which are essential for the lengthening and growth of the right ventricular outflow tract, are abnormally positioned in the Lp2KO. This explains the absence of the aorto-pulmonary septum, which leads to common arterial trunk and ventricular septal defects. Severe blebbing of the epicardial cells covering the ventricles is seen. Epithelial-mesenchymal transition does occur; however, there are fewer WT1-positive epicardium-derived cells in the ventricular wall as compared to normal, coinciding with the myocardial thinning and deep intertrabecular spaces. LRP2 plays a crucial role in cardiovascular development in mice. This corroborates findings of cardiac anomalies in humans with LRP2 mutations. Future studies should reveal the underlying signaling mechanisms in which LRP2 is involved during cardiogenesis.

KEY WORDS: Cardiac outflow tract, Heart development, Lipoprotein-related receptor protein 2, Neural crest, Second heart field

INTRODUCTION

Lipoprotein-related receptor protein 2 (LRP2), also known as megalin, is a transmembrane glycoprotein receptor on the surface of epithelial cells, and belongs to the low-density-lipoprotein receptor (LDLR) family. The gene is mapped to chromosome 2 in the mouse as well as in human (Korenberg et al., 1994). LRP2 is highly expressed throughout development, starting in the eight-cell stage, and is limited to the trophectoderm in the blastocyst stage (Assemat et al., 2005). After implantation, it is expressed on the maternal-fetal interface, including in the trophectoderm, the visceral endoderm of the yolk sac and the placenta (Gueth-Hallonet et al., 1994). During further development, LRP2 is detected in many epithelial tissues, including the neuroepithelium, the kidney proximal tubule epithelial cells and the pericardium (Assemat et al., 2005). LRP2 is a multi-ligand receptor and capable of binding important signaling molecules, such as bone morphogenetic protein 4 (BMP4) (Sooligen et al., 2005), sonic hedgehog (Shh) (McCarthy and Argraves, 2003), retinol binding protein (RBP) (Christensen et al., 1999) and vitamin B12 (Moestrup et al., 1996). Furthermore, LRP2 is a low-density lipoprotein (LDL) receptor on the yolk sac and placenta, and is involved in cholesterol transport as a co-transporter with cubilin (Christensen and Verrout, 2002).

Targeted disruption of the gene in mice results in severely malformed offspring with holoprosencephaly, neural tube defects, limb anomalies and a 100% lethality in utero or within 24 h after birth (Willnow et al., 1996). Disruption in humans is known as the autosomal-recessive Donnai-Barrow syndrome. Individuals with Donnai-Barrow syndrome display severe congenital malformations, including agenesis of the corpus callosum, sensorineural deafness, diaphragmatic hernia, omphalocele and nephritis (Kantarci et al., 2007). Literature reports of a few cases with heart anomalies such as double-outlet right ventricle, persistence of the left caval vein, persistent ductus arteriosus and ventricular septal defects; however, authors assume that because of early fetal loss of severe cases and limited examination of fetuses, the real incidence of heart anomalies could be much higher (Pober et al., 2009).

Interestingly, the effect of depletion of LRP2 on the cardiovascular development has never been described in mice, although expression of LRP2 on the mesothelial cells of the pericardial cavity has been reported together with expression of LRP2 in the neural crest (Assemat et al., 2005; Fisher and Howie, 2006). Previous studies have consistently shown that the epicardium, which arises from the pericardial mesothelium, and neural crest play important roles in the development of different...
cardiac structures such as the ventricular myocardium, the atrioventricular valves, the coronary arteries and the distal part of the outflow tract (Gittenberger-de Groot et al., 2000, 2012; Poelmann et al., 1998; Waldo et al., 2005; Wessels et al., 2012). We examined in a developmental series the hearts of Lrp2 knockout mouse embryos to elucidate a role of LRP2 in cardiac development and to determine which cardiac and/or extracardiac cell population is affected by the absence of the protein. We postulate that LRP2 plays a role in the formation of the compact ventricular myocardium through interaction with epicardium-derived cells (EPDCs). The transcription factor Wilms tumor 1 (WT1) is expressed in nuclei of the epicardium and in EPDCs during their early migration, whereas myocardial cells are negative for this protein (Moore et al., 1999), and is therefore considered to be a valid EPDC marker. We further examined the distribution patterns of cells from the neural crest (NCCs) (Jain et al., 2011; Keyte and Hutson, 2012; Poelmann et al., 1998; Waldo et al., 2005) and the second heart field (SHF) (Dyer and Kirby, 2009; Mjaatvedt et al., 2001; Scherptong et al., 2012; Waldo et al., 2001) because both are known to be crucial for outflow tract (OFT) septation and remodeling. Because LRP2 acts as a receptor in the retinoic acid (RA) and Shh signaling pathways, which are indispensable in SHF and NCC differentiation, we postulate a role for LRP2 in cardiac OFT formation. A valid and useful marker is Nkx2.5, which is highly expressed by cells from the SHF during their migration towards the OFT (Anderson et al., 2012; Scherptong et al., 2012). Extracardiac-positioned SHF cells that are Nkx2.5-positive, but negative for myocardial markers such as tropomyosin and myosin light chain kinase, migrate towards the distal OFT to establish right OFT lengthening, rotation and septation (Anderson et al., 2012). Failure of this process can result in OFT abnormalities (Bajolle et al., 2006). The contribution of NCCs to the OFT in Lrp2 knockout and wild-type mice was studied by using Tlap2α, a transcription factor that is highly expressed by migrating and differentiating NCCs and not by cells of the SHF (Jain et al., 2011).

RESULTS
LRP2 is crucial for development of the cardiac outflow tract and atrioventricular canal
Cardiac malformations were seen in all 17 analyzed Lrp2 knockout embryos from E12.5 to E15.5 and are summarized in Table 1. The most obvious cardiac anomaly was a common arterial trunk (CAT) (Dyer and Kirby, 2009; Mjaatvedt et al., 2001; Scherptong et al., 2012; Waldo et al., 2001) because both are known to be crucial for outflow tract (OFT) septation and remodeling. Because LRP2 acts as a receptor in the retinoic acid (RA) and Shh signaling pathways, which are indispensable in SHF and NCC differentiation, we postulate a role for LRP2 in cardiac OFT formation. A valid and useful marker is Nkx2.5, which is highly expressed by cells from the SHF during their migration towards the OFT (Anderson et al., 2012; Scherptong et al., 2012). Extracardiac-positioned SHF cells that are Nkx2.5-positive, but negative for myocardial markers such as tropomyosin and myosin light chain kinase, migrate towards the distal OFT to establish right OFT lengthening, rotation and septation (Anderson et al., 2012). Failure of this process can result in OFT abnormalities (Bajolle et al., 2006). The contribution of NCCs to the OFT in Lrp2 knockout and wild-type mice was studied by using Tlap2α, a transcription factor that is highly expressed by migrating and differentiating NCCs and not by cells of the SHF (Jain et al., 2011).

Cardiac malformations were seen in all 17 analyzed Lrp2 knockout embryos from E12.5 to E15.5 and are summarized in Table 1. The most obvious cardiac anomaly was a common arterial trunk (CAT) (Dyer and Kirby, 2009; Mjaatvedt et al., 2001; Scherptong et al., 2012; Waldo et al., 2001) because both are known to be crucial for outflow tract (OFT) septation and remodeling. Because LRP2 acts as a receptor in the retinoic acid (RA) and Shh signaling pathways, which are indispensable in SHF and NCC differentiation, we postulate a role for LRP2 in cardiac OFT formation. A valid and useful marker is Nkx2.5, which is highly expressed by cells from the SHF during their migration towards the OFT (Anderson et al., 2012; Scherptong et al., 2012). Extracardiac-positioned SHF cells that are Nkx2.5-positive, but negative for myocardial markers such as tropomyosin and myosin light chain kinase, migrate towards the distal OFT to establish right OFT lengthening, rotation and septation (Anderson et al., 2012). Failure of this process can result in OFT abnormalities (Bajolle et al., 2006). The contribution of NCCs to the OFT in Lrp2 knockout and wild-type mice was studied by using Tlap2α, a transcription factor that is highly expressed by migrating and differentiating NCCs and not by cells of the SHF (Jain et al., 2011).
E15.5, all the embryos presented with abnormal coronary arteries. In 3/5 embryos we found a single left coronary artery, in 1/5 a single right coronary artery and in 1/5 a very wide left coronary artery in combination with a hypoplastic right coronary artery. There was no abnormally high take-off of the arteries seen (data not shown in figure).

Also at E15.5, in 5/5 embryos the right pleural cavity was not separated from the pericardial cavity, resulting in an abnormal position of the right lung next to the heart.

LRP2 is crucial for migration and development of the second heart field and neural crest cells

At E10.5, LRP2 was highly expressed in the mesenchyme surrounding the non-septated cardiac OFT region (Fig. 2A). The LRP2 expression was mainly in the non-myocardial cells (Nkx2.5-negative cells). During OFT septation, the wall of the ascending aorta (Fig. 2B) and pulmonary trunk remained LRP2-positive. The differentiating myocardium (Nkx2.5-positive) and epicardium were LRP2-negative at E11.5. At E12.5, LRP2 in the heart was mainly expressed by the mesenchymal cells in the OFT cushions (Fig. 2C), although the endocardial cells and the epicardium were also slightly positive. At E13.5, the LRP2 expression became more distinct in the developing intercalated cushions (Fig. 2D) and the expression in the epicardium was no longer detectable. In addition to the heart, the mesothelial lining of the pericardial cavity also expressed LRP2 from E10.5 to E13.5 (Fig. 2B-E).

In the non-myocardial OFT area (tropomyosin-negative), from which the aorta and pulmonary trunk develop, both NCCs (Tfap2α-positive) and SHF cells (Nkx2.5-positive) were found to be present (Fig. 3). Tfap2α and Nkx2.5 are not co-expressed in the stages studied. Therefore, these markers can be used to study the two separate cell populations. The non-myocardial Nkx2.5-positive mesenchymal cells are considered as SHF cells and will contribute to part of the smooth-muscle cell population of the ascending aorta and pulmonary trunk, as well as the myocardium of the OFT of both the left and right ventricle (Scherptong et al., 2012). Also, NCCs are present within this area and will contribute to the smooth-muscle cell population and OFT septum, but not to the myocardium of the OFT.

Already at E10.5 we found differences in the patterning and distribution of the Nkx2.5-positive SHF as well as the Tfap2α NCC population in the knockout population as compared to the control embryos (Fig. 4A-L). The SHF population in the wild-type embryos showed a mid-sagittal position in front of the aortic sac and provided a strand of cells between the 3rd and 4th pharyngeal arch artery towards the right (future aortic side) of the aortic sac (Fig. 4A). This population ended in the deep part of the (saddle-shaped) orifice level at the border of the myocardial OFT (Fig. 4A, asterisk). A second population of SHF cells was positioned immediately in front of the foregut, forming a flow divider (FD) in between the 6th pharyngeal arch arteries (Figs 3C and 4E, arrow). Remarkably, this FD was negative for NCCs, which were prominently present around the 6th pharyngeal arch arteries (Fig. 4F,J). The main body of the dorsal mid-sagittal SHF population was positioned in between the left and right 4th and the left and right 6th pharyngeal arch arteries, where it also showed a marked extension to the left side (pulmonary side) of the OFT (Fig. 4B,I, black arrowhead).
At E11.5 this stream of SHF cells coursing between the 4th and 6th pharyngeal arch arteries became even more prominent (Fig. 5A,B,E,F). Together with a leftward and upward shift of the dorsal SHF population, it mainly contributed to the pulmonary OFT. This process has been referred to as ‘pulmonary push’ (Scherptong et al., 2012). During this leftward shift and extension of the SHF, also the FD that separates the left and right 6th pharyngeal arch arteries moves to the left (Fig. 5E,F, arrow). At this stage the NCCs were abundantly present around the pharyngeal arch arteries and extend with prongs into the endocardial cushions (Fig. 5A,B, C).
bright green, and G). They formed the aorto-pulmonary septum separating the aortic sac into the aorta and the pulmonary trunk (Fig. 5E-G, double arrow). The pulmonary trunk was still connected to the 6th pharyngeal arch system and the connecting pulmonary arteries (Fig. 5B).

In the mutant embryos at E10.5 and E11.5, several differences were observed both in the distribution of the SHF and the NCC. The anteriorly located SHF population directed towards the right (aortic side) of the OFT was less well developed and less organized in the mutant embryo compared to the wild type (compare Fig. 4C to 4A). The main difference correlating with the development of a CAT was the diminished development of the left-sided SHF population that leads to the pulmonary push (compare Fig. 4D to 4B and 5D to 5B). The mid-sagittal SHF population remained in the mutant in the low position and did not properly extend anteriorly (D). The mesenchymal NCC population (dark green) on the left (pulmonary) side already reaches the NCCs in the endocardial OFT (D). The midline dorsal FD (arrows in G, H, K, L) protrudes prominently into the AoS and separates the 6th pharyngeal arch arteries. Scale bars: 50 µm. See also the online supplemental videos (E10.5_video_1080p.mpg) and interactive pdf reconstructions (E10.5_WT_revise.pdf and E10.5_KO_revise.pdf) at http://www.caskanatomy.info/research/supplement_megalin.html. S, superior; R, right; L, left; I, inferior; A, anterior; P, posterior; ec, endocardial cushions.
Normally, increase of the compact ventricular myocardium is observed (Fig. 6A) corresponding with normal development. E12.5-E13.5, in contrast to E14.5 and E15.5 where a strong increase of the total volume of the ventricular myocardium hardly increased from E12.5 to E15.5, measured the volumes of the compact ventricular myocardial layer and compare Fig. 1C with 1D). The thickness of the free wall myocardium varied strongly between the knockout mice (Fig. 7). Of all embryos studied, 14 embryos had abnormal epicardial blebbing and most of them (n=13) contained red blood cells on the outer surface of the left ventricle (Table 1, Fig. 7C,D) connecting to deep intertrabecular spaces of the myocardial wall (Fig. 7D,F).

Because Lrp2 knockout mice display hypoplasia of the ventricular myocardium, we investigated whether this was due to abnormal formation of the compact myocardium. Therefore, we measured the volumes of the compact ventricular myocardial layer including the ventricular septum from E12.5 to E15.5, encompassing the total volume of the ventricular myocardium (compact and trabecular myocardium). In wild-type embryos the total volume of the ventricular myocardium hardly increased from E12.5 to E13.5, in contrast to E14.5 and E15.5 where a strong increase is observed (Fig. 6A) corresponding with normal development. Normally, increase of the compact ventricular myocardium is a linear process marked by a regression coefficient that is close to 1 (measured from E12.5 to E15.5) (Fig. 6C, R^2 = 0.98). The amount of compact myocardium measured as the ratio of the total ventricular myocardium increased strongly at E14.5 and E15.5 (Fig. 6B). This ratio was not significantly reduced in the knockout embryos for E12.5 to E13.5, but was significantly reduced for E14.5 and E15.5 (Fig. 6B, P=0.019 and P=0.008). In mutants we did not observe the same linear process of formation of the compact ventricular myocardium. For the younger stages (E12.5, E13.5 and E14.5) it seems to start with a somewhat linear process; however, at E14.5 and E15.5 (light green and green; Fig. 6C), we observed a lower volume of the compact myocardium relative to the total ventricular myocardium, and wider distributions within each gestation age (R^2 = 0.68).

At E13.5 the epicardium and one to two layers of subepicardial cells were equally present in both wild-type and knockout embryos (Fig. 8A,B). At E14.5 a reduced WT1 expression was observed in the free wall of the right ventricle in the knock out compared to the wild type, the reduction being even more prominent in the free wall of the left ventricle (Fig. 8C,D). At E15.5, WT1-expressing EPDCs were still recognizable in the ventricular myocardial wall; however, the WT1 expression was still more reduced in the knock out (Fig. 8E,F).
DISCUSSION

LRP2 is crucial for embryonic development in mice and humans and a role has been mentioned in the NCCs (Spoelgen et al., 2005) and in Shh signaling (McCarthy and Argraves, 2003), but a thorough analysis in cardiac development is lacking. Therefore, we investigated the hearts of Lrp2 knockout embryos in a developmental series between E10.5 and E15.5. For the first time, we were able to demonstrate that LRP2 is crucial for normal cardiac development. Absence of LRP2 resulted in OFT anomalies (CAT, aortic arch and coronary artery anomalies), ventricular septal defects and overriding of the tricuspid valve as well as a marked reduction of the compact layer of the ventricular myocardium, epicardial blebbing and the presence of deep intertrabecular spaces.

Septation of the outflow tract is a process that is orchestrated by the NCCs and the SHF (Anderson et al., 2012; Dyer and Kirby, 2009; Jain et al., 2011; Keyte and Hutson, 2012; Li et al., 2004; Mjaatvedt et al., 2001; Scherptong et al., 2012; Waldo et al., 2001). During lengthening and septation of the OFT, LRP2 is expressed in all the contributing cell populations, i.e. SHF (with the exception of the differentiated myocardium), NCCs, endocardium, epicardium and pericardial mesothelium, indicating the potential for involvement in OFT formation. Here, we show that, in Lrp2 knockout embryos, OFT septation fails. The SHF population does provide, in both wild-type and knockout embryos, a ridge, which we refer to as an FD, in a mid-sagittal position anterior to the gut that separates in the posterior wall of the aortic sac both 6th pharyngeal arch arteries. In the Lrp2 knockout embryos the bulk of the SHF population remains in a mid-sagittal position, resulting in absence of the pulmonary push (Scherptong et al., 2012), which is necessary to create a right-sided space in which NCCs can form the aorto-pulmonary septum that septates the aortic sac into a pulmonary and aortic side. Failure of this septation process as seen in CAT in the mutant embryos shows rerouting of the mesenchymal NCC population towards the left (pulmonary side), where these cells connect to the endocardial cushion tissue of the OFT. The pulmonary side of the aortic sac including the connection of the 6th pharyngeal arch arteries and their intermediate FD remain connected to the common OFT. So no NCC-containing aorto-pulmonary septum is formed at the proper side necessary for fusion of the endocardial OFT cushions, resulting in a CAT in 15/17 Lrp2 knockout embryos. Because our data show that LRP2 is expressed by both the cardiac NCCs and the non-myocardial SHF-derived...
cells, the combined disturbed remodeling of the NCC and SHF population seems to underlie the OFT anomalies seen in Lrp2 knockout embryos. Fig. 9 provides a schematic depiction of the relative disposition of the crucial elements for OFT septation.

More than half of the Lrp2 knockout embryos had an inlet ventricular septal defect (VSD) and overriding tricuspid valve orifice. Also, other studies (e.g. Bartram et al., 2001) have previously demonstrated that OFT abnormalities can coincide with overriding tricuspid valve and inlet VSD, and concluded that this combination of cardiac malformations represents disturbances of the final phase of the cardiac looping process. In our study the defective addition of SHF to the OFT is accompanied by a deficient movement of the atrioventricular canal to the right. This process can be seen as final stage deficient looping and leads to malalignment of the various septal components (Gittenberger-de Groot et al., 2014). In the case of the Lpr2 mutant, the OFT septation can be appreciated as being completely out of line, whereas primary atrial and inlet septum are malaligned with subsequent overriding of the tricuspid orifice accompanied by an inlet VSD.

Secondly, the process of formation of the compact ventricular myocardium was severely disturbed. This process is orchestrated by EPDCs that migrate from the epicardium around E13.5 into the ventricular myocardium, reviewed by Gittenberger-de Groot et al. (2012), where they contribute to the myocardial architecture as interstitial fibroblasts (Gittenberger-de Groot et al., 1998). Normally, at E15.5 the process of formation of the compact myocardium is almost finished. At E15.5, there was a significant reduction in the volume of the compact myocardium in the knockout embryos. The epicardium showed severe blebbing at several sites with blood-filled spaces that seemed to be in contact with deep ventricular intertrabecular spaces. These anomalies coincided with a reduced WT1 expression. These data suggest that downstream signaling that determines EPDC differentiation into fibroblasts and the necessary interaction of EPDCs with the myocardial cells to form the compact layer of the ventricular myocardium is disturbed, as has been described for other animal models (Gittenberger-de Groot et al., 2012). Also, the observation of a great number of single coronary arteries in the CAT cases of the mutant embryos points towards an epicardial disturbance (Eralp et al., 2005). The focus on a possible primary epicardial problem is based on the observation that Lrp2 is temporarily expressed in the epicardium at E13.5 and in the epicardium around the great arteries at E14.5, but not in the differentiated ventricular myocardium.

**Resemblance of the Lrp2 knockout with other animal models points towards disturbed RA and Shh signaling**

Retinoic acid (RA) is crucial for cardiac development, especially for the OFT and the formation of the compaction ventricular myocardium (Jiang et al., 2002; Niederreither et al., 2001). Because LRP2 is a retinol-binding protein receptor (Christensen et al., 1999)
and the Lrp2 knockout mouse displays both OFT anomalies and a less-compact ventricular myocardium as demonstrated in this manuscript, the underlying mechanism is suspected to be disturbed RA signaling. Other knockout models point in the same direction. Retinoic acid receptor (RAR) knockouts, for example RARα1/RARβ double knockouts, have a 100% penetrance of CAT (Jiang et al., 2002). In these mice a marked shortened OFT arises from the right ventricle. RA signaling is carried by the SHF and the cardiac NCCs (Dyer and Kirby, 2009). Interestingly, Jiang et al. showed that, in RARα1/RARβ knockout mice, the number, migration and differentiation of NCCs were normal, highly resembling our Lrp2 knockout mice (Jiang et al., 2002). Furthermore, resemblance of the phenotype of individuals with a STRA6 mutation (Mathew-Woods syndrome) with individuals with an LRP2 mutation (Donnai-Barrow syndrome) is obvious, as is the resemblance of a zebrafish model in which STRA6 function is disrupted (Isken et al., 2008; Scherz et al., 2008; Segel et al., 2009) with the Lrp2 knockout mouse. The phenotype in humans with a STRA6 mutation includes OFT anomalies, whereas extracardiac anomalies highly resemble the phenotype of the Lrp2 knockout mouse, and humans with an LRP2 mutation with diaphragmatic hernia and anophthalmia. STRA6 is the most important RBP receptor in humans (Sun, 2012). Unfortunately, little is known about STRA6 expression during heart development, and the cardiac phenotype in the zebrafish model has not been studied in detail as well. Further studies are necessary to elucidate how co-expression of STRA6 with LRP2 affects RBP uptake during OFT formation and myocardial compaction.

It is possible that the multi-ligand receptor LRP2 influences different pathways during cardiac development or maybe even plays an as-yet-unknown key role in connecting pathways, as has been seen for TBX1 (Garg et al., 2001), which influences both Shh- and RA-dependent signaling (Liao et al., 2008). We know that morphogens like Shh and RA act in parallel with other pathways, such as the BMP and Nodal pathway, in the regulation of shared target genes (Maynard et al., 2013; Wong et al., 2012). Future studies should unravel the position of Lrp2 in these networks of genes.

The Sonic hedgehog (SHH)-GLI pathway plays a role in many syndromes associated with septal defects (reviewed by Gittenberger-de Groot et al., 2014). The primary cilium present on the endocardial cells is involved in this signaling process, which controls the endocardial-mesenchymal transformation in the cardiac cushions. LRP2 has been recently recognized as a cilium-related gene (Li et al., 2015). Our staining for LRP2 expression demonstrated endocardial expression in the OFT cushions but did not reveal a distinct expression in the atrioventricular cushions. Although we cannot rule out a role of LRP2 in the atrioventricular cushions, the described malformations at the inflow seems to be more related to the disturbed looping processes described above than to hampered endothelial-mesenchymal transformation.

**LRP2 as a cholesterol transporter on the yolk sac and placenta as a possible causative factor in the multifactorial model**

Cholesterol is important for embryonic development. Individuals with Smith-Lemli-Opitz syndrome, an autosomal-recessive disease
of endogenous cholesterol synthesis, display a broad range of congenital anomalies resembling those in individuals with Donnai-Barrow syndrome (Digilio et al., 2003; Smith et al., 1964). The underlying mechanism in Smith-Lemli-Opitz syndrome is a disturbed SHH-GLI signal transduction: cholesterol is essential to activate this pathway (Digilio et al., 2003).

It is postulated that besides endogenous cholesterol synthesis also maternal-fetal cholesterol transport across the yolk sac and placenta is important for embryonic development (Baardman et al., 2013; Woollett, 2005). LR2 function is intriguing because it is expressed at all maternal-fetal interfaces throughout development (Woollett, 2005). Furthermore, ligands capable of binding to LR2 are essential for embryonic development as mentioned before. The question raised is whether a disturbed maternal-fetal cholesterol transport can systemically cause the anomalies seen in Lrp2 knockout mice or, alternatively, whether defects in fundamental local signaling cascades such as disturbed Wnt-BMP4 signaling, Shh signaling, RA signaling, or even impaired local uptake of cholesterol by the heart itself are the underlying causes. Targeted disruption of Lrp2 specific for the neural floor plate during embryogenesis resulted in offspring with the same neural tube and forebrain anomalies as seen in the complete knockout, which suggests that the function of LR2 in signaling cascades is likely causing the neurological phenotype (Hammes et al., 2005; Spoelgen et al., 2005). Impaired NCC development and differentiation plays a key role in the neural tube and forebrain defects in the Lrp2 knockout mouse, together with disturbed Shh signaling (Hammes et al., 2005). Possible future studies to reveal the exact underlying mechanism could include a yolk-sac- and placenta-specific Lrp2 knockout in which the heart is thoroughly examined and functional studies conducted that analyze the cholesterol transport function of LR2 in utero.

In summary, Lrp2 knockout mice display a range of cardiovascular abnormalities resulting from failure of OFT septation, failure of a normal ventricular inlet formation, and failure of formation of the compact ventricular myocardium. The underlying process for OFT abnormalities are severe abnormalities in the distribution of both NCCs and SHF-derived cells, and development towards the base of the OFT. The underlying process for failure of the formation of the compact ventricular myocardium remains unclear but seems to be related to deficient EPDC interaction with the ventricular myocardium. Disturbed downstream signaling in which LR2 functions in the RA pathway is a potential causative factor. However, we contemplate that the function of LR2 on cholesterol transport either in the heart itself or at the level of the maternal-fetal interfaces might also play a significant role in cardiac development.

Based on the range of cardiovascular abnormalities found in Lrp2 knockout mice, together with the existence of a human phenotype (Donnai-Barrow syndrome) that includes cardiac anomalies resembling those seen in the Lrp2 knockout mouse, we propose that LR2 also plays an important role in cardiogenesis in humans, thereby regulating the development of multiple cardiac structures such as the OFT and the cardiac septa by orchestrating the migration of both the SHF and the NCCs. Further research must focus on identifying LR2 mutations in humans with syndromic CHD, including CHD with associated extracardiac anomalies.

**MATERIALS AND METHODS**

**Generation of Lrp2 knockout mice and harvesting of embryos**

Animal use was in accordance with the University Medical Center Groningen (UMCG) institutional guidelines. All animals received humane care and all experiments were reviewed and approved by the
local Animal Experimental Committee. Heterozygous Lrp2+/− mice on a CB56BL/6 background were kindly provided by Thomas Willnow (Max Delbrück Center, Berlin, Germany). Lrp2+/− mice were housed in the animal facility under standard conditions. Food and water were provided ad libitum. Lrp2+/− males were crossed with heterozygous females. We observed from 26 litters a normal mendelian ratio of Lrp2+/+ (n=36), Lrp2+/− (n=67) and Lrp2−/− (n=38) embryos. The average number of embryos per pregnancy was 8.2. The day of appearance of the vaginal plug was considered as E0.5. Pregnant females were sacrificed, embryos harvested and the whole embryo (E10.5-E12.5) or solely the thorax (E13.5-E15.5) were isolated.

Per age group we randomly selected wild-type (E12.5: n=3, E13.5: n=4, E14.5: n=6, E15.5: n=5) and knockout (E12.5: n=4, E13.5: n=4, E14.5: n=4, E15.5: n=5) embryos to phenotype the cardiac malformations. Additionally, we studied the contribution of the NCCs and SHF in wild-type embryos and Lrp2 knockout embryos. The day of appearance of the vaginal plug was considered as E0.5. Pregnant females were sacrificed, embryos harvested and the whole embryo (E10.5-E12.5) or solely the thorax (E13.5-E15.5) were isolated.

Morphological examination and 3D reconstruction

3D reconstructions were generated of representative E10.5, E11.5 and E15.5 wild-type and knockout embryos, displaying the morphological differences. The reconstructions were made as described earlier (Scherppong et al., 2012), establishing the various stainings and the relative position of the NCC and SHF populations in each reconstruction, using the AMIRA software package version 5.4 (Template Graphics Software, San Diego, CA). Reconstructions can be downloaded from: http://www.caskanatomy.info/research/supplement_megalin.html.

Myocardial morphometry

We performed morphometry on the ventricular myocardium (E12.5-E15.5) and the endocardial cushions (E10.5-E14.5) using the method of Gunderson et al. based on Cavalleri’s principle (Gunderson et al., 1988). In short, to measure ventricular myocardium we randomly positioned regularly spaced points (49 mm² grid for E12.5 and E13.5, and 100 mm² for E14.5 and E15.5) on the MLC-2a-stained ventricular myocardium of embryos from E12.5 to E15.5. We compared the total ventricular myocardial volumes of wild-type embryos and Lrp2 knockout embryos from different litters. For E15.5 we also analyzed five available heterozygous embryos. After analyzing the five heterozygous embryos of E15.5 and confirming that heterozygous Lrp2 embryos did not have a phenotype, we counted two heterozygous embryos of E12.5 and included them in the wild-type group for this embryonic day. We established the percentage of compact and trabecular myocardium in these wild-type and knockout embryos. The distance between the subsequent sections of the slides was either 0.075 mm or 0.01 mm depending on the number of sections that differed between the embryos. Ten sections were considered to be sufficient for morphometry. Volume measurements were performed using an Olympus microscope with a 40× or 100× magnification. The distance between the subsequent sections of the slides was either 0.025 mm or 0.05 mm.

Statistics

Statistical analysis on the volume measurement was performed with Mann-Whitney U-test because the data were not perfectly normally distributed and because of low sample size. We calculated the regression coefficient for both wild-type and knockout embryos for the ratio of compact myocardium/total ventricular myocardium to analyze the growth of the compact myocardium relative to the growth of the total ventricular myocardium. A regression coefficient close to 1 was considered to be a linear relation between the two. All data of the volume measurements have been presented as average±s.d. in the graphs. Significance was assumed when P<0.05 using the SPSS 20.0 software program (SPSS Inc., Chicago, USA).

Acknowledgements

We are very grateful to Robert Poelmann for critically reading the manuscript and Margot Bartelings for valuable discussions on development of CAT.

Competing interests

The authors declare no competing or financial interests.

Author contributions

M.C.D., R.M.W.H., R.M.F.B., W.S.K.-F. and T.P. conceived and designed the experiments. M.E.B., M.V.Z., A.J., B.P.H. and L.J.W. performed the experiments. M.E.B., L.J.W., M.V.Z., M.R.M.J., A.C.G.-d.G. and M.C.D. analyzed the data.

Funding

T.P. is supported by the Dutch Heart Foundation (grant 2004T4801) and the Netherlands Organization for Scientific Research (TOP ZonMw 40008129811053). Support of the Seabold Fund is gratefully acknowledged.

Supplementary information

Supplementary information available online at http://dmm.biologists.org/lookup/suppl/doi:10.1242/dmm.022053/-/DC1
Wessels, A., van den Hoff, M. J. B., Adamo, R. F., Phelps, A. L., Lockhart, M. M., Sauls, K., Briggs, L. E., Norris, R. A., van Wijk, B., Perez-Pomares, J. M. et al. (2012). Epicardially derived fibroblasts preferentially contribute to the parietal leaflets of the atrioventricular valves in the murine heart. Dev. Biol. 366, 111-124.

Willnow, T. E., Hilpert, J., Armstrong, S. A., Rohlmann, A., Hammer, R. E., Burns, D. K. and Herz, J. (1996). Defective forebrain development in mice lacking gp330/megalin. Proc. Natl. Acad. Sci. USA 93, 8460-8464.

Wong, K. S., Rehn, K., Palencia-Desai, S., Kohli, V., Hunter, W., Uhl, J. D., Rost, M. S. and Sumanas, S. (2012). Hedgehog signaling is required for differentiation of endocardial progenitors in zebrafish. Dev. Biol. 361, 377-391.

Woollett, L. A. (2005). Maternal cholesterol in fetal development: transport of cholesterol from the maternal to the fetal circulation. Am. J. Clin. Nutr. 82, 1155-1161.