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## Epicardial progenitors contribute to the cardiomyocyte lineage in the developing heart

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### Abstract

The heart is formed from cardiogenic progenitors expressing the transcription factors Nkx2-5 and Is11<sup>1,2</sup>. These multipotent progenitors give rise to cardiomyocyte, smooth muscle, and endothelial cells, the major lineages of the mature heart<sup>3,4</sup>. Here we identify a novel cardiogenic precursor marked by expression of the transcription factor *Wt1* and located within the epicardium, an epithelial sheet overlying the heart. During normal heart development, a subset of these *Wt1*<sup>+</sup> precursors differentiated into fully functional cardiomyocytes. *Wt1*<sup>+</sup> proepicardial cells arose from progenitors that express Nkx2-5 and Is11, suggesting that they share a developmental origin with multipotent *Nkx2-5*<sup>+</sup>/*Is11*<sup>+</sup> progenitors (Suppl. Fig 1). These results identify *Wt1*<sup>+</sup> epicardial cells as previously unrecognized cardiomyocyte progenitors, and lay the foundation for future efforts to harness the cardiogenic potential of these progenitors for cardiac regeneration and repair.

Epicardial cells migrate from the proepicardium (PE), an outgrowth of the septum transversum, and spread over the surface of the heart<sup>5,6</sup>. A subset of epicardial cells transition to a mesenchymal phenotype, migrate into the subjacent myocardium, and differentiate into smooth muscle cells (SMCs) and endothelial cells (ECs)<sup>7-13</sup>. *Wt1* was expressed in PE and epicardium, but not in myocardium (Fig. 1a-c). In order to trace the fate of *Wt1*-expressing PE and epicardial cells, we knocked a *GFPCre* cDNA<sup>14</sup> into the endogenous *Wt1* start codon (Suppl. Fig. 2). GFP and Cre expression in *Wt1*<sup>GFPCre/+</sup> embryos co-localized with *Wt1*, indicating that the knockin strategy placed *GFPCre* under control of endogenous *Wt1* regulatory elements (Fig. 1d-f). In the heart, *GFPCre* expression was confined to PE and epicardium from E9.5 to E15.5, and not found in the myocardium (Fig. 1d-g).

We used *Wt1*<sup>GFPCre</sup> and the Cre-activated reporters *Rosa26*<sup>fsLz15</sup> and *Z/Red*<sup>16</sup> to analyze the fate of *Wt1*-expressing cells in the heart. Upon Cre-mediated recombination, these reporters heritably express β-galactosidase (β-gal) or red fluorescent protein (RFP), respectively. Using two different reporters minimized potential artefacts related to unanticipated behavior of Cre-activated reporters, or to false-positive immunostaining. While *Wt1* and *GFPCre* expression were confined to the epicardium, descendants of *Wt1*-expressing progenitors (hereafter called *Wt1*-derived cells), marked by β-gal, were found in a mosaic pattern throughout the

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myocardium (Fig. 2a). Consistent with prior reports showing that epicardially-derived mesenchyme predominately differentiates into SMCs in mammals, most *Wt1*-derived cells adopted a SMC fate, and a minority differentiated into ECs (Suppl. Fig. 3) <sup>8,9</sup>.

Remarkably, we found that some *Wt1*-derived cells differentiated into cardiomyocytes (CMs) during normal heart development, as demonstrated by co-expression of lineage tracers and CM markers cardiac troponin T (*Tnnt2*) and sarcomeric  $\alpha$ -actinin (*Actn1*) (Fig. 2b-d). The *Wt1*-derived CMs also expressed cardiac transcription factors *Gata4* and *Nkx2-5* (Suppl. Fig. 4). *Wt1*-derived CMs were located in the myocardium of all four cardiac chambers and in the interventricular septum, constituting 7-10% of CMs in ventricles and 18% in atria (Suppl. Fig. 5). We further confirmed co-expression of CM and lineage tracers in isolated cells by immunostaining of dissociated fetal heart cultures (Fig. 2e-f). 4% of CMs in dissociated fetal heart cultures were *Wt1*-derived (Suppl. Fig. 5), comparable to the frequency observed in tissue sections.

To determine if *Wt1*-derived CMs had functional properties of CMs, we analyzed dissociated cells from *Wt1<sup>GFP<sup>Cre/+</sup></sup>*; *Z/Red* fetal hearts. A subset of red fluorescent cells exhibited spontaneous contractile activity (Fig. 2g; Suppl. Movie 1). In addition, these contracting RFP<sup>+</sup> cells exhibited calcium oscillations with kinetics, amplitude, and frequency characteristic of CMs (Fig. 2h-j; Suppl. Movie 2). Also characteristic of CMs were calcium sparks preceding calcium waves (Fig. 2h1) and caffeine augmentation of calcium transient amplitude (Fig. 2j), consistent with calcium release from CM ryanodine receptors. Calcium transients of RFP<sup>+</sup> cells were indistinguishable from and synchronous with adjacent RFP<sup>-</sup> cells (Fig. 2i), suggesting electrical coupling between *Wt1*-derived and non-*Wt1*-derived CMs. Consistent with this finding, the gap junction protein connexin 43 (*Cx43*) localized to the membrane between *Wt1*-derived and neighboring CMs (Suppl. Fig. 6a). A similar pattern of *Cx43* expression was observed in *Wt1*-derived CMs in tissue sections of E15.5 hearts (Suppl. Fig. 6b), suggesting that *Wt1*-derived CMs are also electrically coupled to other CMs in vivo. Collectively, these data indicate that during heart development a subset of *Wt1*-expressing cells differentiate into CMs.

To further characterize the *Wt1*-expressing CM precursors, we utilized several independent methods to control the temporal and spatial window during which these precursors were labeled. To temporally regulate Cre-labeling, we knocked a cDNA encoding a Cre-modified estrogen ligand binding domain (CreERT2) into the *Wt1* locus (Suppl. Fig. 7). CreERT2 fusion protein recombinase activity requires tamoxifen <sup>17</sup>. Maternal injection of tamoxifen at E10.5 and E11.5 induced Cre activity and resulted in  $\beta$ -gal expression within *Wt1<sup>CreERT2</sup>*; *Rosa26<sup>flLz</sup>* myocardium (Fig. 3a-e), while *Wt1<sup>CreERT2</sup>* did not activate *Rosa26<sup>flLz</sup>* in the absence of tamoxifen (Fig. 3b and Suppl. Fig. 7). The frequency of *Wt1<sup>CreERT2</sup>*-labeled cells in epicardium and myocardium (Suppl. Fig. 7-8) was reduced compared to constitutive labeling by *Wt1<sup>GFP<sup>Cre</sup></sup>*, likely due to inefficient CreERT2 activation by tamoxifen concentrations compatible with maintenance of pregnancy. Co-staining for differentiation markers showed that  $\beta$ -gal<sup>+</sup> cells differentiated into CM, EC, and SMC lineages (Fig. 3c-e). The distribution of labeled cells between these lineages was comparable between pulse and constitutive labeling approaches. We verified co-expression of CM markers and pulse-labeled lineage tracers at the single cell level by staining cardiomyocytes dissociated from E16.5 *Wt1<sup>CreERT2/+</sup>*; *Z/Red* hearts, pulsed with tamoxifen at E10.5 and E11.5 (Fig. 3f-h). We consistently observed these pulse-labeled CMs, although the frequency ( $0.02 \pm 0.01\%$ ) was notably less than with constitutive labeling with *Wt1<sup>GFP<sup>Cre</sup></sup>*. Within the window of the tamoxifen pulse, cardiac *Wt1* expression was confined to the epicardium (Fig. 1). Based on these data, we conclude that a subset of epicardial cells expressing *Wt1* differentiate into CMs.

To further delimit the location of *Wt1*-expressing cells that differentiate into CMs, we microdissected E11.5 *Wt1<sup>GFP<sup>Cre/+</sup></sup>*; *Z/Red* fetal hearts. Serial enzymatic digestion of intact hearts yielded epicardial cells preferentially in the early fractions, due to their location on the exterior of the heart. Early (epicardial) and late (negative control) digestion fractions were sorted for GFP fluorescence, yielding a population enriched for active GFP expression (Fig. 3i). These GFP<sup>+</sup> cells were plated on either mitotically inactivated cardiac feeders or untreated tissue culture dishes (Fig. 3j-k). In both conditions a subset of the sorted *Wt1<sup>GFP<sup>Cre/+</sup></sup>*; *Z/Red* cells differentiated into CMs, identified by co-expression of the RFP lineage tracer and CM markers (Fig. 3j-k). These data provide further evidence that a subset of heart cells actively expressing *Wt1*, confined within epicardium at E10-11.5 (Fig. 1), differentiated into cardiomyocytes.

We obtained additional independent evidence that epicardial cells differentiate into cardiomyocytes by selective dye labeling of epicardium in E11.5 explanted hearts (Fig. 3l). Explanted hearts were briefly incubated in culture media containing the dye CMFDA and then place in culture media without dye. This resulted in selective labeling of epicardium, as confirmed in sections of hearts fixed immediately after CMFDA incubation (Fig. 3l, 0 hours). After two days of explant culture, labeled epicardial cells were found within myocardium, and a subset expressed the CM markers *Nkx2-5* (Fig. 3l, 48 hours). Presence of dye and the CM markers *Nkx2-5*, *Tnnt2*, and *Actn1* within the same cell was further demonstrated in single cells isolated by dissociating heart explants two days after labeling (Fig. 3m). Collectively, these data indicate that precursors actively expressing *Wt1* within E10.5-E11.5 epicardium differentiate into CMs.

Reported cardiac precursors derive from multipotent *Isl1<sup>+</sup>/Nkx2-5<sup>+</sup>* progenitors<sup>1-4</sup>. We used Cre-based lineage tracing to ask if *Wt1<sup>+</sup>* PE cells are related to these progenitors, or represent a different cardiogenic lineage. Using an *Nkx2-5<sup>IRE5-Cre</sup>* knockin allele<sup>18</sup>, we found that *Nkx2-5*-driven Cre activated *Rosa26<sup>fsLz</sup>* in a subset of PE cells (Fig. 4a-b), suggesting descent of *Wt1<sup>+</sup>* PE cells from *Nkx2-5*-expressing cells.

We independently corroborated this result using a different *Nkx2-5* knockin allele, *Nkx2-5<sup>Cre</sup>*<sup>19</sup>, and a novel Cre-activated reporter, *Gata4<sup>flap</sup>*. *Gata4* is expressed in CM, SMC, and EC compartments of the myocardium, as well as in PE (Suppl. Fig. 9)<sup>20-22</sup>. Therefore, within this domain endogenous *Gata4* regulatory elements can be used to drive expression of a Cre-dependent reporter gene, alkaline phosphatase (AP). We generated such a reporter, *Gata4<sup>flap</sup>* (Suppl. Fig. 10). In the absence of Cre, *Gata4<sup>flap</sup>* did not express AP (Suppl. Fig. 10). In the presence of well-characterized Cre transgenes, *Gata4<sup>flap</sup>* expressed AP in patterns consistent with the expected sites of Cre activity (cardiac troponin Cre (*cTNTCre*) and myosin heavy chain  $\alpha$ -Cre (*MHC $\alpha$ Cre*), myocardium; or *Tie2Cre*, endothelium; Suppl. Fig. 10). Quantitative analysis revealed that *MHC $\alpha$ Cre* activated *Gata4<sup>flap</sup>* in a greater percentage of cardiomyocytes than *Rosa26<sup>fsLz</sup>* (*Gata4<sup>flap</sup>* 93  $\pm$  3%, versus *Rosa26<sup>fsLz</sup>* 72  $\pm$  6%,  $p < 0.005$ ,  $n=4$ ), suggesting that *Gata4<sup>flap</sup>* has greater sensitivity to Cre recombination than *Rosa26<sup>fsLz</sup>*. Therefore, we asked if *Gata4<sup>flap</sup>* would show a greater contribution of *Nkx2-5<sup>+</sup>* cells to PE than suggested by *Nkx2-5<sup>IRE5-Cre</sup>*; *Rosa26<sup>fsLz</sup>*.

*Gata4<sup>flap</sup>* demonstrated a robust contribution of *Nkx2-5*-expressing progenitors to PE (asterisk, Fig. 4c). The *Nkx2-5<sup>Cre</sup>*-labeled PE cells expressed *Wt1* (Fig. 4d), indicating that *Wt1<sup>+</sup>* cells in PE are derived from *Nkx2-5*-expressing precursors. *Gata4<sup>flap</sup>* also showed a robust contribution of *Isl1*-expressing precursors to the *Wt1<sup>+</sup>* cells in PE (Suppl. Fig. 11b). Supporting this finding, at E8.0 *Wt1* and *Isl1* were expressed in adjacent regions, and a subset of cells were positive for both markers (Suppl. Fig. 12a-b).

While *Wt1*<sup>+</sup> cells in PE were labeled by *Nkx2-5*-driven Cre, they did not actively co-express *Nkx2-5* at E9.5 (Fig. 4e). In E8.0 embryos, *Nkx2-5* and *Wt1* were expressed in adjacent cells, but were not co-expressed (Suppl. Fig. 12c), suggesting that *Nkx2-5* and *Wt1* are expressed sequentially, or transiently co-expressed. To further investigate the relationship of *Nkx2-5* and *Wt1* expression, we studied the expression of *Wt1* in *Nkx2-5*<sup>+</sup> cells during embryoid body differentiation of embryonic stem (ES) cells. Using transgenic *Nkx2-5<sup>gfp</sup>* ES cells<sup>4</sup> and FACS, we isolated *Nkx2-5*-expressing cells at several time points during embryoid body differentiation. *Wt1* was transiently upregulated in *Nkx2-5*<sup>+</sup> cells during ES cell differentiation (Fig. 4f). This result was specific, as we did not detect significant *Wt1* expression in parallel experiments with *Mef2c-AHF-GFP* ES cells<sup>23</sup> (data not shown). In embryos, the activity domain of the *Mef2c-AHF* enhancer (on in anterior heart field, off in PE) did not overlap with the *Wt1* expression domain<sup>24</sup> (yellow arrowhead, Fig. 4e). Collectively, these data suggest that *Nkx2-5* and *Wt1* are sequentially expressed, or transiently co-expressed, in a subset of PE precursors.

We have shown that *Wt1*<sup>+</sup> PE/epicardial cells contribute to the CM lineage during normal heart development (Suppl. Fig. 1). *Wt1*<sup>+</sup> cells located on the heart at E10.5-E11.5 differentiate into functional CMs. Although differentiation of PE cells into CMs was previously noted in vitro<sup>25</sup>, prior fate-mapping studies of PE cells, using retroviral labels in chick or transgene labels in mice, did not describe PE contribution to the CM lineage in vivo<sup>8-10,12,13</sup>. This may be attributable to differences in methodology, species, or domains of transgene activity. Consistent with the capacity of *Wt1*-expressing cells to differentiate into CMs, *Wt1*<sup>+</sup> PE/epicardial cells are derived from progenitors that express *Nkx2-5* and *Isl1*, suggesting that they share a common developmental origin with previously described multipotent cardiogenic progenitors<sup>1,2</sup>. These experiments identify a previously unrecognized CM progenitor population in the developing heart that may be of use for cardiac regeneration or repair.

## Methods Summary

Gene targeting and mouse lines are described in the Full Methods. Fetal hearts were dissociated by serial digestion with collagenase and trypsin. For calcium imaging, dissociated fetal heart cultures were loaded with Fluo-4 AM and imaged with an Olympus FV1000 confocal microscope. Cardiac feeders were prepared and mitotically inactivated as described<sup>26</sup>. *Nkx2-5<sup>gfp</sup>* and *Mef2c-AHF-GFP* ES cells were differentiated in embryoid body culture and sorted for GFP fluorescence as described previously<sup>4,23</sup>. Immunohistochemistry was performed according to standard methods, using primary and secondary antibodies listed in Supplementary Table 1.

## Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

## Acknowledgements

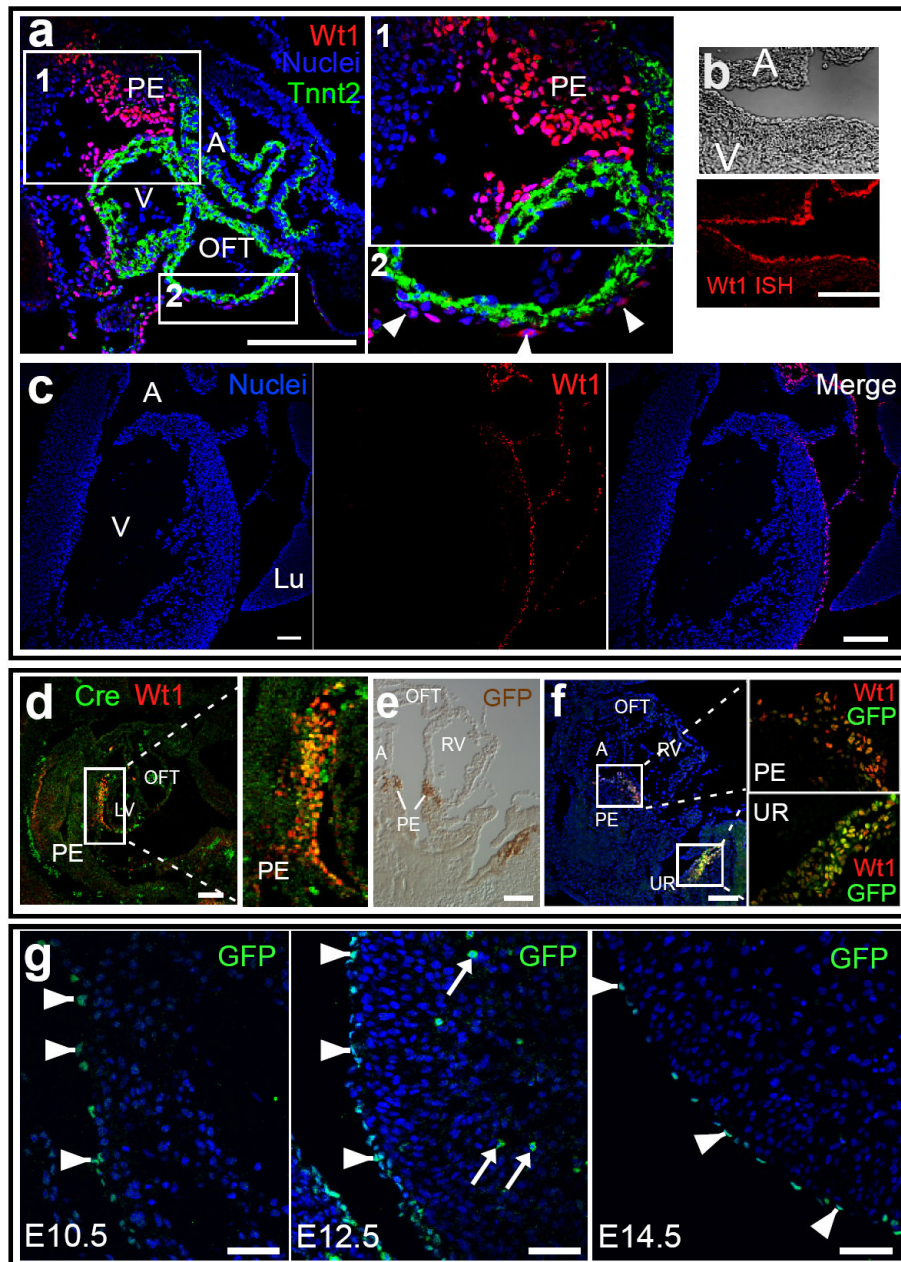
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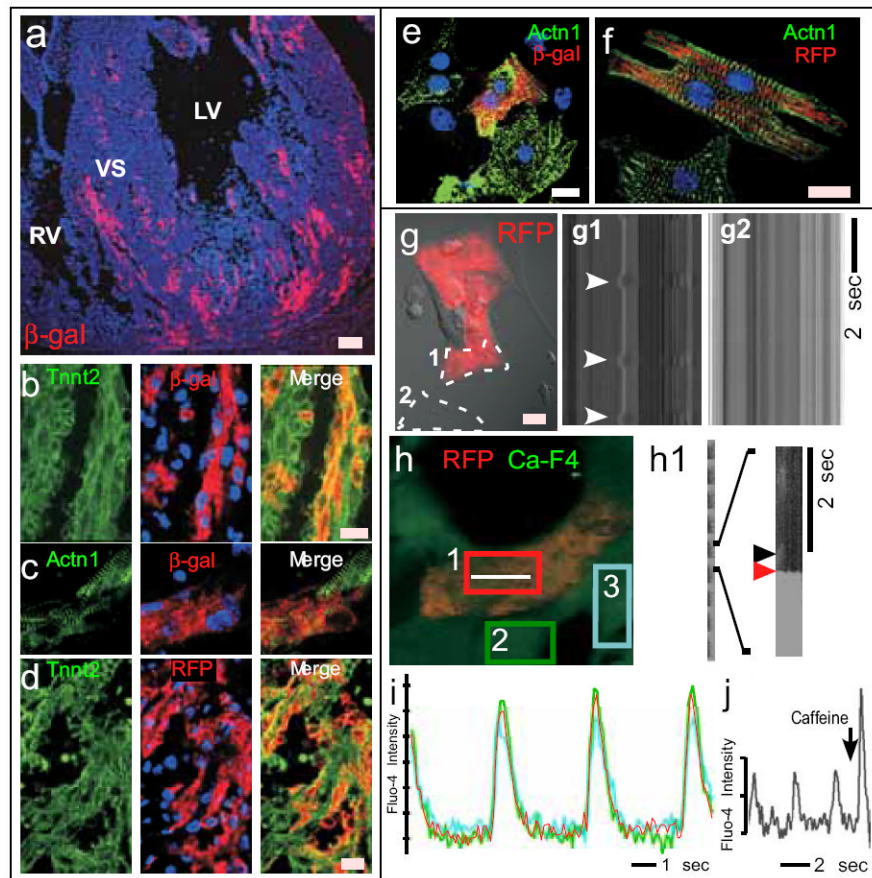
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**Fig. 1. Cardiac *Wt1* and *Wt1*-driven *GFPCre* expression**

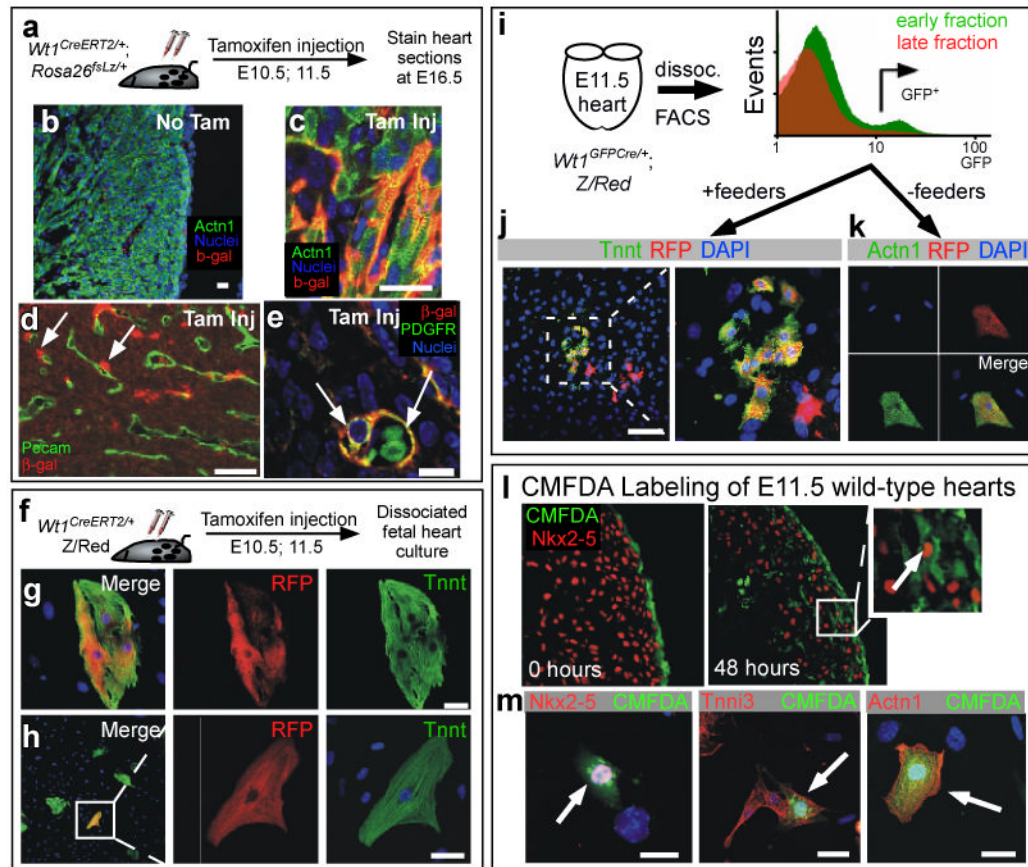
**a.** At E9.5, *Wt1* was expressed in proepicardium (PE) and in scattered pericardial cells over the surface of the heart (arrowheads, a2). **b-c.** At E15.5, *Wt1* expression was confined to the epicardium, as determined by in situ hybridization (ISH, **b**) and immunohistochemistry. **d-f.** Co-expression of *Wt1* and *GFPCre* expression in *Wt1*<sup>*GFPCre*+</sup> E9.5 embryos. *Wt1* and *GFPCre* were co-expressed in PE and urogenital ridge (UR). **g.** *Wt1*-driven *GFPCre* was confined to epicardium (arrowheads), and not detected within myocardium, at E9.5-E15.5 (representative images for E10.5, E12.5, and E14.5 are shown). Arrows indicate autofluorescent red blood cells. All images except **b** show immunohistochemical staining. A, atrium. V, ventricle. LV and RV, left and right ventricle. Lu, lung. OFT, outflow tract. Scale bars: 50  $\mu$ m.



### Fig. 2. *Wt1*-derived cells differentiate into cardiomyocytes

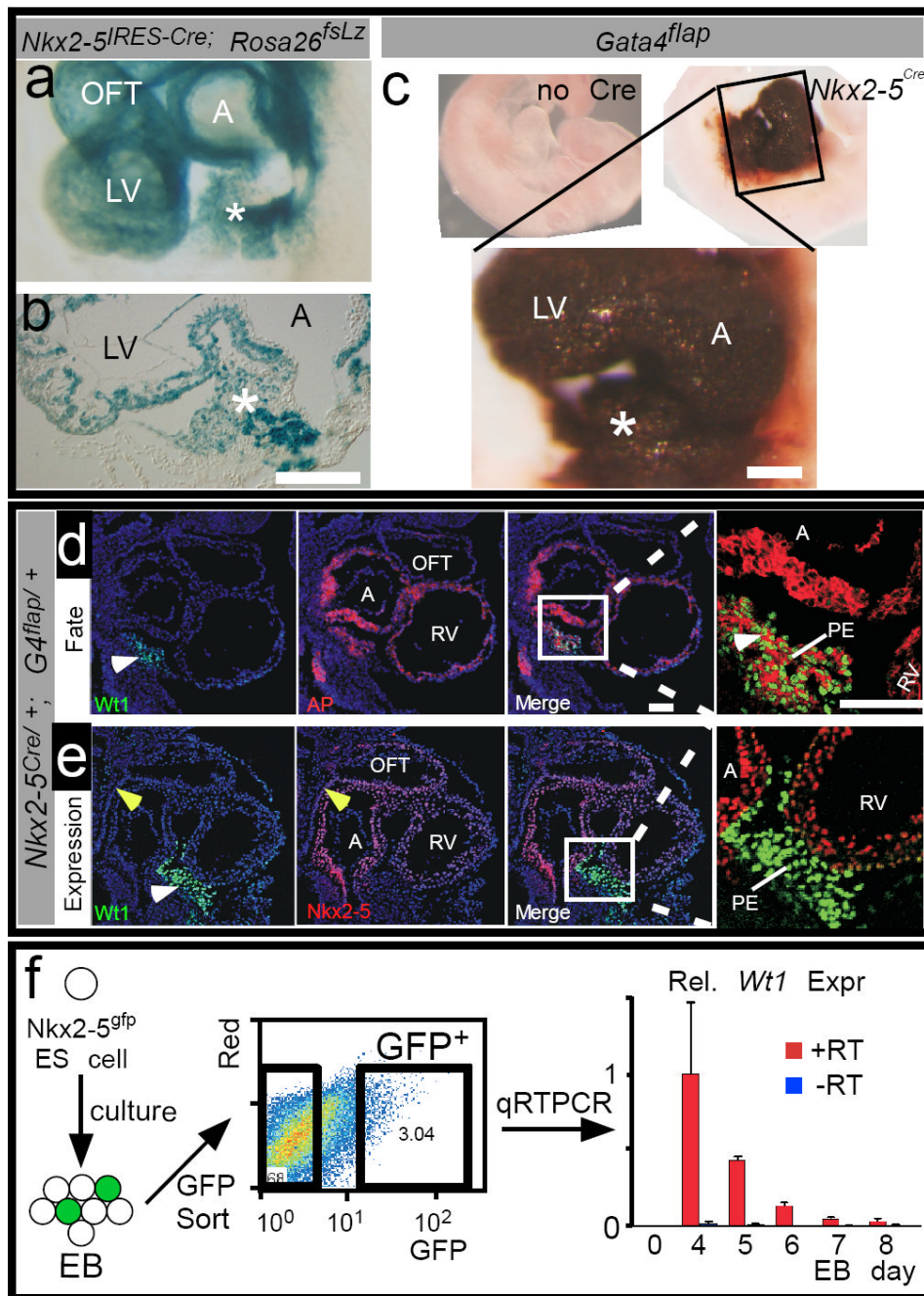
The fate of *Wt1*-derived cells, marked by  $\beta$ -gal expression in *Wt1*<sup>GFP*Cre*+</sup>; *Rosa26*<sup>fs*Lz*</sup> hearts (a-c,e), or RFP expression in *Wt1*<sup>GFP*Cre*+</sup>; *Z/Red* hearts (d,f,g-j), was analyzed by immunohistochemistry (a-f) and live cell imaging (g-j). **a.** Overview of contribution of *Wt1*-derived cells to myocardium. **b-d.** Cryosections of E15.5 heart stained for genetic lineage tracers ( $\beta$ -gal or RFP) and cardiomyocyte-specific markers *Tnt2* and *Actn1*. **e-f.** Co-expression of genetic lineage tracers with cardiomyocyte markers in dissociated E15.5 heart culture. **g-j.** Live cell imaging of *Wt1*-derived cells, identified by RFP fluorescence. **g.** Spontaneous contraction of an RFP<sup>+</sup> cells. Transmission line scan along the paths indicated by dashed lines. Arrowheads indicate contraction of an RFP<sup>+</sup> cell (**g1**). A non-contractile cell is shown in **g2**. **h.** Fluo-4 AM calcium imaging of a cluster of beating cells, one of which was RFP<sup>+</sup>. Calcium oscillations in the RFP<sup>+</sup> cell, measured by line scan along the white line, showed calcium sparks (black arrowhead, **h1**) preceding calcium waves (red arrowhead). **i.** Calcium oscillations in the RFP<sup>+</sup> cell was synchronized with adjacent cardiomyocytes, indicating electrical coupling. Plots correspond to scans within colored boxes shown in **h**. **j.** Caffeine augmentation of the amplitude of calcium oscillations in an RFP<sup>+</sup> cell. Blue staining, a-f, DAPI. Scale bars: **a**, 50  $\mu$ m, others 10  $\mu$ m.





### Fig. 3. *Wt1*-expressing epicardial cells differentiate into cardiomyocytes

**a-h.** Temporal restriction of Cre labeling with *Wt1<sup>CreERT2</sup>*. *Wt1<sup>CreERT2</sup>* did not recombine the fate mapping reporter *Rosa26<sup>fsLz</sup>* in the absence of tamoxifen (**b**). Tamoxifen treatment at E10.5 and E11.5 induced *Wt1<sup>CreERT2</sup>* labeling of cardiomyocytes, as determined in both tissue sections (**c**) and in dissociated heart culture (**g-h**). Also labeled were SMCs adjacent to endothelial tubes (**d-e**). **i-k.** E11.5 *Wt1<sup>GFP<sup>Cre/+</sup></sup>* heart cells actively expressing *Wt1*, as determined by GFP fluorescence, differentiated into cardiomyocytes. *Wt1<sup>+</sup>* epicardial cells were enriched in early digestion fractions compared to late digestion fractions. GFP<sup>+</sup> cells from early fractions were plated either with mitotically inactivated feeders (**j**) or without feeders (**k**). **l-m.** CMFDA dye, selectively incorporated into E11.5 epicardium, was found in CMs at 48 hours. Brief incubation of E11.5 heart explants with the dye CMFDA resulted in selective labeling of epicardium (0 hours culture). After 48 hours in explant culture, dye labeled cells were present in the myocardial wall. A subset of dye-labeled cells co-expressed the CM markers Nkx2-5, cardiac troponin I (Tnni3), and Actin1. Arrows indicate co-expression. Scale bars: 10  $\mu$ m.



**Fig. 4. Proepicardium arises from *Nkx2-5*<sup>+</sup> precursors**

**a-c.** The *Nkx2-5* fate map was determined in *Nkx2-5<sup>IRES-Cre</sup>; Rosa26<sup>flSLZ</sup>* (**a-b**) or *Nkx2-5<sup>Cre</sup>; Gata4<sup>flap</sup>* (**c**) embryos. *Nkx2-5*-driven Cre activated lineage tracer expression in PE (asterisk). **d.** *Nkx2-5* fate, marked by expression of alkaline phosphatase (AP, red membrane staining) compared to *Wt1* immunohistochemistry (green nuclear staining). *Wt1* was expressed in *Nkx2-5*-derived cells in PE (white arrowhead). **e.** Immunohistochemistry comparing *Nkx2-5* (red) and *Wt1* expression (green). Yellow arrowhead indicates anterior heart field, which did not express *Wt1*. **f.** *Nkx2-5<sup>GFP</sup>* ES cells were differentiated in embryoid body (EB) culture for the indicated number of days. The cells were then dissociated, and GFP<sup>+</sup> cells were isolated by flow cytometry. *Wt1* expression in GFP<sup>+</sup> cells was measured by quantitative reverse

transcription PCR. Bars indicate mean  $\pm$  sd. n=3. Scale bars: 50  $\mu$ m. A, atrium. RV, right ventricle. LV, left ventricle. OFT, outflow tract.