An FGF-driven feed-forward circuit for spatiotemporal patterning of the cardiopharyngeal mesoderm in a simple chordate

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Abstract

In embryos, pluripotent stem cells and multipotent progenitors must divide and produce distinct progeny to express their full developmental potential. In vertebrates, mounting evidence point to the existence of multipotent cardiopharyngeal progenitors that produce second-heart-field-derived cardiomyocytes, and branchiomeric skeletal head muscles. However, the cellular and molecular mechanisms underlying these early fate choices remain largely elusive. The tunicate Ciona has emerged as an attractive model to study early cardiopharyngeal development at high spatial and temporal resolution: through two asymmetric and oriented cell divisions, defined multipotent cardiopharyngeal progenitors produce distinct first and second heart precursors, and pharyngeal muscle (aka atrial siphon muscle, ASM) precursors. Here, we demonstrate that differential FGF/MAPK signaling distinguishes between MAPK-negative heart precursors, and MAPK-positive multipotent progenitors and ASM precursors. We characterize an FGF/MAPK-driven feed-forward circuit that promotes the successive activations of essential cardiopharyngeal determinants, Tbx1/10 and Ebf. Finally, we show that coupling FGF/MAPK restriction and cardiopharyngeal network deployment with cell divisions permits the emergence of diverse cell types from common multipotent progenitors.
Introduction

In the past few years, studies guided by developmental genetics knowledge progressed towards driving mammalian stem cells into forming pure cultures of selected cell types in vitro (e.g. Kattman et al., 2011; Mazzoni et al., 2011; Peljto and Wichterle, 2011). By contrast, in their embryonic context, pluripotent cells generate diverse cell types in defined proportions. This simple observation implies that pluripotent stem cells and multipotent embryonic progenitors must divide before individual cells among their progeny adopt distinct fates, as a result of differential exposure to inducing signals and/or inheritance of cell autonomous determinants.

Subsets of the heart and head/neck myocytes recently emerged as related derivatives of multipotent progenitors located in the mesodermal cardiopharyngeal field (Diogo et al., 2015; Tzahor, 2009; Tzahor and Evans, 2011). Specifically, early lineage tracing, transplantations and controlled explant culture experiments demonstrated that the anterior splanchnic/pharyngeal mesoderm of amniote embryos can produce either skeletal muscles or heart tissue, depending upon exposure to growth factors and signaling molecules (Nathan et al., 2008; Tirosh-Finkel et al., 2006; Tzahor et al., 2003; Tzahor and Lassar, 2001). Clonal analyses in the mouse further revealed the existence of common Mesp1-expressing progenitors for subsets of the second heart field-derived cardiomyocytes and branchiomeric facial, jaw, neck and even esophageal muscles (Gopalakrishnan et al., 2015; Lescroart et al., 2014; Lescroart et al., 2015; Lescroart et al., 2010; Lescroart et al., 2012). In vitro studies using pluripotent stem cells indicated that controlled Mesp1 expression can drive mesodermal progenitors towards cardiac and/or skeletal muscle fates (Bondue et al., 2008; Chan et al., 2016; Chan et al., 2013). Genetic labeling and functional studies showed that proper development of the pharyngeal apparatus and second heart field derivatives require shared inputs from Tbx1, Nkx2-5 and Islet1 transcription factors (e.g. (Cai et al., 2003; George et al., 2015;
Taken together, a growing body of evidence point to the existence of a mesodermal field of multipotent progenitors capable of producing either SHF-derived cardiomyocytes or branchiomeric skeletal muscles in early amniote embryos (Diogo et al., 2015; Mandal et al., 2017). However, the mechanisms that distinguish fate-restricted heart and head muscle precursors remain largely elusive.

The tunicate *Ciona*, which is among the closest living relatives to the vertebrates (Delsuc et al., 2006; Putnam et al., 2008), has emerged as a simple chordate model to characterize multipotent cardiopharyngeal progenitors and the mechanisms that initiate heart vs. pharyngeal muscle fate choices (Kaplan et al., 2015; Razy-Krajka et al., 2014; Stolfi et al., 2010; Tolkin and Christiaen, 2016; Wang et al., 2013). *Ciona* tailbud embryos possess two multipotent cardiopharyngeal progenitors on either side. Like their vertebrate counterparts, these cells emerge from *Mesp*+ progenitors towards the end of gastrulation; they are induced by FGF/MAPK signaling and have been termed *trunk ventral cells* (aka TVCs; (Christiaen et al., 2008; Davidson and Levine, 2003; Davidson et al., 2006; Davidson et al., 2005; Satou et al., 2004; Stolfi et al., 2010)). TVCs activate conserved cardiac markers, including *Hand*, *Gata4/5/6* and *Nk4/Nkx2-5*, and migrate as bilateral polarized pairs of cells, until the left and right pairs meet at the ventral midline and begin to divide asymmetrically along the mediolateral axis (Figure 1A; (Christiaen et al., 2008; Davidson et al., 2005; Satou et al., 2004; Stolfi et al., 2010)).

The first oriented asymmetric divisions produce small median first heart precursors (FHPs), and large lateral second trunk ventral cells (STVCs), which specifically activate *Tbx1/10* expression (Davidson et al., 2005; Stolfi et al., 2010; Wang et al., 2013). STVCs later divide again to produce small median second heart precursors (SHPs), and large
lateral atrial siphon muscle founder cells (ASMFs), which activate Ebf (aka COE; (Razy-Krajka et al., 2014; Stolfi et al., 2010; Stolfi et al., 2014c)). The transcription factors Hand-related (Hand-r)/Notrlc, which is expressed in the TVCs and maintained in the STVCs and ASMFs after each division, and Tbx1/10 are required for Ebf activation in the ASMFs, whereas Nk4/Nkx2-5 represses Tbx1/10 and Ebf expression in the second heart precursors (SHPs)(Razy-Krajka et al., 2014; Tolkin and Christiaen, 2016; Wang et al., 2013). Conversely, Tbx1/10 and Ebf inhibit cardiac markers, and likely determinants, such as Gata4/5/6 and Hand (Razy-Krajka et al., 2014; Stolfi et al., 2010; Stolfi et al., 2014a; Wang et al., 2013). These regulatory cross-antagonisms presumably underlie the transition from transcriptionally primed multipotent progenitors to separate fate-restricted precursors, by limiting the deployment of the heart- and pharyngeal-muscle-specific programs to their corresponding specific precursors (Kaplan et al., 2015).

Here, we identify regulatory mechanisms ensuring the emergence of diverse fate-restricted precursors from multipotent progenitors. We show that differential FGF/MAPK signaling, feed-forward regulatory mechanisms and coupling with the cell cycle control the spatially restricted activation of Tbx1/10 and Ebf, successively, thus permitting the emergence of both first and second heart precursors, and ASM/pharyngeal muscle precursors from common multipotent progenitors.
Results

MAPK signaling is active in the multipotent cardiopharyngeal progenitors and progressively restricted to the pharyngeal muscle precursors.

During the earliest stages of cardiopharyngeal development in ascidians, multipotent progenitors co-express early regulators of both the heart and ASM programs, a phenomenon referred to as multilineage transcriptional priming, (Razy-Krajka et al., 2014; Stolfi et al., 2014b). Subsequent regulatory cross-antagonisms lead to the segregation of these distinct cardiopharyngeal programs to their corresponding fate-restricted progenitors (Stolfi et al., 2010; Wang et al., 2013); reviewed in (Kaplan et al., 2015)). ASM-specific expression of Ebf is necessary and sufficient to terminate the heart program and impose a pharyngeal muscle fate (Razy-Krajka et al., 2014; Stolfi et al., 2010). Antagonistic Tbx1/10 and Nk4 activities determine ASM-specific Ebf activation (Wang et al., 2013); however, the symmetry-breaking events leading to cardiopharyngeal mesoderm patterning and ASM-specific expression of Ebf remain unknown. In particular, we surmised that differential signaling inputs determine the stereotyped spatio-temporal patterning of early cardiopharyngeal progenitors.

The Ciona homologs of specific FGF/MAPK pathway components, including FGF receptor substrate 2/3 (Frs2/3; (Gotoh et al., 2004)), Ets.b, and Fgf4/5/6, are preferentially expressed in the TVCs, in the STVCs and in the ASMFs as cells transition from a multipotent progenitor state to distinct heart vs. ASM fate-restricted precursors (Razy-Krajka et al., 2014). This patterned expression of MAPK effector genes prompted us to evaluate a role for FGF/MAPK pathway in cardiopharyngeal fate decisions.

We first used an antibody specific to the dual phosphorylated form of Extracellular Regulated Kinase (dpERK) to monitor Mitogen Activated Protein Kinase (MAPK) activity in the cardiopharyngeal mesoderm. We detected dpERK staining in the newly
born TVCs, marked by the B7.5-lineage-specific Mesp>H2B::mCherry transgene, as previously observed (Davidson et al, 2006). We also detected weaker but persistent dpERK staining in the TVCs during migration (Figs. 1 and S1). Following the first and second asymmetric divisions of the TVCs and STVCs, dpERK staining was successively restricted to the more lateral STVCs and ASMFs, respectively (Figures 1A, B; S1).

**The canonical FGF/Ras/MEK/ERK pathway is necessary and sufficient to promote pharyngeal muscle specification in the cardiopharyngeal lineage.**

This exclusion of MAPK activity from the medial first and second heart precursors opened the possibility that differential ERK activity is required for proper STVC and ASMF vs. heart precursors fate decisions. In *Ciona*, signaling through the sole FGF receptor (FGFR) governs ERK activity in several developmental processes, including neural induction (Bertrand et al., 2003; Hudson et al., 2003) and central nervous system patterning (Haupaix et al., 2014; Racioppi et al., 2014; Stolfi et al., 2011; Wagner et al., 2014), early endomesoderm and notochord fate specification (Imai et al., 2002; Picco et al., 2007; Shi and Levine, 2008; Shi et al., 2009; Yasuo and Hudson, 2007). Notably, FGF/MAPK signaling is active in the only Mesp+ cardiogenic B7.5 blastomeres (Imai et al., 2006; Shi and Levine, 2008), where targeted misexpression of a dominant negative form of FGFR (dnFGFR) using a B7.5-lineage-specific Mesp driver blocks TVC induction (Davidson et al., 2006). We used a TVC-specific FoxF enhancer (*FoxF(TVC):bpFOG-1*dnFGFR::mCherry, hereafter called *FoxF>dnFGFR*; (Beh et al., 2007)), to bypass early effects and achieve later misexpression of dnFGFR in the TVCs and their progeny. *FoxF>dnFGFR* prevented neither TVC migration nor asymmetric divisions, but it abolished the expression of both *Tbx1/10* in the STVCs and *Ebf* in the ASMFs (Figure 1C). This data indicate that FGF/MAPK signaling is required in the cardiopharyngeal
progenitors and/or their progeny for ASM fate specification, beyond the initial TVC induction.

Upon FGF/MAPK-dependent induction, the TVCs express *Hand-related/Hand-r* (renamed after *Notrlc/Hand-like*; (Christiaen et al., 2008; Davidson and Levine, 2003; Davidson et al., 2006; Satou et al., 2004; Stolfi et al., 2014c; Woznica et al., 2012)), which encodes a basic helix-loop-helix (bHLH) transcription factor necessary for *Ebf* expression in the ASMFs (Razy-Krajka et al., 2014). Moreover, the *Hand-r* TVC enhancer contains putative Ets1/2 binding sites, which are necessary for reporter gene expression, and presumably mediate the transcriptional inputs of FGF/MAPK (Woznica et al., 2012).

Since *Hand-r* and *FoxF* expressions start at approximately the same time in newborn TVCs, we used FoxF>dnFGFR to test whether the maintenance of *Hand-r* expression in migratory TVCs requires prolonged FGF/MAPK inputs after initial TVC induction. FoxF>dnFGFR inhibited *Hand-r* expression in late TVCs (Figure 1C), indicating that sustained *Hand-r* expression requires continuous FGF/MAPK signaling.

To test whether the spatial restriction of MAPK activity explains the patterned expressions of *Hand-r*, *Tbx1/10* and *Ebf* following asymmetric cell divisions, we used gain-of-function perturbations to force FGF/MAPK activity throughout the cardiopharyngeal mesoderm and assayed gene expression (Figure 2). We focused on the canonical FGF/MAPK pathway where signal transduction involves Ras, Raf, MEK and ERK downstream of FGFR and upstream of transcriptional effectors (Lemmon and Schlessinger, 2010). We first used M-RasG22V, a defined constitutively active form of M-Ras, which mediates FGF signaling in *Ciona*, where other classical *Ras* genes are missing (Keduka et al., 2009). To assay the transcriptional consequences of forced M-Ras activity in the cardiopharyngeal lineage, we first focus on *Htr7* and *Tbx1/10* expression following the first asymmetric TVC division in 15 hours post-fertilization (hpf) embryos. *Htr7* encodes a trans-membrane G-protein coupled receptor and, like *Hand-r*, its expression
and maintenance in the TVCs require MAPK activity (Figure S2; (Razy-Krajka et al., 2014)), and become restricted to the lateral STVC following asymmetric division. However, \(Htr7\) mRNAs appear to be cleared more rapidly from the FHPs, making the patterned expression easier to analyze than that of \(Hand-r\) (Figures 2 and 3D; (Razy-Krajka et al., 2014)). Importantly, misexpression of M-Ras\(^{G22V}\) using the TVC-specific \(FoxF\) enhancer did not alter the cell division patterns, allowing us to identify large lateral STVCs and small median FHPs. Compared to control embryos overexpressing wild-type M-Ras (M-Ras\(^{WT}\)), TVC-specific gain of M-Ras function caused both persistent \(Htr7\) expression and ectopic activation of \(Tbx1/10\) in the first heart precursors following asymmetric divisions. Similarly, \(FoxF>\)M-Ras\(^{G22V}\)-expressing 18hpf larvae displayed ectopic \(Ebf\) activation throughout the cardiopharyngeal mesoderm (Figure 2B, C). These results indicated that forced M-Ras activation throughout the cardiopharyngeal lineage is sufficient to ectopically activate STVC and ASMF markers. This is consistent with the idea that spatially defined signaling upstream of M-Ras restricts MAPK activity, thus localizing STVC- and ASM-specific gene activities.

To further probe the signal transduction pathway, we engineered a constitutively active version of the \(Ciona\) Mek1/2 protein by introducing phosphomimetic mutations of two conserved serine residues in the catalytic domain, as previously shown for the mammalian homolog (Cowley et al., 1994; Mansour et al., 1994). Early misexpression of this Mek\(^{S220E,S216D}\) construct in the B7.5 blastomeres using a \(Mesp\) enhancer caused ectopic TVC induction, mimicking the effects of published gain of Ets1/2 function (Figure S3; (Davidson et al., 2006)). Mirroring the effects of M-Ras\(^{G22V}\) gain-of-function experiments, TVC-specific misexpression of Mek\(^{S220E,S216D}\) using the \(FoxF\) enhancer also caused ectopic expression of \(Htr7\) and \(Tbx1/10\), and \(Ebf\) in 15 and 18hpf larvae, respectively (Figure 2B, C). Taken together, these results indicate that activity of the canonical FGF-Ras-MEK-ERK pathway is progressively restricted to the STVC and
ASMF, and is both necessary and sufficient to promote STVC- and ASMF-specific gene expressions.

Continuous FGF/MAPK activity is required for the successive activations of Tbx1/10 and Ebf.

FGF/MAPK signaling is sufficient and necessary to maintain Hand-r expression in late TVCs (Figure 1), and Hand-r is necessary for Ebf expression in the ASMF (Razy-Krajka et al., 2014). Therefore, it is possible that later FGF/MAPK signaling is dispensable for Tbx1/10 and Ebf activation and ASM specification, as long as STVC and ASMF cells inherit sustained levels of Hand-r mRNAs and/or proteins. To disentangle late from early requirements of FGF/MAPK signaling for TVC progeny specification, we incubated embryos at different stages with the MEK/Mapkk inhibitor U0126, which abolishes dual ERK phosphorylation and the initial MAPK-dependent TVC induction in Ciona embryos (Figure S1; (Davidson et al., 2006; Hudson et al., 2003)). MEK inhibition during TVC migration (i.e. between 9.5 and 12.5 hpf, Figure 3A) blocked the expression of Hand-r and Htr7 in late TVCs (Figure 3B, E). Similarly, U0126 treatments in late TVCs, and through the first asymmetric division (i.e. between 12 and 15 hpf, Figure 3A) blocked both the maintenance of Hand-r and Htr7, and the activation of Tbx1/10 in the STVCs (Figure 3C, D, F, G). Finally, MEK inhibition in late STVCs and through asymmetric divisions (i.e. between 15 and 18 hpf) blocked the ASMF-specific expression of Ebf (Figure 3H). These results indicate that continuous MEK activity is required throughout cardiopharyngeal development to successively activate TVC-, STVC-, and ASMF-expressed genes.

Since Ebf expression is maintained for several days in the ASMF derivatives as they differentiate into body wall and siphon muscles (Razy-Krajka et al., 2014), we tested whether continued MEK activity is also required for the maintenance of Ebf expression
past its initial onset and cells' commitment to an ASM fate. Using both regular and intron-specific antisense probes, which specifically detect nascent transcripts (Wang et al., 2013), we showed that later MEK inhibition (i.e. U0126 incubation between 17 and 20 hpf) did not block the maintenance of Ebf transcription in the ASMPs (Figure 3I, J). This indicates that sustained MEK activity is required until the onset of Ebf expression, but not beyond, the maintenance of Ebf expression during ASM development is independent of MAPK.

Since U0126 treatments affect the whole embryo, we sought to further confirm the later roles for FGF/MAPK signaling specifically in the cardiopharyngeal mesoderm. To this aim, we used an STVC-specific enhancer from the Tbx1/10 locus (termed T12; Figure 3K, L; (Tolkin and Christiaen, 2016); Racioppi et al., in preparation) to drive expression of either dnFGFR or the constitutively active M-Ras$^{G22V}$ starting at ~14hpf, and assayed Ebf expression at 18hpf (Figure 3K, L). These perturbations minimally affected the cell division patterns, such that cells corresponding to FHP, SHP and ASMF could be identified by their position relative to the midline (Figure 3K). M-Ras$^{G22V}$ misexpression caused conspicuous ectopic Ebf expression in the SHPs, whereas dnFGFR-mediated inhibition of MAPK activity blocked Ebf activation in the lateral ASMFs. These results support the notion that localized FGF/MAPK activity is necessary and sufficient for ASMF-specific expression of Ebf.

**Coherent feed-forward circuits for cardiopharyngeal mesoderm patterning and ASM fate specification.**

The above results indicate that Hand-$r$, Tbx1/10 and Ebf require ongoing FGF/MAPK activity for their successive activations in the TVCs, STVCs and ASMFs, respectively. We previously showed that RNAi and/or CRISPR-mediated inhibition of
either Hand-r or Tbx1/10 function blocks Ebf activation in the ASMFs, where both
Hand-r and Tbx1/10 expressions are maintained (Razy-Krajka et al., 2014; Tolkin and
Christiaen, 2016; Wang et al., 2013). Therefore, observations such as the loss of Ebf
expression upon FoxF>dνFGFR electroporation could be due to an early loss of Hand-r
and/or Tbx1/10. We used epistasis assays to systematically test whether early regulators
mediate the effects of FGF/MAPK on later gene expression and ASM fate specification,
or whether FGF/MAPK signaling acts both upstream and in parallel to early regulators
in a more complex regulatory circuit.

We first revisited the regulatory relationships between FGF/MAPK, Hand-r and
Tbx1/10 in late TVCs and early STVCs. We validated single guide RNAs (sgRNAs) for
CRISPR/Cas9-mediated mutagenesis of Hand-r (Table S1; (Gandhi et al., 2017)), and
determined that Hand-r function is necessary for Tbx1/10 activation in the STVCs
(Figure 4A). Co-expression of a modified Hand-r cDNA containing wobble base
mutations that disrupt the sgRNA protospacer adjacent motif (PAM; Hand-rPAMmis)
rescued Tbx1/10 expression in the STVCs, indicating that Tbx1/10 down-regulation in
this CRISPR "background" is specifically due to Hand-r loss-of-function (Figure 4A). To
further probe if Hand-r activity is necessary for FGF/MAPK-dependent Tbx1/10
expression, we used gain of M-Ras function in a Hand-r CRISPR "background".
Whereas, misexpression of the constitutively active M-RasG22V caused ectopic Tbx1/10
expression, concomitant loss of Hand-r function diminished both endogenous and
ectopic Tbx1/10 expression in the STVC and FHP, respectively (Figure 4A). Although,
remaining ectopic activation could still be observed, possibly because M-RasG22V could
boost Hand-r expression in heterozygous cells where CRISPR/Cas9 disrupted only one
copy of the gene. This data indicate that Hand-r is necessary for FGF/MAPK-induced
activation of Tbx1/10.
To further probe the epistatic relationships between *Hand-r* and MAPK signaling upstream of *Tbx1/10*, we attempted to rescue *Tbx1/10* expression in U0126-treated embryos, by over-expressing *Hand-r* with the TVC-specific *FoxF* enhancer. Neither did *Hand-r* over-expression cause ectopic *Tbx1/10* activation (in the FHPs), nor was it sufficient to rescue *Tbx1/10* expression in 15hpf STVCs (Figure 4B). Taken together, these data indicate that both *Hand-r* and MAPK activities are required to activate *Tbx1/10* in the STVCs. These results also imply that MAPK signaling is restricted to the STVC independently of *Hand-r* activity, which suffice to explain the STVC-specific activation of *Tbx1/10*.

Next, we investigated the epistatic relationship between FGF/MAPK, *Hand-r*, and *Tbx1/10* upstream of *Ebf* in the ASMFs. We first used previously validated CRIPSR/Cas9 reagents targeting the *Tbx1/10* coding region (Tolkin and Christiaen, 2016), to confirm that B7.5-lineage-specific loss of *Tbx1/10* function inhibited *Ebf* activation, and verified that this effect could be rescued by over-expression of a CRISPR/Cas9-resistant *Tbx1/10* cDNA, expressed with a minimal TVC-specific *FoxF* enhancer (Figure 4C; *Tbx1/10*\textsuperscript{PAMmis}). In these rescue experiments, we observed ectopic *Ebf* activation in the SHP, as previously described when driving *Tbx1/10* expression with a TVC-specific *FoxF* enhancer (Wang et al., 2013). As explained below, this ectopic activation could be attributed to a precocious expression of *Ebf* in the STVCs (Figure 4E). To test whether *Tbx1/10* was also required for ectopic *Ebf* expression in response to MAPK activation, we combined CRISPR/Cas9-mediated *Tbx1/10* knockout with constitutive MAPK activation using the M-Ras\textsuperscript{G22V} mutant and observed a significant inhibition of both endogenous and ectopic *Ebf* expression in the 18hpf ASMF and SHP, respectively (Figure 4C). Taken together, these results show that *Tbx1/10* function is necessary for FGF/MAPK-induced expression of *Ebf* in the ASMFs.
To further test whether *Tbx1/10* acts in parallel and/or downstream of MAPK to activate *Ebf*, we combined gain of *Tbx1/10* function with perturbations of FGF/MAPK signaling and assayed *Ebf* expression. We realized that *FoxF*-driven misexpression of *Tbx1/10* caused precocious *Ebf* activation in 15hpf STVCs (Figure 4D, E). This precocious expression remained remarkably patterned, suggesting that STVC-restricted FGF/MAPK activity prevented *Ebf* expression in the dpERK-negative, small median FHPs (Figures 1B, 4E, S1). Indeed, co-expression of both *Tbx1/10* and M-Ras<sup>G22V</sup> caused both precocious and ectopic *Ebf* expression in the 15hpf medial and lateral TVC derivatives, which would be FHPs and STVCs in control embryos, respectively. This data confirms that *Tbx1/10* misexpression does not suffice to cause ectopic *Ebf* expression in the FHPs, because the latter presumably lack FGF/MAPK activity, as is the case in control embryos.

U0126-mediated MEK inhibition from 12 to 15hpf, i.e. after the onset of *FoxF>*Tbx1/10* misexpression, further confirmed that MAPK activity is required in parallel to *Tbx1/10* for precocious *Ebf* activation in 15hpf STVCs (Figure 4D, E). Taken together, these results indicate that *Tbx1/10* and MAPK are both required to activate *Ebf* in the cell cycle following that of *Tbx1/10* onset.

Since *Hand-r* expression is maintained in the ASMF, and CRISPR/Cas9- or RNAi-mediated *Hand-r* knockdown blocked both *Tbx1/10* (Figure 4A) and *Ebf* expression (Razy-Krajka et al., 2014), we reasoned that *Hand-r* could also act both upstream and in parallel to *Tbx1/10* for *Ebf* activation. To test this possibility, we assayed *Ebf* expression in 18hpf ASMF following defined perturbations of *Hand-r* and *Tbx1/10*. As expected, CRISPR/Cas9-mediated *Hand-r* mutagenesis strongly inhibited *Ebf* expression, and this effect could be rescued by a CRISPR-resistant *Hand-r* cDNA (Figure 4F). To test whether this effect was mediated by a loss of *Tbx1/10* expression, we attempted to rescue the *Hand-r* loss-of-function by over-expressing *Tbx1/10* using the *FoxF* enhancer. As
explained above, FoxF-mediated Tbx1/10 misexpression caused precocious and ectopic 
Ebf expression in larvae co-electroporated with control sgRNAs (Figure 4D, E, F). By 
contrast, combining loss of Hand-r function with Tbx1/10 misexpression inhibited both 
the endogenous and ectopic Ebf expression (Figure 4F), indicating that Hand-r is also 
required in parallel to Tbx1/10 for Ebf activation in the ASMFs.

Taken together, these analyses of the epistatic relationships between FGF/MAPK 
signaling, Hand-r, Tbx1/10 and Ebf suggest that coherent feed-forward circuits govern 
the sequential activation of Hand-r, Tbx1/10 and Ebf in response to continuous but 
progressively restricted FGF/MAPK inputs (Figure 4G), thus linking spatial patterning to 
the temporal deployment of the regulatory cascade leading to localized Ebf activation 
and pharyngeal muscle specification.

The cell cycle entrains the temporal deployment of the cardiopharyngeal 
gene regulatory network.

In principle, the feed-forward circuit described above is sufficient to explain the 
successive activations of Hand-r, Tbx1/10 and Ebf. However, Tbx1/10 and Ebf do not 
turn on until after oriented and asymmetrical divisions of the TVCs and STVCs, 
respectively. Notably, even when we misexpressed Tbx1/10 in the TVCs, Ebf was 
activated only after TVC division and in the lateral-most cells, where FGF/MAPK 
signaling is normally maintained (Figures 1B, 4E). This sequence of events -divisions 
followed by gene activation- is paramount in the cardiopharyngeal mesoderm, as it 
permits the birth of first and second heart precursors, whose fates are antagonized by 
Tbx1/10 and Ebf (Razy-Krajka et al., 2014; Stolfi et al., 2010; Wang et al., 2013). 
Therefore, we sought to investigate the role(s) of the cell cycle in controlling the timing of 
Tbx1/10 and Ebf activations.
We first evaluated the effects of cytochalasin B, a classic inhibitor of cytokinesis widely used to study cell fate specification in ascidians (Figure 5A; (Whittaker, 1973)). Treatments starting before TVC divisions (12 hpf) did not block Tbx1/10 or Ebf expression in embryos fixed after their normal onset at either 16 or 19 hpf, respectively (Figure 5B). Similarly, treatment starting between the first and second asymmetric divisions (15 hpf) did not block localized Ebf expression at 19 hpf (Figure 5B). This indicates that Tbx1/10 and Ebf activations occur by default in the absence of cytokinesis, most likely because FGF/MAPK signaling persists throughout the shared cytoplasm. This data thus illustrates how the spatial restriction of FGF/MAPK signaling, following cell divisions, leads to the localized activations of Tbx1/10 and Ebf, and permits the emergence of first and second cardiac precursors.

Cytochalasin treatments usually lead to the formation of polynucleated cells (e.g. Figure 5B, middle panel), because the cell cycle and nucleokinesis continue in these artificial endoreplicating cells. To alter cell cycle progression more comprehensively, and specifically in the cardiopharyngeal lineage, we used genetically encoded inhibitors of cell cycle transitions: Cdkn1b.a and Cdkn1b.b (also known as Noto16), the ortholog of which is a potent inhibitor of the G1/S transition in the ascidian species Halocynthia roretzi (Kuwajima et al., 2014), and Wee1, a G2/M inhibitor, as previously described (Dumollard et al., 2017). We used the TVC-specific FoxF enhancer to misexpress these negative regulators of cell cycle progression, monitored cell divisions and assayed Tbx1/10 expression at 15 hpf, when control TVCs have divided and the lateral-most STVCs normally express Tbx1/10. Each perturbation efficiently inhibited TVC divisions, such that only two cells were visible on either side of the embryos (Figure 5C). In these delayed TVCs, Tbx1/10 expression was strongly reduced compared to control STVCs (Figure 5C; compare to Figure 4A, B). However, approximately 40% of the delayed TVCs expressed Tbx1/10 to variable extents. This suggests that the cardiopharyngeal
regulatory network can qualitatively unfold independently of cell cycle progression, but the latter is necessary for Tbx1/10 expression to its wild-type levels.

We next used the STVC-specific Tbx1/10 T12 enhancer, to misexpress Cdkn1b.a, Noto16 and Wee1, and assay Ebf expression at later stages. Inhibitors of the G1/S transition failed to block STVC divisions (data not shown), most likely because T12-driven products did not accumulate quickly enough to interfere with the G1/S transition in STVCs (this cell cycle lasts only ~2 hours compared to ~6 hours for the TVC interphase), suggesting that the G1 phase is too short for T12-driven gene products to accumulate before the G1/S transition. Therefore, we focused the analyses of Ebf response to cell cycle perturbations on misexpression of the G2/M inhibitor Wee1.

Preliminary analyses of 18hpf larvae, fixed approximately 2 hours after the documented onset of Ebf expression in ASMFs (Razy-Krajka et al., 2014), indicated that Ebf can turn on in arrested STVCs that failed to divide upon Wee1 misexpression (Figure 5D).

Because ~30% of the embryos showed variable expression, as was the case for Tbx1/10 in the previous experiment, we reasoned that perturbations of the G2/M transition could alter the dynamics of Ebf upregulation. We investigated this possibility using embryos fixed every 30 minutes between 15.5hpf and 18hpf, when cells transition from a late Tbx1/10+; Ebf- STVC state to a committed Ebf+, Mrf+ ASMF state (Razy-Krajka et al., 2014; Wang et al., 2013). First, we observed that the proportion of embryos with conspicuous ASMFs increased from ~20% to >90% between 15.5 and 16.5 hpf in control embryos (Figure 4E). By contrast, Wee1-expressing cells had divided in only ~35% of the embryos by 16.5hpf, and that proportion gradually increased to ~70% by 18hpf (Figure 4E), indicating that Wee1 misexpression strongly delays cell cycle progression, blocking cell divisions in a substantial fraction of embryos.

Focusing on ASMFs, we found that the proportion of Ebf+ cells in control embryos progressively increased from ~20% showing "weak" expression at 15.5hpf to ~90%
showing "strong" expression by 18hpf (Figure 5F; see Figure 5D for examples of "weak" and "strong" expression). This semi-quantitative analysis revealed an under-appreciated dynamic at the onset of Ebf expression, which appears to take at least one hour to be "strongly" expressed in >75% of newborn ASMFs (Figure 4F).

To evaluate the impact of Wee1-induced mitosis inhibition on Ebf accumulation, we focused on undivided STVCs at each time point (hence the lower numbers in Figure 4F compare to Figure 4E). By 17hpf, wee1-expressing delayed STVCs showed "strong" Ebf expression in comparably high proportions of embryos. However, these proportions were significantly lower at 16 and 16.5hpf (Chi-square tests, $P=0.002$ and $P=0.0003$, respectively), with ~1.5 and ~1.2 times less "strongly" expressing cells than in the control distributions (hypergeometric tests, $P=0.0005$ and $P=0.0001$, respectively). These semi-quantitative data suggest that the cardiopharyngeal network can eventually unfold and lead to high levels of Ebf expression independently of cell divisions, albeit with a delay revealing that cell divisions probably entrain Ebf upregulation in early ASMFs.

**Transition from a MAPK-dependent to a MAPK-independent and autoregulative mode of Ebf expression in early ASMFs.**

Given the semi-quantitative nature of our analysis, and the relatively subtle effects observed on Ebf dynamics, we sought to further probe the mechanisms that regulate the onset and upregulation of Ebf expression in early ASMFs, and the biological significance for cell-fate specification. Since we observed a progressive accumulation of Ebf mRNAs, and a transition from a MAPK-dependent onset to a MAPK-independent maintenance of Ebf transcription (Figure 3I, J), we reasoned that the window of MAPK-dependence might coincide with the accumulation of Ebf mRNAs between 16 and 17hpf. To test this possibility, we treated embryos with the MEK inhibitor U0126 at successive time points, assayed ongoing transcription using intronic probes and counted the numbers of Ebf
transcribing cells (Figure 6A). This analysis revealed that Ebf transcription gradually lost its sensitivity to MAPK inhibition between 16 and 17hpf, i.e. during the first hour of the ASMF cycle when Ebf mRNAs normally accumulate (as shown in Figure 5E, F).

Because Ebf transcription becomes independent from MAPK by the time Ebf mRNA have accumulated to "high" levels, and because Ebf expression lasts for several days in the progeny of the ASMFs, we reasoned that autoregulation might suffice to maintain high levels of Ebf mRNA past the MAPK-dependent onset. To test this possibility, we misexpressed the Ebf coding sequence using the STVC-specific T12 enhancer as described (Tolkin and Christiaen, 2016). Assaying endogenous Ebf transcription using intronic probes demonstrated that, in addition to its normal expression in the ASMFs, Ebf misexpression caused precocious and ectopic activation of the endogenous locus in the STVCs, and in the MAPK-negative SHPs, respectively (Figure 6C-F). This result suggests that Ebf transcription bypasses both requirements for cell-division coupling and MAPK inputs if high levels of Ebf gene products are present in the cell.

We reasoned that, if high levels of Ebf expression can promote its own transcription independently of MAPK signaling, then Ebf misexpression should be sufficient to rescue a chemical inhibition of MAPK at a critical stage. We tested this possibility by combining Ebf misexpression using the STVC-specific T12 enhancer and U0126 treatments starting at 16hpf, which normally block Ebf expression (Figure 6A, D-F). We observed that transcription of the endogenous Ebf locus became independent of early MAPK activity upon misexpression of an Ebf cDNA, further supporting the notion that high levels of Ebf expression suffice to maintain Ebf transcription independently of MAPK activity.

A potentially important implication of this transient MAPK-dependence of is to render Ebf expression initially reversible. For instance, Ebf occasionally turns on precociously in the STVCs of a small proportion of embryos (Figure S4). Given the
powerful anti-cardiogenic effects of Ebf (Razy-Krajka et al., 2014; Stolfi et al., 2010),
persistent Ebf expression would have dramatic consequences for SHP development
(Wang et al., 2013). However, because MAPK activity is excluded from the SHPs, and the
early phase of Ebf expression depends upon continuous MAPK activity, we surmise that
Ebf cannot be maintained in the SHPs. For instance, when embryos from the same
electroporated batch were fixed at the time of early U0126 treatment (i.e. 15.75 and
16.25hpf) and ~4 hours later, at 20hpf, and assayed for Ebf transcription using intronic
probes, initially wild-type patterns of Ebf transcription could not be maintained (Figure
S5A). This suggests that, although Ebf can be activated precociously in a MAPK-
dependent manner, its expression shuts off in the SHPs upon MAPK inhibition following
STVC division.

We further addressed the interplay between cell division, MAPK signaling and Ebf
expression. We reasoned that, if cell divisions entrain Ebf accumulation and the
transition to a MAPK-independent autoregulative mode, then delaying STVC divisions
should extend the period of MAPK-dependent Ebf transcription. We tested this
possibility by expressing Wee1 under the control of the STVC-specific T12 enhancer, and
treated embryos with U0126 at 17hpf, which inhibited the maintenance of Ebf
transcription in only 15% to 20% of the control embryos (Figures 6A, S5B). The
proportion of embryos showing U0126-sensitive Ebf transcription increased to almost
50% upon T12>Wee1 expression (Figure S5B), which is consistent with our hypothesis
that inhibiting the G2/M transition delayed the accumulation of Ebf gene products thus
postponing the transition from a low level/MAPK-dependent to an high level/MAPK-
independent and self-activating mode of Ebf regulation.

Taken together these data lead us to propose a model for Ebf regulation whereby
Hand-r, Tbx1/10, ongoing MAPK signaling and cell-cycle-regulated transcriptional
input(s) govern the onset and initial accumulation of Ebf gene products during the first
hour of the ASMF cycle, whereas the maintenance of Ebf expression relies primarily on MAPK-independent autoactivation, following initial accumulation (Figure 7).
Discussion

Here, we demonstrated that the progressive restriction of FGF/MAPK signaling follows asymmetric cell divisions of multipotent progenitors and patterns the ascidian cardiopharyngeal mesoderm in space and time. This leads to the localized expression of Hand-\(r\), Tbx1/10 and Ebf in fate-restricted pharyngeal muscle precursors, and their concomitant exclusion for first and second heart precursors. We show that coherent feedforward circuits encode the successive activations of Hand-\(r\), Tbx1/10 and Ebf, whereas cell divisions entrain the progression of this regulatory sequence and thus define the timing of gene expression. Finally, we provide evidence that the initiation of Ebf expression depends on MAPK activity in early ASMF, until Ebf accumulation permits MAPK-independent auto-activation. Given the potent anti-cardiogenic, and pro-pharyngeal muscle effects of Ebf (Razy-Krajka et al., 2014; Stolfi et al., 2010), we surmise that the latter switch corresponds to the transition from a cardiopharyngeal multipotent state to a committed pharyngeal muscle identity.

Spatial patterning by localized maintenance of FGF/MAPK signaling.

Our results demonstrate that MAPK signaling is maintained only in the lateral-most daughter cells following each asymmetric division of multipotent cardiopharyngeal progenitors - the TVCs and STVCs. This asymmetric maintenance is necessary and sufficient for the progressive and localized deployment of the pharyngeal muscle network. Notably, the TVCs themselves are initially induced by similar polarized FGF/MAPK signaling coincidental to asymmetric cell divisions of their mother cells, aka the B8.9 and B8.10 founder cells (Davidson et al., 2006). Detailed analyses have since indicated that asymmetrical maintenance of sustained FGF/MAPK signaling involves intrinsic Cdc42-dependent polarity of the founder cells, which promotes polarized cell-matrix adhesion of the prospective TVC membrane to the ventral epidermis. The latter
differential integrin-mediated adhesion promotes localized MAPK activation, leading to TVC induction (Cooley et al., 2011; Norton et al., 2013). It has been proposed that adhesion- and caveolin-dependent polarized FGFR recycling during mitosis accounts for the localized activation of MAPK in the prospective TVCs (Cota and Davidson, 2015). Whereas similar mechanisms could in principle account for asymmetric maintenance of FGF/MAPK signaling in STVCs and ASMFs, this has not been formally tested and there are notable differences opening the possibility that other mechanisms may be at work: during TVC induction, MAPK signaling is maintained in the smaller daughter cell that contacts the epidermis, whereas in the following divisions, MAPK activity persists in the larger daughter cells and all cells maintain contact with the epidermis (Nicole Kaplan and Lionel Christiaen, data not shown). Moreover, using an FGFR::mKate2 fusion protein similar to that used in previous studies, we could not observe a marked polarized distribution of FGFR molecules to the lateral-most cells (the STVCs and ASMFs; Yelena Bernadskaya and Lionel Christiaen, data not shown). However, the fact that constitutively active forms of M-Ras and Mek1/2 were sufficient to bypass the loss of MAPK activity, and impose pharyngeal muscle specification, indicates that differential FGF/MAPK activity is regulated upstream of M-Ras. Further work is needed to elucidate the cellular and molecular mechanisms governing the spatiotemporal patterns of FGF/MAPK signaling in the cardiopharyngeal mesoderm. In particular, it will be important to disentangle the relative impacts of extrinsic (i.e. tissues, contacts) vs. intrinsic (i.e. asymmetric cell division) effects onto FGF/MAPK signaling and the downstream transcriptional inputs.

**Transcriptional effects of differential FGF/MAPK signaling.**

Because differential FGF/MAPK signaling rapidly impacts cell-specific gene expression, we surmise that transcriptional effectors are dynamically regulated. For
instance, even though we have not formally identified the downstream DNA-binding transcription factor (see discussion below), it would be conceivable that the phosphorylated forms of either transcriptional effector persist through cell division upon maintenance of FGF/MAPK activity. However, we have shown that continuous MAPK activity is needed following each division. Therefore, we must invoke elusive phosphatase activities, such as dual-specificity phosphatases (DUSPs; (Patterson et al., 2009), which would reset transcriptional effectors to a dephosphorylated state, thus rendering steady-state FGF/Ras/MAPK inputs necessary.

Systematic dephosphorylation of FGF/MAPK transcriptional effectors is likely to be particularly important for heart fate specification. For instance, whole genome analyses indicate that heart-specific de novo gene expression requires MAPK inhibition (Wang et al., 2017). Although the molecular mechanisms remain elusive, one simple possibility is that, lest fate-restricted heart precursors inhibit MAPK activity, they will activate Tbx1/10 and Ebf, which will block the cardiac program (Razy-Krajka et al., 2014; Stolfi et al., 2010; Wang et al., 2013). Finally, we previously proposed that repressor inputs from Nk4 are needed in the second heart precursors to avoid ectopic activation of Ebf (Wang et al., 2013). The observation that Nk4 transcripts are detected in all cardiopharyngeal cells opened the question as to how Ebf would escape repression by Nk4 in the ASMFs. Differential MAPK activity offers an intriguing possibility: for instance, Nk4/Nkx2-5-mediated repression in other species involves the co-repressor Groucho/TLE (Choi et al., 1999), which is strongly expressed in the cardiopharyngeal mesoderm (Razy-Krajka et al., 2014); and, in flies, MAPK-mediated phosphorylation of Groucho inhibits its repressor function (Cinnamon et al., 2008; Cinnamon and Paroush, 2008; Hasson et al., 2005). Therefore, it is possible that persistent MAPK signaling dampens Groucho/TLE-mediated repressive inputs on cell-specific regulatory genes like Ebf. Future studies will
determine whether such mechanisms provide bistable switches underlying MAPK-
dependent fate choices in the cardiopharyngeal mesoderm.

Temporal deployment of the pharyngeal muscle network

The localized and successive activation of Tbx1/10 and Ebf in STVCs, and ASMFs, respectively, are important features of the cardiopharyngeal network that permit the emergence of diverse cell fates: first and second heart precursors, and atrial siphon muscle precursors. Experimental misexpression of Ebf throughout the cardiopharyngeal mesoderm suffice to inhibit heart development (Razy-Krajka et al., 2014; Stolfi et al., 2010), illustrating how important it is for Ebf expression to be restricted to the ASMF, once the first and second heart precursors are born and have terminated MAPK activity.

Our analyses indicate that the sequential activations of Hand-r, Tbx1/10 and Ebf is encoded in the feed-forward structure of this sub-circuit, whereas the continuous requirement for MAPK inputs and their progressive exclusion from heart progenitors restrict the competence to activate Tbx1/10 and Ebf to the most lateral cells, after each division. Our model implies that each gene may directly respond to transcriptional inputs from MAPK signaling. We have not formally identified the transcription factors(s) that mediate the transcriptional response to FGF/MAPK signaling. However, multipotent cardiopharyngeal progenitors express Ets1/2 and Elk, two common transcriptional effectors of FGF/MAPK signaling in Ciona (Bertrand et al., 2003; Christiaen et al., 2008; Davidson et al., 2006; Gainous et al., 2015). Moreover, Ets1/2 has been implicated in the initial FGF/MAPK-dependent induction of multipotent TVCs (Christiaen et al., 2008; Davidson et al., 2006), and it expression is also progressively restricted to the lateral-most progenitors following each division (Razy-Krajka et al., 2014). Taken together, Ets1/2 and, to some extend, Elk are intriguing candidate transcriptional effectors of FGF/MAPK signaling in cardiopharyngeal development.
The binding preferences of Ets-family factors have been extensively studied in *Ciona*, and they do not depart markedly from conserved Ets-family binding sites with a GGAW core (Bertrand et al., 2003; Farley et al., 2015; Farley et al., 2016; Gueroult-Bellone et al., 2017; Khoueiry et al., 2010). Putative Ets-family binding sites in the TVC-specific *Hand-r* enhancer are conserved between *Ciona intestinalis* and its sibling species *C. robusta* and *C. savignyi*, and necessary for its activity in reporter assays (Woznica et al., 2012).

Similarly, minimal STVC and ASM enhancers for *Tbx1/10* and *Ebf*, respectively, contain conserved putative Ets-family binding sites, although their function has not been tested (Razy-Krajka et al., 2014; Wang et al., 2013 and data not shown). Taken together, these observations suggest that the proposed feed-forward sub-circuit involves direct transcriptional inputs from FGF/MAPK-regulated Ets-family factors on the cardiopharyngeal enhancers of *Hand-r*, *Tbx1/10* and *Ebf*.

Whereas the regulatory architecture of the MAPK; Hand-r; *Tbx1/10*; *Ebf* sub-circuit explains the sequence of activation events, it is also crucial for its correct deployment, and the generation of diverse cell identities, that genes are not fully activated before successive cell divisions. While divisions are not absolutely required for *Ebf* to eventually turn on, cell cycle progression appears to entrain the deployment of this network, especially for *Tbx1/10* and *Ebf* activation in STVCs and ASMFs, respectively. These observations imply that, while the network can eventually unfold, its intrinsic dynamic is slower than observed. This allows first and second heart precursors to be born prior to the onset of *Tbx1/10* and *Ebf*, respectively. The latter sequence is essentially for the heart progenitors to escape the anti-cardiogenic effects of *Tbx1/10* (Wang et al., 2013), and *Ebf* (Razy-Krajka et al., 2014).

Initial *Ebf* expression in early ASMFs is also labile and MAPK-dependent for approximately one hour. This continued requirement for MAPK inputs ensures that, in
the rare instances when Ebf expression starts in the multipotent STVC progenitors and/or expands to the nascent SHPs, inhibition of MAPK shuts off Ebf expression before it reaches the levels needed for commitment to an ASM fate. Indeed, our results indicate that, once Ebf mRNAs have accumulated to high levels, its expression becomes auto-regulative and MAPK-independent. We surmise that this transition coincides with a fundamental switch from a multipotent cardiopharyngeal state to a committed pharyngeal muscle identity.

From this standpoint, the observed entrainment of Ebf expression by the cell cycle can be seen as acceleration of the transition to commitment following asymmetric division of multipotent progenitors. Although the mechanisms remain elusive, it is likely that this requires the M/G1 transition, as the G1 phase has been shown to be particularly conducive to the expression of fate-specific regulators in mammalian pluripotent stem cells (Dalton, 2015; Pauklin et al., 2016; Pauklin and Vallier, 2013; Soufi and Dalton, 2016).

**Conserved dual effects of FGF/MAPK signaling on heart development in chordates**

Previous studies highlighted how FGF/MAPK signaling is necessary along side Mesp during early cardiac development in *Ciona* (Christiaen et al., 2008; Davidson, 2007; Davidson et al., 2006), and how this early requirement also exists in vertebrates (Abu-Issa et al., 2002; Alsan and Schultheiss, 2002; Barron et al., 2000; Brand, 2003; Reifers et al., 2000; Zaffran and Frasch, 2002). We now know that these early FGF/MAPK inputs induce and maintain multipotent cardiopharyngeal states in *Ciona*, including the *Tbx1/10*+ multipotent progenitors that eventually produce the second heart lineage ((Razy-Krajka et al., 2014; Stolfi et al., 2010; Wang et al., 2013; Wang et al., 2017), and this study). Similarly, in vertebrates, regulatory interplay between Fgf8 and Fgf10
signaling and Tbx1 is required for development of both pharyngeal arch and second heart field derivatives, presumably in part by maintaining an undifferentiated and proliferative state (Abu-Issa et al., 2002; Aggarwal et al., 2006; Brown et al., 2004; Chen et al., 2009; Hu et al., 2004; Ilagan et al., 2006; Kelly and Papaioannou, 2007; Park et al., 2006; Park et al., 2008; Vitelli et al., 2002b; Watanabe et al., 2010; Watanabe et al., 2012). Notably, FGF signaling acts in successive phases, and its inhibition is necessary for final myocardial specification and differentiation (Hutson et al., 2010; Marques et al., 2008; Tirosh-Finkel et al., 2010; van Wijk et al., 2009). Conversely, continued FGF signaling beyond the multipotent mesodermal progenitor stages was shown to promote smooth muscle and epicardial differential in the heart (Hutson et al., 2010; van Wijk et al., 2009), and also myoblast specification and/or skeletal muscle differentiation in the head, with the expression of FGF ligands being maintained in the pharyngeal arches (Bothe et al., 2011; Buckingham and Vincent, 2009; Michailovici et al., 2015; Michailovici et al., 2014; von Scheven et al., 2006). Taken together, these and our data suggest that FGF/MAPK signaling plays evolutionary conserved roles during chordate cardiopharyngeal development, by promoting the specification of successive mesodermal and Tbx1+ multipotent states, and a fate-restricted non-cardiac muscle identity, while MAPK inhibition is required for myocardial specification and differentiation in the first and second heart field, successively.
Material and methods

Animals, electroporations, and chemical treatments

Gravid wild *Ciona intestinalis* type A, now called *Ciona robusta* (Pennati et al., 2015), were obtained M-REP (Carlsbad, CA, USA), and kept under constant light to avoid spawning. Gametes from several animals were collected separately for *in vitro* cross-fertilization followed by dechorionation and electroporation as previously described (Christiaen et al., 2009a, b). Different quantities of plasmids were electroporated depending on the constructs. Typically, 50 µg of DNA was electroporated for NLS::lacZ or plain mCherry driving constructs but only 15 µg for *Mesp-1>H2B::mCherry*. For perturbation constructs, 70 µg were usually electroporated, except for *Mesp>NLS::Cas9::NLS* (30 µg) and pairs of U6>sgRNA plasmids (25 µg each). U0126 (Cell Signaling Technology, Danvers, MA) was used at 5µM in artificial seawater from a stock solution of 20mM in DMSO. Cytochalasin B (Sigma, Saint Louis, MO) was used at ~3 µg/mL from a 10 mg/mL stock solution in DMSO as previously performed (Jeffery et al., 2008). Control embryos were incubated in parallel with corresponding concentrations of DMSO alone.

In situ hybridization

*In situ* hybridizations were carried out essentially as described previously (Christiaen et al., 2009c; Razy-Krajka et al., 2014), using DIG labeled riboprobes, anti-DIG-POD Fab fragments (Roche, Indianapolis, IN), and Tyramide Amplification Signal coupled to Fluorescein (Perkin Elmer, MA). Reporters expressed in the lineage of interest were marked using anti-β-galactosidase monoclonal mouse antibody (1:1000; Promega, Fitchburg, WI) or anti-mCherry rabbit polyclonal antibody (1:500; BioVision 5993-100), respectively targeted with anti-mouse or anti-rabbit secondary antibody coupled with
Alexa 648 (1:500; Invitrogen, Carlsbad, CA). The different probes used in this study were described previously (Razy-Krajka et al., 2014; Stolfi et al., 2010; Wang et al., 2013).

**dpERK/mcherry double fluorescent immunostaining**

Samples were fixed, as for in situ hybridizations, in MEM-PFA with Tween 20 (0.05%) but only for 30 minutes at room temperature, washed three times in PBSt (Tween 20 0.01%) for 10 minutes, gradually dehydrated every 10 minutes in Ethanol/PBS series (33%, 50%, 80%) and Methanol 100%. Samples were then gradually rehydrated every 10 minutes in Methanol/PBSt series, rinsed three times in PBSt, permeabilized with PBS Triton-100 (0.2%) for 30 minutes and incubated for 2 hours at room temperature with anti-dpERK mouse monoclonal antibody (1:200; Sigma, Saint Louis, MO) and anti-mCherry polyclonal antibody from rabbit (1:500; Biovision, Milpitas, CA) in PBS 0.01% Triton-100 (T-Pbs) supplemented with 2% normal goat serum. Samples were then washed three times in T-PBS and incubated in anti-mouse and anti-rabbit antibodies (1:500 each), respectively coupled with Alexa 488 and Alexa 568 (Invitrogen, Carlsbad, CA), overnight at 4°C or for 2 hours at room temperature. Finally, samples were rinsed three times in T-PBS for 15 minutes and mounted in Prolong Gold (Molecular Probes, Eugene, OR).

**Molecular cloning**

Coding sequences for wild-type M-Ras (KH.L172.2), Mek1/2 (KH.L147.22), Cdkn1b.a (Cdkn1b, KH.C14.564), and Cdkn1b.b (Noto16, KH.S643.6) were PCR-amplified from cDNA libraries prepared by reverse transcription of total RNA from mixed developmental stages. Insertion of the products into expressing vectors was performed using regular restriction/ligation or In-fusion (Clontech, Mountain View, CA) procedure. Oligonucleotide directed mutagenesis or two-step overlap PCRs were used to generate
the point mutated forms M-Ras\(^{G22V}\) and Mek\(^{S220E.S216D}\) from the corresponding wild-type sequences. We also used oligonucleotide directed mutagenesis to generate mismatches in the PAM sequences adjacent to the sgRNA targets for Hand-r (153C>T 574C>T for Hand-r\(^{PAMmis}\)) and Tbx1/10 (325G>A and 579G>A for Tbx1/10\(^{PAMmis}\)). Due to the absence of a correct PAM sequence (NGG, (reverse complement CCN)), overexpressed Hand-r\(^{PAMmis}\) and Tbx1/10\(^{PAMmis}\) are resistant to the Cas9 nuclease activity. Primer sequences are listed in Supplementary Table 1.

CRISPR/Cas9-mediated loss of Hand-r function

The pair of single guide RNA (sgRNA) targeting Tbx1/10 (sgTbx1/10) has been validated previously (Tolkin and Christiaen, 2016). Rescue of the Tbx1/10 loss-of-function was achieved by TVC-specific overexpression of Tbx1/10\(^{PAMmis}\) driven by a FoxF enhancer (FoxF-1>Tbx1/10\(^{PAMmis}\)). For Hand-r loss of function, sgRNAs were first designed to avoid genomic off-targets and tested as described (Gandhi et al., 2017). In short, sgRNA expressing cassettes (U6>sgRNA) were assembled by single step overlap PCR. Individual PCR products (~25 µg) were electroporated with EF1a>NLS::Cas9::NLS (30µg), Myod905>Venus (50 µg), driving ubiquitous expression of Cas9 and a widely expressed fluorescent reporter construct, respectively. Efficient electroporation was confirmed by observation of fluorescence before genomic DNA extraction around 16 hpf (18°C) using QIAamp DNA Micro kit (Qiagen, German Town, MD). Mutagenesis efficacy of individual sgRNAs, as a linear function of Cas9-induced indel frequency, was estimated from electrophoregrams following Singer sequencing of the targeted regions amplified from extracted genomic DNA by PCR. Result of the relative quantification of the indel frequency (“corrected peakshift” of 22% and 24%) was considered high enough for both sgRNAs targeting Hand-r, which were finally selected. The corresponding cassettes were cloned into plasmid for repeated electroporations to study the loss of function of Hand-r.
Rescue of Hand-r loss-of-function was achieved by overexpression of Hand-r<sup>PAMmis</sup> driven by a FoxF TVC specific enhancer (FoxF-1>Hand-r<sup>PAMmis</sup>). In order to control the specificity of the CRISPR/Cas9 system, sgRNAs targeting Neurogenin, a gene not expressed in the TVC and their progeny, was electroporated in parallel. Sequences of the DNA targets and oligonucleotides used for the sgRNAs are listed in Supplementary Table 1.

**Observation and imaging**

Samples were usually scored under a DM2500 epifluorescent microscope (Leica Microsystems, Wetzlar, Germany). Imaging was performed using a TCS SP8 X inverted confocal microscope equipped with a white light laser, AOBS and HyD detectors (Leica Microsystems).

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Figure 1. Spatio-temporal restriction of ERK activity reflects FGF requirement for the specification of cardiopharyngeal progenitors. (A) Schematic of Ciona development showing asymmetric cell divisions and resulting cell fates of the cardiopharyngeal mesoderm (CPM). Embryonic and larval stages (St) according to (Hotta et al., 2007) with hours post fertilization (hpf) at 18°C. Anterior tail muscle (ATM, gray), trunk ventral cell (TVC, green), secondary TVC (STVC, green), first heart precursor (FHP, red), second heart precursor (SHP, orange), atrial siphon founder cell (ASMF, blue). Black bars link sister cells. Dashed lines: ventral midline. The first stage presents a quasi-lateral view while the second and third stages present quasi-ventral views. Anterior is to the left. Scale bar, 50 µm. (B) ERK activity visualized by anti-dpERK antibody (green). TVCs and their progeny are marked by mCherry driven by Mesp and revealed by anti-mCherry antibody (red). H2B::mCherry and hCD4::mCherry accumulate in the nuclei and at the cell membrane, respectively. Arrowheads indicate STVCs and ASMFs at 14 and 16 hpf, respectively. Arrows indicate FHPs and open arrowheads mark SHPs. Anterior to the left. Scale bar, 10 µm. See also Figure S1 for broader time series of dpERK immunostaining in the B7.5 lineage. (C, D) TVC-specific overexpression of dnFGFR induces loss of expression of key lateral CPM markers visualized by in situ hybridization. (C) Representative expression patterns of key CPM genes (Hand-related, Tbx1/10, Ebf) in control embryos (control, electroporated with FoxF(TVC):bpFOG-1>Venus) and TVC-specific dnFGFR expression (electroporated with FoxF(TVC):bpFOG-1>dnFGFR::mCherry) individuals. TVCs and progeny are marked with Mesp>NLS::lacZ (red). Loss of expression in half of the TVC progeny, as presented for Ebf, is assumed to be due to left-right mosaicism. Arrowheads mark the ASMFs. Anterior is to the left. Scale bar, 10 µm. (D) Corresponding histograms with the phenotype proportions. For simplicity, loss of gene expression in half or all of the TVCs and their progeny were combined in the same category. "n" corresponds to the number of individual halves documented per condition.
Figure 2. Constitutively active M-Ras and MEK are sufficient to impose a pharyngeal muscle fate in the cardiopharyngeal lineage. (A) Diagram of the FGF/MAPK transduction pathway with constitutive activation by M-Ras\(^{G22V}\) and MEK\(^{S216D,S220E}\) mutants. (B) Expression patterns of markers of the lateral TVC progeny, Htr7 (a, b, c), Tbx1/10 (d, e, f) and Ebf (g, h, i), visualized by in situ hybridization following TVC-specific over-expression of M-Ras\(^{WT}\) (as control), M-Ras\(^{G22V}\) and MEK\(^{S216D,S220E}\). M-Ras\(^{WT}\) overexpression (a, d, g) does not alter the wild-type spatial expression patterns of Htr7, Tbx1/10 and Ebf in lateral TVC derivatives (STVC and ASMF) and excluded from the median heart precursors. TVC-specific over-expression of M-Ras\(^{G22V}\) (b, e, h) or MEK\(^{S216D,S220E}\) (c, f, i) induces ectopic expression of STVC and/or ASMF markers (Htr7, Tbx1/10 and Ebf) in the more median cells, that normally form cardiac precursors. Solid arrowheads indicate STVCs and ASMFs at 15 and 18 hpf, respectively. Arrows indicate FHPs and open arrowheads mark SHPs. At 18 hpf, the FHPs start dividing or have divided into 4 cells. Anterior to the left. Scale bar, 10 µm. (C) Corresponding histograms: Larvae with TVC-specific over-expression of MEK\(^{WT}\) retain the wild-type expression patterns. For simplicity, ectopic expressions in half to all of the cardiac precursors were combined in the same phenotype category. "n" corresponds to the number of embryo halves documented per condition. See also Figure S2.
Figure 3. Temporal requirement for MAPK activity permits the progressive deployment of the cardiopharyngeal regulatory program. (A) Summary of the CPM cell lineage showing the
different U0126 treatments with regard to the timing of the cell divisions. Abbreviations and color codes as in Figure 1. (B, C) Proportions of embryo halves with wild-type or downregulated expression of Hand-r at 12.5 hpf (B) and 15 hpf (C) following 3-hour incubations in U0126 (with DMSO as control treatment). (D) Hand-r expression visualized by in situ hybridization at 15 hpf in control (DMSO treated) and U0126 treated embryos. In control embryos, Hand-r remains expressed in the STVCs and downregulated in the FHPs. In U0126 (12-15 hpf) treated embryos, downregulation of Hand-r expression is observed throughout the TVC progeny (STVCs and FHPs), suggesting inhibition of transcription and inheritance of remnant transcripts following TVC divisions. (E, F) Proportions of embryo halves with wild-type or downregulated expression of Htr7 at 12.5 hpf (E) and 15 hpf (F) following 3-hour incubations in U0126 (with DMSO as control treatment). (G) Proportions of larvae with wild-type expression or downregulated expression of Tbx1/10 at 16 hpf following 4-hour incubation in U0126 (with DMSO as control). (H) Proportions of larvae with wild-type or downregulated expression of Ebf at 18 hpf following a three hour incubation in U0126 (with DMSO as control). (I) Proportions of larvae with wild-type or downregulated transcription of Ebf at 18 hpf following a 3-hour incubation in U0126 (DMSO as vehicle control). (J) Pattern of nascent Ebf transcripts visualized by in situ hybridization with intronic probes (green) at 20 hpf. The nuclear dots reveal the active transcription sites in the four ASMPs per side in larvae, both control/DMSO- and U0126-treated from 17 to 20 hpf. (K) Ebf expression (green) in 18hpf larvae expressing control M-RasWT, constitutively active M-RasG22V or dominant negative dnFGFR under the control of the T12 element, an STVC-specific Tbx1/10 enhancer. Arrows: first heart precursors (FHP); open arrowhead: second heart precursors (SHPs); closed arrowheads: ASM founder cells (ASMFs); dotted line: midline. (L) Proportions of larvae with wild-type or downregulated expression of Ebf at 18 hpf in larvae with Venus (control), M-RasWT, M-RasG22V, or dnFGFR driven by Tbx1/10 cis-regulatory sequence and overexpressed in the STVCs. "n" : number of individual halves documented per condition.
Figure 4. M-Ras/MAPK-driven feed-forward subcircuits control the successive activations of Hand-r, Tbx1/10 and Ebf. (A) Proportions of embryo halves with indicated Tbx1/10 expression patterns following TVC-specific CRISPR/Cas9-mediated mutagenesis of Neurogenin/Neurog as a control (sgCtrl), and Hand-r (sgHand-r). TVC-specific overexpression of a CRISPR/Cas9-resistant form of Hand-r with mutation in the PAM sequence (Hand-r\textsuperscript{PAMmis}) rescued Tbx1/10 expression in the sgHand-r "background". TVC-specific overexpression of a constitutively active M-Ras mutant (M-Ras\textsuperscript{G22}) (control: M-Ras\textsuperscript{WT}) was sufficient to induce ectopic expression of Tbx1/10 in the FHPs in sgCtrl embryos but not in sgHand-r embryos indicating that Hand-r is necessary for M-Ras-dependent activation of Tbx1/10 transcription. (B) Proportions of embryo halves with indicated Tbx1/10 expression patterns following TVC-specific overexpression of Hand-r or a neutral reporter (Venus) and treated from 12 to 15hpf with the MEK inhibitor U0126 (+) or with DMSO (-) as control. Hand-r overexpression is not sufficient to rescue loss of Tbx1/10 expression due to MAPK inhibition indicating that M-Ras/MAPK activity is required in parallel of Hand-r expression to activate Tbx1/10 transcription in the TVC progeny. (C) Tbx1/10 is necessary downstream of M-Ras/MAPK activity to activate Ebf transcription in the TVC progeny. Shown are proportions of Ebf expression phenotypes following TVC-specific CRISPR/Cas9-mediated loss of Tbx1/10 function (sgTbx1/10), with Neurog-targeting sgRNA as control (sgCtrl). Specificity of Tbx1/10 loss of function was validated through rescue of Ebf expression with TVC-specific overexpression of a CRISPR/Cas9 resistant form of Tbx1/10 (Tbx1/10\textsuperscript{PAMmis}). Ectopic Ebf expression in SHPs in Tbx1/10\textsuperscript{PAMmis} larvae is explained by precocious misexpression of Tbx1/10 in the TVC as described in Wang et al, 2013. TVC-specific overexpression of M-Ras\textsuperscript{G22} (M-Ras\textsuperscript{G22}), with wild type M-Ras (M-Ras\textsuperscript{WT}) as control, was sufficient to induce ectopic expression of Ebf in the cardiac precursors in sgCtrl embryos but not in sgTbx1/10 embryos indicating that Tbx1/10 is necessary for M-Ras-dependent activation of Ebf transcription. (D, E) Proportions (D) and examples (E) of 15hpf larvae halves showing induced Ebf expression phenotypes in sgCtrl and sgHand-r CRISPR/Cas9 conditions combined with TVC-specific overexpression of a neutral reporter (Venus), Hand-r\textsuperscript{PAMmis}, or Tbx1/10, and with MEK inhibition by U0126 (+) or not (DMSO control (-)). Arrowhead: STVCs, Arrows: FHPs, dotted line: ventral midline (F) Loss of Hand-r function impaired the ability of Tbx1/10 to induce ectopic Ebf expression. For simplicity, ectopic expressions in half to all of the cardiac precursors were combined in the same phenotype category. "n": number of individual halves documented per condition. (G) Summary model of the temporal deployment of FGF/MAPK-driven feed-forward subcircuits leading to the sequential activations of Tbx1/10 and Ebf in the STVCs and ASMFs, respectively.
**Figure 5. Temporal deployment of the cardiopharyngeal network is partially coupled with cell cycle progression.** (A) Schematic representation of the canonical eukaryotic cell cycle, and actions of the perturbations used in this study. (B) Tbx1/10 and Ebf expression at indicated time points, and following inhibition of cytokinesis by cytochalasin B treatment at indicated time points. Note that 15 to 19hpf treatment is applied AFTER the first division and birth the FHPs, which do not activate Ebf at 19hpf (right panel, arrows). (C) Inhibition of G1/S or G2/M blocks TVC division, and reduces Tbx1/10 expression. Picture shows left-right mosaic embryo, with TVCs that have not divided on the electroporated side (marked by Mesp->H2B::mCherry, red), one cell turned on Tbx1/10, but not the other. Left: the proportions of embryos showing strong Tbx1/10 expression is substantially reduced compared to control embryos (e.g. Figure 1, and (Wang et al., 2013)). (D) Inhibition of G2/M in the STVCs by misexpression of Wee1 using the Tbx1/10 T12 enhancer inhibits STVC division, and has a mild impact on Ebf expression at 18hpf. Open arrows indicate STVCs that have not divided, but express high (middle) or low (right) levels of Ebf. Left: control larva showing high Ebf expression in the ASMF (closed arrowheads), but neither in the SHPs (open arrowheads) nor in the FHPs (Arrows). (E) Proportions of larva halves fixed at successive time points and showing undivided STVCs, or ASMFs with or without ectopic Ebf expression in the SHPs following STVC-specific expression of the G2/M inhibitor Wee1 (+), or a control construct (-). See Figure S4C for an example of ectopic Ebf expression in the SHPs (grey labels). Note the sharp increase in % of larva with ASMF between 15.5 and 16hpf, indicating that mitosis occurs primarily during this time window, but is delayed in a majority of larvae upon Wee1 misexpression. (F) Proportions of larva halves with cells showing indicated Ebf expression. The numbers (n) for cells expressing Wee1 focus on cells that have not divided (% shown in E), to estimate the dynamics of Ebf activation in G2/M-inhibited cells. Control cells consist mostly ASMFs after 15.5hpf as shown in (E). Wee1 and controls distributions differ significantly only at 16 and 16.5hpf (**, p<0.01, Chi² test), suggesting that Wee1 merely delays the accumulation of Ebf transcripts. In all image panels, dotted line : ventral midline.
**Figure 6. Ebf regulation transitions from MAPK-dependent to autoregulative during the early phase of ASMF cycle.** (A) Proportions of 20hpf larva halves showing the indicated number of Ebf-expressing cells following U0126 treatments started at the indicated time points. This indicates that, by 17hpf, Ebf expression, which started at ~16hpf, has become largely insensitive to loss of MAPK activity. (B) Summary lineage diagram and time scale indicating the approximate stages for U0126 and DMSO (control) treatments for the results shown in (C, D). (C) Control (Ctrl) and Ebf-misexpressing embryos fixed at 16hpf, prior to chemical treatments, and stained for nascent transcripts with an intronic Ebf probe. In controls, the ASMFs (solid arrowhead), but neither the SHPs (open arrowheads) nor the FHPs (arrows), actively transcribe Ebf (green nuclear dots). In Larvae misexpressing the Ebf cDNA under the control of the STVC-specific Tbx1/10 enhancer, divisions are delayed and STVCs (solid arrowheads) activated transcription of endogenous Ebf loci (green nuclear dots). (D) After 4 hours, U0126 treated ASMFs no longer transcribe Ebf (top right image, solid arrowheads), whereas control DMSO-treated ASMFs do (top left, green nuclear dots). Upon misexpression of the Ebf cDNA in the STVCs and derivatives, ongoing Ebf transcription is detected at 20hpf in both DMSO and U0126-treated cells, and it persists in both ASMFs (solid arrowheads), and SHPs (open arrowheads). (E, F). Proportions of larva halves showing the indicated Ebf transcription patterns, in indicated experimental conditions, as illustrated in C and D, respectively.
**Figure 7. Summary model.** (A) Schematic representation of cardiopharyngeal lineage cells at successive time points representing the main fate transitions. hpf: hours post-fertilization; TVC: trunk ventral cells; L: Leader T: trailer; migr.: migration; STVC: second trunk ventral cells; FHP: first heart precursors; dotted line: midline; black bars link sister cells; ASMF: atrial siphon muscle founder cells; SHP: second heart precursors; iASMP: inner atrial siphon muscle precursors; oASMP: outer atrial siphon muscle precursor (these cells correspond to stem-cell-like Mrf-; Notch+ precursors and Mrf+; Notch- differentiating myoblasts, respectively; see (Razy-Krajka et al., 2014) for details). (B) Lineage diagram and documented regulatory relationships between indicated genes and pathways, as showing here and in (Razy-Krajka et al., 2014; Wang et al., 2013). In TVCs, primed heart and ASM markers are coexpressed, and maintenance of the STVC and ASM markers requires ongoing FGF/MAPK signaling. Following the first oriented and asymmetric cell division, FGF/MAPK is maintained only in the STVCs, where it permits the continued expression of Hand-r and the activation of Tbx1/10. Cell division, presumably through G1-specific inputs, contributes to Tbx1/10 activation, and Tbx1/10 function antagonizes Gata4/5/6 expression (Wang et al., 2013). In the FHPs, termination of FGF/MAPK signaling inhibits Hand-r expression and prevents Tbx1/10 activation. Following oriented and asymmetric division of the STVCs, FGF/MAPK signaling persists only in the ASMFs, where it permits the transient maintenance of Hand-r and Tbx1/10, both of which act in parallel to FGF/MAPK to activate Ebf expression, together with contributions from presumed G1 inputs. Ebf activities further antagonize the cardiac program (marked by Gata4/5/6, Nk4/Nkx2-5 and Hand expression; (Razy-Krajka et al., 2014; Stolfi et al., 2010; Wang et al., 2013)). Once Ebf expression reaches "high levels", its regulation becomes MAPK-independent and self-activating (this study). It also feeds back negatively on early activators such as Hand-r, and promotes the expression of the muscle determinant Mrf (Razy-Krajka et al., 2014; Tolkin and Christiaen, 2016). We propose that this transition represents commitment to an ASM fate. In the SHPs, termination of FGF/MAPK signaling prevents maintenance of Hand-r and Tbx1/10 expression, which, together with repressive inputs from Nk4/Nkx2-5, inhibits Ebf activation (Wang et al., 2013), and permits heart fate specification (Wang et al., 2017).
Figure S1. Detailed patterns of MAPK activity during early cardiopharyngeal development.

(A) MAPK activation during TVC induction. Close-up views of B7.5 lineage cells marked with Mesp-H2B::mCherry (nuclei) and Mesp-hCD4::mCherry (membranes) and immunostained for dpERK at indicated successive time points between 7 and 10hpf. DpERK staining was not detected in the founder cells at 7hpf, but increased sharply and specifically in the smaller trunk ventral cells (TVCs, open arrows) at 7.5hpf, but not in the larger anterior tail muscles (ATMs). DpERK staining persisted throughout TVC migration (see also B). (B) MAPK activation patterns during cardiopharyngeal fate diversification. DpERK staining was clearly detected in migrating TVCs (open arrows, 11 to 13hpf); in lateral large STVCs (open arrows, 14 to 15hpf), but not in the small median first heart precursors (FHPs, arrows, 14 to 15hpf); in the large lateral atrial siphon muscle founder cells (ASMFs, solid arrowheads, 16 to 17hpf), but neither in the FHPs (arrows), nor in the second heart precursors (SHPs, open arrowheads). (C) Treatment with the MEK inhibitor U0126 between abolished dpERK staining in the lateral STVCs, compared to a control treatment with DMSO. Numbers of embryos showing the presented pattern out of the total numbers of embryos are shown.
Figure S2. Other markers expressed in the TVC need continuous FGF/MAPK inputs for maintenance. All panels show the proportions of 12.5hpf embryos halves showing expression of the indicated genes in late TVCs, following electroporation of either a FoxF(TVC)>Venus control of a FoxF(TVC)>dnFGFR construct that inhibits signaling through FGFR. Wild-type pattern were first reported in (Razy-Krajka et al., 2014).
Figure S3. The constitutively active MEK<sup>S216D,S220E</sup> mutant is sufficient to impose a TVC identity to the whole B7.5 lineage. (A) Control late tailbud embryo showing the left side B7.5 lineage expressing GFP and a MEK<sup>WT</sup> control under the control of the Mesp enhancer. Two TVCs and two ATMs are normally induced, and TVCs migrated into the trunk. (B) Late tailbud embryo showing the left side B7.5 lineage expressing GFP and a MEK<sup>S216D,S220E</sup> mutant under the control of the Mesp enhancer. Four cells are observed as having migrated into the trunk, indicating that they have been induced to acquire a TVC fate and migrate, replicating FGF/MAPK gain-of-function phenotypes as described in (Davidson et al., 2006). (C) Proportions of embryo halves showing the indicated phenotypes. Extra migration is interpreted as ectopic induction of the TVC fate in all B7.5 lineage cells. Scale bar ~ 20µm.
Figure S4. Rare precocious activation of Ebf transcription in STVCs. (A) 15.5hpf Cardiopharyngeal lineage cells expressing Mesp>H2B::mCherry (red) and control Tbx1/10>unc76::GFP construct (not visible). (A.a) Green nuclear dot indicates nascent Ebf transcription in an STVC (open arrow), but not the other, and not in the first heart precursors (FHP; arrow). (A.b) Left pair of nuclei shows an STVC (open arrow) and an FHP (arrow), neither of which express Ebf, whereas the cousin ASMF (solid arrowhead) shows nascent Ebf transcription (green dot). Dotted line: midline. (B) Proportions of STVCs and ASMFs showing indicated Ebf expression patterns. Note that >90% of STVCs do not express Ebf, which turns on almost exclusively in ASMFs. (C) Cardiopharyngeal lineage cells with Ebf expression in the ASMFs (solid arrowheads), and ectopically in the SHP (open arrowheads), but not in the FHPs (arrows), following misexpression of Wee1 using the STVC-specific Tbx1/10 T12 enhancer. Dotted line: midline.
Figure S5. MAPK signaling is necessary for Ebf expression only in early ASMF, and cell cycle inputs shorten the MAPK-dependent period. (A) Proportions of larva halves showing the indicated Ebf transcriptional activity (assayed using intronic probes). Batches of larvae expressing Mesp>H2B::mCherry were split to be fixed for WMFISH or treated with U0126 at 3 successive time points (15.75hpf, 16hpf or 16.25hpf), and the treated larvae were fixed at 20hpf. This data shows that, although all batches expressed Ebf at the beginning of the experiment, only when MEK was inhibited later (16.25hpf) did Ebf transcription persist in 20hpf larvae. (B) Proportions of larva halves showing the indicated numbers of Ebf+ cells at 20hpf, following expression of the G2/M inhibitor Wee1 in the STVCs, under the control of the Tbx1/10(T12)>Venus construct. Negative controls (-) were electroporated with a Tbx1/10(T12)>Venus construct. Larvae were also treated with U0126 (+) or DMSO (as negative control, (-)), starting at 17hpf, which corresponds to the transition from a MAPK-dependent to a MAPK-independent autoregulative mode of Ebf expression (see Figure 6A). Wee1-induced delays in cell cycle progression increased the sensitivity of late Ebf expression to MAPK inhibition, further supporting the notion that cell divisions accelerate the transition from MAPK-dependent to MAPK-independent self-activating regulation of Ebf transcription.
Table S1. oligonucleotides sequences
REFERENCES

Abu-Issa, R., Smyth, G., Smoak, I., Yamamura, K., Meyers, E.N., 2002. Fgf8 is required for pharyngeal arch and cardiovascular development in the mouse. Development 129, 4613-4625.

Aggarwal, V.S., Liao, J., Bondarev, A., Schimmang, T., Lewandoski, M., Locker, J., Shanske, A., Campione, M., Morrow, B.E., 2006. Dissection of Tbx1 and Fgf interactions in mouse models of 22q11DS suggests functional redundancy. Human molecular genetics 15, 3219-3228.

Alsan, B.H., Schultheiss, T.M., 2002. Regulation of avian cardiogenesis by Fgf8 signaling. Development 129, 1935-1943.

Barron, M., Gao, M., Lough, J., 2000. Requirement for BMP and FGF signaling during cardiogenic induction in non-precardiac mesoderm is specific, transient, and cooperative. Developmental dynamics : an official publication of the American Association of Anatomists 218, 383-393.

Beh, J., Shi, W., Levine, M., Davidson, B., Christiaen, L., 2007. FoxF is essential for FGF-induced migration of heart progenitor cells in the ascidian Ciona intestinalis. Development 134, 3297-3305.

Bertrand, V., Hudson, C., Caillol, D., Popovici, C., Lemaire, P., 2003. Neural tissue in ascidian embryos is induced by FGF9/16/20, acting via a combination of maternal GATA and Ets transcription factors. Cell 115, 615-627.

Bondue, A., Lapouge, G., Paulissen, C., Semeraro, C., Iacovino, M., Kyba, M., Blanpain, C., 2008. Mesp1 acts as a master regulator of multipotent cardiovascular progenitor specification. Cell stem cell 3, 69-84.

Bothe, I., Tenin, G., Oseni, A., Dietrich, S., 2011. Dynamic control of head mesoderm patterning. Development 138, 2807-2821.

Brand, T., 2003. Heart development: molecular insights into cardiac specification and early morphogenesis. Developmental biology 258, 1-19.

Brown, C.B., Wenning, J.M., Lu, M.M., Epstein, D.J., Meyers, E.N., Epstein, J.A., 2004. Cre-mediated excision of Fgf8 in the Tbx1 expression domain reveals a critical role for Fgf8 in cardiovascular development in the mouse. Developmental biology 267, 190-202.

Buckingham, M., Vincent, S.D., 2009. Distinct and dynamic myogenic populations in the vertebrate embryo. Current opinion in genetics & development 19, 444-453.

Cai, C.L., Liang, X., Shi, Y., Chu, P.H., Pfaff, S.L., Chen, J., Evans, S., 2003. Isl1 identifies a cardiac progenitor population that proliferates prior to differentiation and contributes a majority of cells to the heart. Developmental cell 5, 877-889.

Chan, S.S., Hagen, H.R., Swanson, S.A., Stewart, R., Boll, K.A., Aho, J., Thomson, J.A., Kyba, M., 2016. Development of Bipotent Cardiac/Skeletal Myogenic Progenitors from MESP1+ Mesoderm. Stem cell reports 6, 26-34.
Chan, S.S., Shi, X., Toyama, A., Arpke, R.W., Dandapat, A., Iacovino, M., Kang, J., Le, G., Hagen, H.R., Garry, D.J., Kyba, M., 2013. Mesp1 patterns mesoderm into cardiac, hematopoietic, or skeletal myogenic progenitors in a context-dependent manner. Cell stem cell 12, 587-601.

Chen, L., Fulcoli, F.G., Tang, S., Baldini, A., 2009. Tbx1 regulates proliferation and differentiation of multipotent heart progenitors. Circulation research 105, 842-851.

Choi, C.Y., Lee, Y.M., Kim, Y.H., Park, T., Jeon, B.H., Schulz, R.A., Kim, Y., 1999. The homeodomain transcription factor NK-4 acts as either a transcriptional activator or repressor and interacts with the p300 coactivator and the Groucho corepressor. The Journal of biological chemistry 274, 31543-31552.

Christiaen, L., Davidson, B., Kawashima, T., Powell, W., Nolla, H., Vranizan, K., Levine, M., 2008. The transcription/migration interface in heart precursors of Ciona intestinalis. Science 320, 1349-1352.

Christiaen, L., Wagner, E., Shi, W., Levine, M., 2009a. Electroporation of transgenic DNAs in the sea squirt Ciona. Cold Spring Harbor protocols 2009, pdb prot5345.

Christiaen, L., Wagner, E., Shi, W., Levine, M., 2009b. Isolation of individual cells and tissues from electroporated sea squirt (Ciona) embryos by fluorescence-activated cell sorting (FACS). Cold Spring Harbor protocols 2009, pdb prot5349.

Christiaen, L., Wagner, E., Shi, W., Levine, M., 2009c. Whole-mount in situ hybridization on sea squirt (Ciona intestinalis) embryos. Cold Spring Harbor protocols 2009, pdb prot5348.

Cinnamon, E., Helman, A., Ben-Haroush Schyr, R., Orian, A., Jimenez, G., Paroush, Z., 2008. Multiple RTK pathways downregulate Groucho-mediated repression in Drosophila embryogenesis. Development 135, 829-837.

Cinnamon, E., Paroush, Z., 2008. Context-dependent regulation of Groucho/TLE-mediated repression. Current opinion in genetics & development 18, 435-440.

Cooley, J., Whitaker, S., Sweeny, S., Fraser, S., Davidson, B., 2011. Cytoskeletal polarity mediates localized induction of the heart progenitor lineage. Nature cell biology 13, 952-957.

Cota, C.D., Davidson, B., 2015. Mitotic Membrane Turnover Coordinates Differential Induction of the Heart Progenitor Lineage. Developmental cell 34, 505-519.

Cowley, S., Paterson, H., Kemp, P., Marshall, C.J., 1994. Activation of MAP kinase kinase is necessary and sufficient for PC12 differentiation and for transformation of NIH 3T3 cells. Cell 77, 841-852.

Dalton, S., 2015. Linking the Cell Cycle to Cell Fate Decisions. Trends in cell biology 25, 592-600.

Davidson, B., 2007. Ciona intestinalis as a model for cardiac development. Seminars in cell & developmental biology 18, 16-26.
Davidson, B., Levine, M., 2003. Evolutionary origins of the vertebrate heart: Specification of the cardiac lineage in Ciona intestinalis. Proceedings of the National Academy of Sciences of the United States of America 100, 11469-11473.

Davidson, B., Shi, W., Beh, J., Christiaen, L., Levine, M., 2006. FGF signaling delineates the cardiac progenitor field in the simple chordate, Ciona intestinalis. Genes & development 20, 2728-2738.

Davidson, B., Shi, W., Levine, M., 2005. Uncoupling heart cell specification and migration in the simple chordate Ciona intestinalis. Development 132, 4811-4818.

Delsuc, F., Brinkmann, H., Chourrout, D., Philippe, H., 2006. Tunicates and not cephalochordates are the closest living relatives of vertebrates. Nature 439, 965-968.

Diogo, R., Kelly, R.G., Christiaen, L., Levine, M., Ziermann, J.M., Molnar, J.L., Noden, D.M., Tzahor, E., 2015. A new heart for a new head in vertebrate cardiopharyngeal evolution. Nature 520, 466-473.

Dumollard, R., Minc, N., Salez, G., Aicha, S.B., Bekkouche, F., Hebras, C., Besnardeau, L., McDougall, A., 2017. The invariant cleavage pattern displayed by ascidian embryos depends on spindle positioning along the cell's longest axis in the apical plane and relies on asynchronous cell divisions. eLife 6.

Farley, E.K., Olson, K.M., Zhang, W., Brandt, A.J., Rokhsar, D.S., Levine, M.S., 2015. Suboptimization of developmental enhancers. Science 350, 325-328.

Farley, E.K., Olson, K.M., Zhang, W., Rokhsar, D.S., Levine, M.S., 2016. Syntax compensates for poor binding sites to encode tissue specificity of developmental enhancers. Proceedings of the National Academy of Sciences of the United States of America 113, 6508-6513.

Gainous, T.B., Wagner, E., Levine, M., 2015. Diverse ETS transcription factors mediate FGF signaling in the Ciona anterior neural plate. Developmental biology 399, 218-225.

Gandhi, S., Haeussler, M., Razy-Krajka, F., Christiaen, L., Stolfi, A., 2017. Evaluation and rational design of guide RNAs for efficient CRISPR/Cas9-mediated mutagenesis in Ciona. Developmental biology 425, 8-20.

George, V., Colombo, S., Targoff, K.L., 2015. An early requirement for nrx2.5 ensures the first and second heart field ventricular identity and cardiac function into adulthood. Developmental biology 400, 10-22.

Gopalakrishnan, S., Comai, G., Sambasivan, R., Francou, A., Kelly, R.G., Tajbakhsh, S., 2015. A Cranial Mesoderm Origin for Esophagus Striated Muscles. Developmental cell 34, 694-704.

Gotoh, N., Laks, S., Nakashima, M., Lax, I., Schlessinger, J., 2004. FRS2 family docking proteins with overlapping roles in activation of MAP kinase have distinct spatial-temporal patterns of expression of their transcripts. FEBS letters 564, 14-18.
Gueroult-Bellone, M., Nitta, K.R., Kari, W., Jacox, E., Beule Dauzat, R., Vincentelli, R., Diarra, C., Rothbacher, U., Dantec, C., Cambillau, C., Piette, J., Lemaire, P., 2017. Spacer sequences separating transcription factor binding motifs set enhancer quality and strength. bioRxiv.

Hasson, P., Egoz, N., Winkler, C., Volohonsky, G., Jia, S., Dinur, T., Volk, T., Courey, A.J., Paroush, Z., 2005. EGFR signaling attenuates Groucho-dependent repression to antagonize Notch transcriptional output. Nature genetics 37, 101-105.

Haupaix, N., Abitura, P.B., Sirour, C., Yasuo, H., Levine, M., Hudson, C., 2014. Ephrin-mediated restriction of ERK1/2 activity delimits the number of pigment cells in the Ciona CNS. Developmental biology 394, 170-180.

Hotta, K., Mitsuhara, K., Takahashi, H., Inaba, K., Oka, K., Gojobori, T., Ikeo, K., 2007. A web-based interactive developmental table for the ascidian Ciona intestinalis, including 3D real-image embryo reconstructions: I. From fertilized egg to hatching larva. Developmental dynamics : an official publication of the American Association of Anatomists 236, 1790-1805.

Hu, T., Yamagishi, H., Maeda, J., McAnally, J., Yamagishi, C., Srivastava, D., 2004. Tbx1 regulates fibroblast growth factors in the anterior heart field through a reinforcing autoregulatory loop involving forkhead transcription factors. Development 131, 5491-5502.

Hudson, C., Darras, S., Caillol, D., Yasuo, H., Lemaire, P., 2003. A conserved role for the MEK signalling pathway in neural tissue specification and posteriorisation in the invertebrate chordate, the ascidian Ciona intestinalis. Development 130, 147-159.

Hutson, M.R., Zeng, X.L., Kim, A.J., Antoon, E., Harward, S., Kirby, M.L., 2010. Arterial pole progenitors interpret opposing FGF/BMP signals to proliferate or differentiate. Development 137, 3001-3011.

Ilagan, R., Abu-Issa, R., Brown, D., Yang, Y.P., Jiao, K., Schwartz, R.J., Klingensmith, J., Meyers, E.N., 2006. Fgf8 is required for anterior heart field development. Development 133, 2435-2445.

Imai, K.S., Levine, M., Satoh, N., Satou, Y., 2006. Regulatory blueprint for a chordate embryo. Science 312, 1183-1187.

Imai, K.S., Satoh, N., Satou, Y., 2002. Early embryonic expression of FGF4/6/9 gene and its role in the induction of mesenchyme and notochord in Ciona savignyi embryos. Development 129, 1729-1738.

Jeffery, W.R., Chiba, T., Krajka, F.R., Deyts, C., Satoh, N., Joly, J.S., 2008. Trunk lateral cells are neural crest-like cells in the ascidian Ciona intestinalis: insights into the ancestry and evolution of the neural crest. Developmental biology 324, 152-160.

Jerome, L.A., Papaioannou, V.E., 2001. DiGeorge syndrome phenotype in mice mutant for the T-box gene, Tbx1. Nature genetics 27, 286-291.
Kaplan, N., Razy-Krajka, F., Christiaen, L., 2015. Regulation and evolution of cardiopharyngeal cell identity and behavior: insights from simple chordates. Current opinion in genetics & development 32, 119-128.

Kattman, S.J., Witty, A.D., Gagliardi, M., Dubois, N.C., Niapour, M., Hotta, A., Ellis, J., Keller, G., 2011. Stage-specific optimization of activin/nodal and BMP signaling promotes cardiac differentiation of mouse and human pluripotent stem cell lines. Cell stem cell 8, 228-240.

Keduka, E., Kaiho, A., Hamada, M., Watanabe-Takano, H., Takano, K., Ogasawara, M., Satou, Y., Satoh, N., Endo, T., 2009. M-Ras evolved independently of R-Ras and its neural function is conserved between mammalian and ascidian, which lacks classical Ras. Gene 429, 49-58.

Kelly, R.G., Jerome-Majewska, L.A., Papaioannou, V.E., 2004. The del22q11.2 candidate gene Tbx1 regulates branchiomeric myogenesis. Human molecular genetics 13, 2829-2840.

Kelly, R.G., Papaioannou, V.E., 2007. Visualization of outflow tract development in the absence of Tbx1 using an FgfF10 enhancer trap transgene. Developmental dynamics: an official publication of the American Association of Anatomists 236, 821-828.

Khoeiry, P., Rothbacher, U., Ohtsuka, Y., Daian, F., Frangulian, E., Roure, A., Dubchak, I., Lemaire, P., 2010. A cis-regulatory signature in ascidians and flies, independent of transcription factor binding sites. Current biology : CB 20, 792-802.

Kuwajima, M., Kumano, G., Nishida, H., 2014. Regulation of the number of cell division rounds by tissue-specific transcription factors and Cdk inhibitor during ascidian embryogenesis. PloS one 9, e90188.

Lemmon, M.A., Schlessinger, J., 2010. Cell signaling by receptor tyrosine kinases. Cell 141, 1117-1134.

Lescroart, F., Chabab, S., Lin, X., Rulands, S., Paulissen, C., Rodolosse, A., Auer, H., Achouri, Y., Dubois, C., Bondue, A., Simons, B.D., Blanpain, C., 2014. Early lineage restriction in temporally distinct populations of Mesp1 progenitors during mammalian heart development. Nature cell biology 16, 829-840.

Lescroart, F., Hamou, W., Francou, A., Theveniau-Ruissy, M., Kelly, R.G., Buckingham, M., 2015. Clonal analysis reveals a common origin between nonsomite-derived neck muscles and heart myocardium. Proceedings of the National Academy of Sciences of the United States of America 112, 1446-1451.

Lescroart, F., Kelly, R.G., Le Garrec, J.F., Nicolas, J.F., Meilhac, S.M., Buckingham, M., 2010. Clonal analysis reveals common lineage relationships between head muscles and second heart field derivatives in the mouse embryo. Development 137, 3269-3279.

Lescroart, F., Mohun, T., Meilhac, S.M., Bennett, M., Buckingham, M., 2012. Lineage tree for the venous pole of the heart: clonal analysis clarifies controversial genealogy based on genetic tracing. Circulation research 111, 1313-1322.
Mandal, A., Holowiecki, A., Song, Y.C., Waxman, J.S., 2017. Wnt signaling balances specification of the cardiac and pharyngeal muscle fields. Mechanisms of Development 143, 32-41.

Mansour, S.J., Resing, K.A., Candi, J.M., Hermann, A.S., Gloor, J.W., Herskind, K.R., Wartmann, M., Davis, R.J., Ahn, N.G., 1994. Mitogen-activated protein (MAP) kinase phosphorylation of MAP kinase kinase: determination of phosphorylation sites by mass spectrometry and site-directed mutagenesis. Journal of biochemistry 116, 304-314.

Mansour, S.J., Lee, Y., Poss, K.D., Yelon, D., 2008. Reiterative roles for FGF signaling in the establishment of size and proportion of the zebrafish heart. Developmental biology 321, 397-406.

Mazzoni, E.O., Mahony, S., Iacovino, M., Morrison, C.A., Mountoufaris, G., Closser, M., Whyte, W.A., Young, R.A., Kyba, M., Gifford, D.K., Wichterle, H., 2011. Embryonic stem cell-based mapping of developmental transcriptional programs. Nature methods 8, 1056-1058.

Merscher, S., Funke, B., Epstein, J.A., Heyer, J., Puech, A., Lu, M.M., Xavier, R.J., Demay, M.B., Russell, R.G., Factor, S., Tokooya, K., Jore, B.S., Lopez, M., Pandita, R.K., Lia, M., Carrion, D., Xu, H., Schorle, H., Kobler, J.B., Scambler, P., Wynshaw-Boris, A., Skoulitchi, A.I., Morrow, B.E., Kucherlapati, R., 2001. TBX1 is responsible for cardiovascular defects in velo-cardio-facial/DiGeorge syndrome. Cell 104, 619-629.

Michailovici, I., Eigler, T., Tzahor, E., 2015. Craniofacial Muscle Development. Current topics in developmental biology 115, 3-30.

Michailovici, I., Harrington, H.A., Azogui, H.H., Yahalom-Ronen, Y., Plotnikov, A., Ching, S., Stumpf, M.P., Klein, O.D., Seger, R., Tzahor, E., 2014. Nuclear to cytoplasmic shuttling of ERK promotes differentiation of muscle stem/progenitor cells. Development 141, 2611-2620.

Mosimann, C., Panakova, D., Werdich, A.A., Musso, G., Burger, A., Lawson, K.L., Carr, L.A., Nevis, K.R., Sabeh, M.K., Zhou, Y., Davidson, A.J., DiBiase, A., Burns, C.E., Burns, C.G., MacRae, C.A., Zon, L.I., 2015. Chamber identity programs drive early functional partitioning of the heart. Nature communications 6, 8146.

Nathan, E., Monovich, A., Tirosh-Finkel, L., Harrelson, Z., Rousso, T., Rinon, A., Harel, I., Evans, S.M., Tzahor, E., 2008. The contribution of Islet1-expressing splanchnic mesoderm cells to distinct branchiomeric muscles reveals significant heterogeneity in head muscle development. Development 135, 647-657.

Nevis, K., Obregon, P., Walsh, C., Guner-Ataman, B., Burns, C.G., Burns, C.E., 2013. Tbx1 is required for second heart field proliferation in zebrafish. Developmental dynamics: an official publication of the American Association of Anatomists 242, 550-559.

Norton, J., Cooley, J., Islam, A.F., Cota, C.D., Davidson, B., 2013. Matrix adhesion polarizes heart progenitor induction in the invertebrate chordate Ciona intestinalis.

Development 140, 1301-1311.
Park, E.J., Ogden, L.A., Talbot, A., Evans, S., Cai, C.L., Black, B.L., Frank, D.U., Moon, A.M., 2006. Required, tissue-specific roles for Fgf8 in outflow tract formation and remodeling. Development 133, 2419-2433.

Park, E.J., Watanabe, Y., Smyth, G., Miyagawa-Tomita, S., Meyers, E., Klingensmith, J., Camenisch, T., Buckingham, M., Moon, A.M., 2008. An FGF autocrine loop initiated in second heart field mesoderm regulates morphogenesis at the arterial pole of the heart. Development 135, 3599-3610.

Patterson, K.I., Brummer, T., O’Brien, P.M., Daly, R.J., 2009. Dual-specificity phosphatases: critical regulators with diverse cellular targets. The Biochemical journal 418, 475-489.

Pauklin, S., Madrigal, P., Bertero, A., Vallier, L., 2016. Initiation of stem cell differentiation involves cell cycle-dependent regulation of developmental genes by Cyclin D. Genes & development 30, 421-433.

Pauklin, S., Vallier, L., 2013. The cell-cycle state of stem cells determines cell fate propensity. Cell 155, 135-147.

Peljto, M., Wichterle, H., 2011. Programming embryonic stem cells to neuronal subtypes. Current opinion in neurobiology 21, 43-51.

Pennati, R., Ficetola, G.F., Brunetti, R., Caiicci, F., Gasparini, F., Grigio, F., Sato, A., Stach, T., Kaul-Strehlow, S., Gissi, C., Manni, L., 2015. Morphological Differences between Larvae of the Ciona intestinalis Species Complex: Hints for a Valid Taxonomic Definition of Distinct Species. PloS one 10, e0122879.

Picco, V., Hudson, C., Yasuo, H., 2007. Ephrin-Eph signalling drives the asymmetric division of notochord/neural precursors in Ciona embryos. Development 134, 1491-1497.

Putnam, N.H., Butts, T., Ferrier, D.E., Furlong, R.F., Hellsten, U., Kawashima, T., Robinson-Rechavi, M., Shoguchi, E., Terry, A., Yu, J.K., Benito-Gutierrez, E.L., Dubchak, I., Garcia-Fernandez, J., Gibson-Brown, J.J., Grigoriev, I.V., Horton, A.C., de Jong, P.J., Jurka, J., Kapitonov, V.V., Kohara, Y., Kuroki, Y., Lindquist, E., Lucas, S., Osoegawa, K., Pennacchio, L.A., Salamov, A.A., Satou, Y., Sauka-Spengler, T., Schmutz, J., Shin, I.T., Toyoda, A., Bronner-Fraser, M., Fujiyama, A., Holland, L.Z., Holland, P.W., Satoh, N., Rokhsar, D.S., 2008. The amphioxus genome and the evolution of the chordate karyotype. Nature 453, 1064-1071.

Racioppi, C., Kamal, A.K., Razy-Krajka, F., Gambardella, G., Zanetti, L., di Bernardo, D., Sanges, R., Christiaen, L.A., Ristoratore, F., 2014. Fibroblast growth factor signalling controls nervous system patterning and pigment cell formation in Ciona intestinalis. Nature communications 5, 4830.
Razy-Krajka, F., Lam, K., Wang, W., Stolfi, A., Joly, M., Bonneau, R., Christiaen, L., 2014. Collier/OLF/EBF-Dependent Transcriptional Dynamics Control Pharyngeal Muscle Specification from Primed Cardiopharyngeal Progenitors. Developmental cell 29, 263-276.

Reifers, F., Walsh, E.C., Leger, S., Stainier, D.Y., Brand, M., 2000. Induction and differentiation of the zebrafish heart requires fibroblast growth factor 8 (fgf8/acerebellar). Development 127, 225-235.

Satou, Y., Imai, K.S., Satoh, N., 2004. The ascidian Mesp gene specifies heart precursor cells. Development 131, 2533-2541.

Shi, W., Levine, M., 2008. Ephrin signaling establishes asymmetric cell fates in an endomesoderm lineage of the Ciona embryo. Development 135, 931-940.

Shi, W., Peyrot, S.M., Munro, E., Levine, M., 2009. FGF3 in the floor plate directs notochord convergent extension in the Ciona tadpole. Development 136, 23-28.

Soufi, A., Dalton, S., 2016. Cycling through developmental decisions: how cell cycle dynamics control pluripotency, differentiation and reprogramming. Development 143, 4301-4311.

Stolfi, A., Gainous, T.B., Young, J.J., Mori, A., Levine, M., Christiaen, L., 2010. Early chordate origins of the vertebrate second heart field. Science 329, 565-568.

Stolfi, A., Gandhi, S., Salek, F., Christiaen, L., 2014a. Tissue-specific genome editing in Ciona embryos by CRISPR/Cas9. Development 141, 4115-4120.

Stolfi, A., Lowe, E.K., Racioppi, C., Ristoratore, F., Brown, C.T., Swalla, B.J., Christiaen, L., 2014b. Divergent mechanisms regulate conserved cardiopharyngeal development and gene expression in distantly related ascidians. eLife 3, e03728.

Stolfi, A., Sasakura, Y., Satou, Y., Christiaen, L., Dantec, C., Endo, T., Naville, M., Nishida, H., Swalla, B., Volff, J.-N., Voskoboynik, A., Dauga, D., Lemaire, P., 2014c. Guidelines for the Nomenclature of Genetic Elements in Tunicate Genomes. Genesis under review.

Stolfi, A., Wagner, E., Taliaferro, J.M., Chou, S., Levine, M., 2011. Neural tube patterning by Ephrin, FGF and Notch signaling relays. Development 138, 5429-5439.

Tirosh-Finkel, L., Elhanany, H., Rinon, A., Tzahor, E., 2006. Mesoderm progenitor cells of common origin contribute to the head musculature and the cardiac outflow tract. Development 133, 1943-1953.

Tirosh-Finkel, L., Zeisel, A., Brodt-Ivenshitz, M., Shamai, A., Yao, Z., Seger, R., Domany, E., Tzahor, E., 2010. BMP-mediated inhibition of FGF signaling promotes cardiomyocyte differentiation of anterior heart field progenitors. Development 137, 2989-3000.

Tolkin, T., Christiaen, L., 2016. Rewiring of an ancestral Tbx1/10-Ebf-Mrf network for pharyngeal muscle specification in distinct embryonic lineages. Development in press.
Tzahor, E., 2009. Heart and craniofacial muscle development: a new developmental theme of distinct myogenic fields. Developmental biology 327, 273-279.

Tzahor, E., Evans, S.M., 2011. Pharyngeal mesoderm development during embryogenesis: implications for both heart and head myogenesis. Cardiovascular research 91, 196-202.

Tzahor, E., Kempf, H., Mootoosamy, R.C., Poon, A.C., Abzhanov, A., Tabin, C.J., Dietrich, S., Lassar, A.B., 2003. Antagonists of Wnt and BMP signaling promote the formation of vertebrate head muscle. Genes & development 17, 3087-3099.

Tzahor, E., Lassar, A.B., 2001. Wnt signals from the neural tube block ectopic cardiogenesis. Genes & development 15, 255-260.

van Wijk, B., van den Berg, G., Abu-Issa, R., Barnett, P., van der Velden, S., Schmidt, M., Ruijter, J.M., Kirby, M.L., Moorman, A.F., van den Hoff, M.J., 2009. Epicardium and myocardium separate from a common precursor pool by crosstalk between bone morphogenetic protein- and fibroblast growth factor-signaling pathways. Circulation research 105, 431-441.

Vitelli, F., Morishima, M., Taddei, I., Lindsay, E.A., Baldini, A., 2002a. Tbx1 mutation causes multiple cardiovascular defects and disrupts neural crest and cranial nerve migratory pathways. Human molecular genetics 11, 915-922.

Vitelli, F., Taddei, I., Morishima, M., Meyers, E.N., Lindsay, E.A., Baldini, A., 2002b. A genetic link between Tbx1 and fibroblast growth factor signaling. Development 129, 4605-4611.

von Scheven, G., Alvares, L.E., Mootoosamy, R.C., Dietrich, S., 2006. Neural tube derived signals and Fgf8 act antagonistically to specify eye versus mandibular arch muscles. Development 133, 2731-2745.

Wagner, E., Stolfi, A., Gi Choi, Y., Levine, M., 2014. Islet is a key determinant of ascidian palp morphogenesis. Development 141, 3084-3092.

Wang, W., Razy-Krajka, F., Siu, E., Ketcham, A., Christiaen, L., 2013. NK4 antagonizes Tbx1/10 to promote cardiac versus pharyngeal muscle fate in the ascidian second heart field. PLoS biology 11, e1001725.

Wang, W., Xiang, S., Jullian, E., Kelly, R.G., Satija, R., Christiaen, L., 2017. A single cell transcriptional roadmap for cardiopharyngeal fate diversification. BioRxiv.

Watanabe, Y., Miyagawa-Tomita, S., Vincent, S.D., Kelly, R.G., Moon, A.M., Buckingham, M.E., 2010. Role of mesodermal FGF8 and FGF10 overlaps in the development of the arterial pole of the heart and pharyngeal arch arteries. Circulation research 106, 495-503.

Watanabe, Y., Zaffran, S., Kuroiwa, A., Higuchi, H., Ogura, T., Harvey, R.P., Kelly, R.G., Buckingham, M., 2012. Fibroblast growth factor 10 gene regulation in the second heart field by Tbx1, Nkx2-5, and Islet1 reveals a genetic switch for down-regulation in the myocardium. Proceedings of the National Academy of Sciences of the United States of America 109, 18273-18280.
Whittaker, J.R., 1973. Segregation during ascidian embryogenesis of egg cytoplasmic information for tissue-specific enzyme development. Proceedings of the National Academy of Sciences of the United States of America 70, 2096-2100.

Witzel, H.R., Cheedipudi, S., Gao, R., Stainier, D.Y., Dobreva, G.D., 2017. Isl2b regulates anterior second heart field development in zebrafish. Scientific reports 7, 41043.

Woznica, A., Haeussler, M., Starobinska, E., Jemmett, J., Li, Y., Mount, D., Davidson, B., 2012. Initial deployment of the cardiogenic gene regulatory network in the basal chordate, Ciona intestinalis. Developmental biology 368, 127-139.

Yagi, H., Furutani, Y., Hamada, H., Sasaki, T., Asakawa, S., Minoshima, S., Ichida, F., Joo, K., Kimura, M., Imamura, S., Kamatani, N., Momma, K., Takao, A., Nakazawa, M., Shimizu, N., Matsuoka, R., 2003. Role of TBX1 in human del22q11.2 syndrome. Lancet 362, 1366-1373.

Yasuo, H., Hudson, C., 2007. FGF8/17/18 functions together with FGF9/16/20 during formation of the notochord in Ciona embryos. Developmental biology 302, 92-103.

Zaffran, S., Frasch, M., 2002. Early signals in cardiac development. Circulation research 91, 457-469.

Zhang, Z., Huynh, T., Baldini, A., 2006. Mesodermal expression of Tbx1 is necessary and sufficient for pharyngeal arch and cardiac outflow tract development. Development 133, 3587-3595.