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## Single nucleotide variants within heart enhancers increase binding affinity and disrupt heart development

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### Summary:

Transcriptional enhancers direct precise gene expression patterns during development and harbor the majority of variants associated with phenotypic diversity, evolutionary adaptations, and disease. Pinpointing which enhancer variants contribute to changes in gene expression and phenotypes is a major challenge. Here we find that suboptimal or low-affinity binding sites are necessary for precise gene expression during heart development. Single nucleotide variants can optimize the affinity of ETS binding sites, causing gain-of-function gene expression, cell migration defects, and phenotypes as severe as extra beating hearts in the marine chordate *Ciona robusta*. In human iPSC-derived cardiomyocytes, a single nucleotide variant within a human

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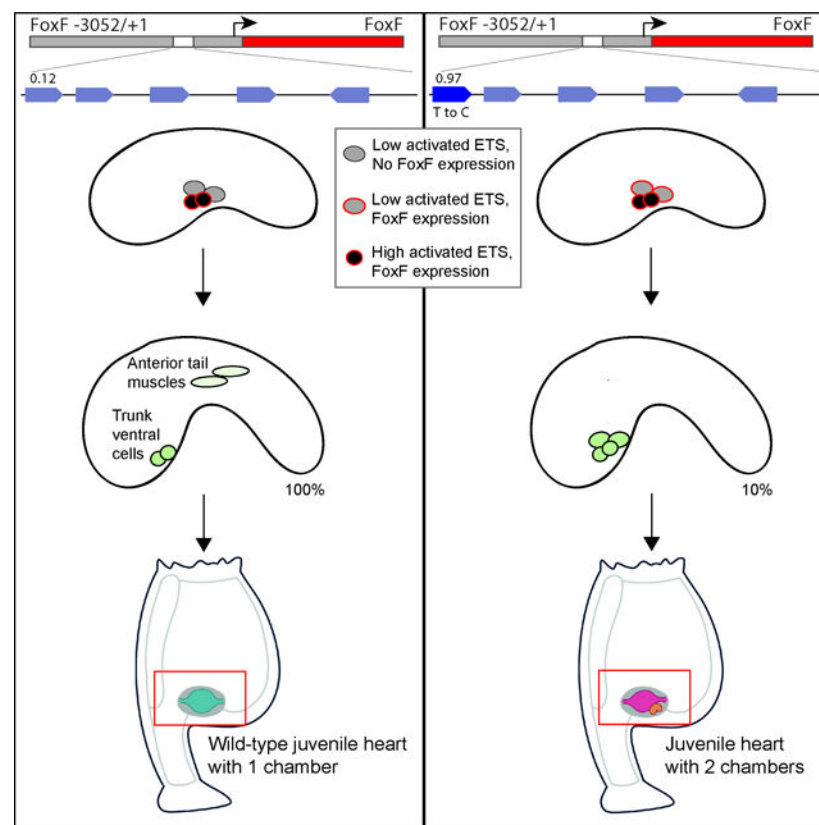
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*GATA4* enhancer increases ETS binding affinity and causes gain-of-function enhancer activity. The prevalence of suboptimal-affinity sites within enhancers creates a vulnerability whereby affinity-optimizing SNVs can lead to gain-of-function gene expression, changes in cellular identity, and organismal-level phenotypes that could contribute to the evolution of novel traits or diseases.

## In Brief:

Jindal et al. show that low-affinity ETS sites are critical for tissue-specific enhancer activity within the developing heart, and that single nucleotide variants can increase binding affinity, causing gain-of-function enhancer activity disrupting heart development. These findings have implications for understanding how enhancer variants alter phenotypes in health, evolution, and disease.

## Graphical Abstract



## Introduction

Enhancers are genomic elements that act as switches to control the location, level, and timing of gene expression to ensure the successful development and integrity of an organism<sup>1,2</sup>. Sequence changes within enhancers can lead to changes in gene expression, phenotypic diversity, novel traits, and disease<sup>3-5</sup>. Indeed, most disease-associated variants within the genome lie within enhancers<sup>6-8</sup>. Yet, pinpointing causal base-pair changes within enhancers is challenging as they are typically associated with many inert variants due to

linkage disequilibrium furthermore it is hard to functionally validate variants at scale in the relevant biological context<sup>9</sup>. Our inability to predict causal enhancer variants is stalling efforts to harness the full potential of genomic data to understand the genetic basis of health and disease. Here we use a mechanistic understanding of the regulatory principles governing enhancers to pinpoint causal enhancer variants that alter gene expression and heart development.

FGF (Fibroblast Growth Factor) signaling plays a critical and conserved role in heart development across bilaterians<sup>10,11</sup>. FGF signaling mediates changes in gene expression through binding of activated ETS transcription factors (TFs) to enhancers<sup>12</sup>. Loss and gain of FGF signaling pathway components cause heart defects in organisms as diverse as *Drosophila* and human<sup>13–15</sup>. For example, gain of FGF is implicated in cardiac hypertrophy<sup>16</sup> while manipulation of ETS-1 in mice recapitulates some of the most common congenital heart defects in Jacobsen syndrome<sup>17</sup>. In *Ciona*, expression of a constitutively active form of ETS (ETS-VP16) in non-heart muscle cells leads to development of a multi-chambered heart by recruitment of more cardiac progenitor cells to the ventral midline<sup>18</sup>.

*Ciona robusta* (*Ciona*) is a marine chordate and member of the urochordates, the sister group to vertebrates<sup>19</sup>. Like vertebrates, *Ciona* have a pumping heart within a pericardium, and early heart specification involves the TFs ETS and MesP<sup>20</sup>. Across bilaterians, migration of the heart cells to the ventral midline is dependent on FGF signaling. In *Ciona*, the Trunk Ventral Cells (TVCs) give rise to the heart and pharyngeal cells<sup>21,22</sup>. FGF signaling and MesP activate expression of *FoxF* in the TVCs, this triggers migration of the TVCs to the ventral midline. *Ciona* is experimentally tractable, allowing us to investigate enhancer activity in hundreds of developing embryos and easily visualize the migration of the TVCs to the ventral midline<sup>20</sup>. Thus, *Ciona* heart development and the *FoxF* enhancer provide an ideal system to investigate the relationship between enhancer sequence, tissue-specific gene expression and organismal-level phenotypes.

Developmental enhancers contain suboptimal-affinity sites (also referred to as submaximal and low-affinity sites), which ensure tissue-specific gene expression<sup>23–28</sup>. While the use of suboptimal-affinity binding sites to encode precise gene expression has been seen in a variety of contexts, it has not been explored within heart development. Additionally, in the previous studies, many nucleotide changes are made to the sequences to turn low-affinity sites into high-affinity sites, but these large numbers of changes do not reflect the sequence changes naturally arising within genomic enhancers. Here we seek to determine if low-affinity binding sites are important in heart development, and if single nucleotide variants (SNVs) that increase binding affinity (affinity-optimizing SNVs) could cause gain-of-function gene expression that contributes to phenotypic changes in heart development.

## Results

### The *FoxF* enhancer drives expression in TVCs and contains low-affinity ETS binding sites

*FoxF* is expressed within the TVCs and triggers migration of the TVCs to the ventral midline. A 295bp *FoxF* enhancer upstream of a minimal promoter (bpFog) driving expression of nuclear mCherry recapitulates this TVC specific expression pattern (Figures

1A,E,S1A)<sup>29,30</sup>. An E-box site, ATTA sites and three ETS sites are important for activity of the FoxF enhancer (Figure S1B)<sup>21,31</sup>. To identify additional ETS sites within the FoxF enhancer, we search for ETS cores (GGAW) within the enhancer and identify three additional putative ETS sites. We assign affinities of all six ETS sites from Protein Binding Microarray (PBM) data, which measures the interaction of ETS1 with all possible 8-mers<sup>32,33</sup>. We use PBM data for mouse ETS1 as a proxy for *Ciona* ETS1 binding, as the DNA-binding domain and binding specificity of ETS1 is highly conserved<sup>25,34</sup>. We calculate the relative affinity as the ratio of each 8-mer's fluorescence to the fluorescence of CCGGAAGT, the 8-mer with the highest affinity for ETS1 in *Drosophila*, *Ciona*, mouse, and human<sup>25,33,34</sup>. The putative ETS sites within the FoxF enhancer bind ETS with low relative affinities ranging from 0.09 to 0.24 (Figure S1C).

### Low-affinity ETS sites are necessary for tissue-specific expression

To determine if these six low-affinity ETS sites are necessary for expression within the TVCs, we ablate these ETS sites using point mutations that disrupt hydrogen bonding between the ETS TF and the binding site (GGAW to GCAW)<sup>35</sup>. Ablating all of the ETS sites in parallel leads to complete loss of enhancer activity (Figures 1B,E,S1C). To investigate the contribution of each ETS site to FoxF enhancer activity, we ablate individual ETS sites with point mutations (GGAW to GCAW) and test the effect of these changes in reporter assays by scoring mCherry expression levels in embryos. To the best of our knowledge, these ablations do not create or ablate motifs for TFs present within the TVCs (Table S1). Ablating individual sites in the enhancer leads to a significant reduction in enhancer activity for five ETS sites (Figures S1C,S2). The only site without a reduction in expression upon ablation is ETS6, which has the lowest affinity at 0.09. While loss of the ETS6 site doesn't alter expression, we cannot rule out that this site is redundantly involved in enhancer activity<sup>36–38</sup>. These results demonstrate that five low-affinity ETS sites are necessary for FoxF enhancer activity in the TVCs and illustrate the functional role of low-affinity ETS sites within a developmental heart enhancer.

To determine if the ETS sites within the FoxF enhancer need to be low-affinity for tissue-specific expression, we increase the affinity of the five ETS sites necessary for activity, keeping the GGAW core constant. Optimizing the affinity of all five ETS sites within the enhancer requires 16 nucleotide changes and leads to ectopic expression in tissues receiving FGF signaling, including the anterior tail muscle cells (ATMs), mesenchyme, nervous system, and notochord (Figure 1C,E). To demonstrate that it is the increase of ETS affinity driving the expression change, we made a FoxF enhancer with the 16 nucleotides that optimize ETS affinity in combination with mutations which ablate ETS binding (GGAW to GCAW) (FoxF all opt ablated). This enhancer has no activity (Figures 1D,E). Thus, low-affinity sites are important for maintaining TVC-specific expression of the FoxF enhancer. Similar results have been seen upon optimizing the affinity of sites within neural and notochord enhancers, indicating that the role of low-affinity sites to encode tissue-specific expression is a general regulatory principle governing developmental enhancers<sup>25,26</sup>.

## Affinity-optimizing SNVs lead to loss of tissue-specific expression

To determine if any Single Nucleotide Variants (SNVs) could optimize ETS binding affinity and alter gene expression, we make every possible SNV in the flanking region of the ETS sites of the FoxF enhancer *in silico* and calculate the resulting affinity change. Within this manuscript, we use SNVs to denote single base-pair changes, regardless of whether they are synthetic or naturally found within *Ciona* or human populations. We find that SNVs in three of the five ETS sites cause a 3-fold increase in ETS affinity (Figure 2A).

The greatest affinity change occurs within the ETS1 site which has a starting affinity of 0.12, a SNV increases affinity 8-fold creating an almost consensus site, with 0.97 affinity. We demonstrate the affinity change using electrophoretic mobility shift assay (EMSA) to detect differential binding of *Ciona* ETS1 to the WT and affinity-optimized ETS1 sequences (Figure S3). When we make this SNV within the FoxF enhancer (FoxF-ETS1-T-to-C), the enhancer drives ectopic mCherry expression in the ATMs and mesenchyme, both cell types which receive FGF signaling (Figures 2C,H).

We also find affinity-optimizing SNVs within the ETS3 and ETS4 sites. A SNV within the ETS3 site causes a 3-fold increase in affinity, from 0.11 to 0.34. This affinity-optimizing SNV drives ectopic expression in ATMs, mesenchyme, and neural tissues (Figures 2D,H). The affinity-optimizing SNV within the ETS4 site increases affinity from 0.17 to 0.52, this 3-fold affinity increase causes ectopic expression in the ATMs and mesenchyme (Figures 2E,H). These results suggest that affinity-optimizing SNVs are sufficient to cause loss of tissue-specific gene expression.

A FoxF enhancer containing all three affinity-optimizing SNVs (FoxF 3 SNVs) drives strong activity in tissues responding to FGF signaling, similar to the FoxF all opt enhancer and the effect of these three SNVs appears to be synergistic (Figures 2F,H). A FoxF enhancer with all three affinity-optimizing SNVs and ablated ETS cores (GGAW to GCAW) drives no expression, demonstrating that the effect of these SNVs is ETS-dependent (Figures 2G,H).

## Affinity-optimizing SNVs cause migration defects and disrupt heart development

All three enhancers containing a single affinity-optimizing SNV show ectopic reporter expression in the ATMs (Figure 2). The ATMs and TVCs are related cell types that both receive FGF9 ligand; however, the TVCs contain higher levels of activated ETS due to FGF receptor enrichment<sup>39,40</sup>. The high levels of activated ETS within the TVCs causes TVC-specific expression of *FoxF* and migration of the TVCs to the ventral midline<sup>18</sup>. As the ATMs have low levels of activated ETS, they do not express *FoxF* and do not migrate (Figure 3A; Movie S1)<sup>18</sup>. Overexpression of constitutively active ETS or FoxF in the ATMs causes the ATMs to migrate with the TVCs<sup>18,21</sup>. We therefore hypothesize that the ectopic ATM expression driven by FoxF enhancers containing affinity-optimizing SNVs could cause migration of the ATMs to the ventral midline.

To test this hypothesis, we co-electroporate a *Mesp*-2kb>GFP reporter to mark both the TVCs and ATMs and a construct containing the FoxF 3kb regulatory element, including the endogenous promoter, driving *FoxF* mRNA (Figure S4A). We use the 3Kb regulatory element rather than a minimal enhancer to better recapitulate the regulatory region. When

the wild-type FoxF enhancer drives *FoxF*, we observe no migration of ATMs to the ventral midline (Figures 3A,C,S4; Movie S1) and juvenile *Ciona* have normal hearts (Figure 3E,G; Movie S2). When we electroporate the FoxF-ETS1-T-to-C enhancer driving *FoxF*, we see migration defects in 9.3% of embryos (Figures 3B,D,S4; Movie S1). The FoxF-ETS3-T-to-G enhancer driving *FoxF* causes migration defects in 9.9% of embryos, despite the fact this SNV only increases the affinity of the site to 0.34 (Figure S4). The FoxF-ETS4-T-to-C driving *FoxF* causes migration defects in 8.0% of embryos (Figure S4). Thus, SNVs that lead to 3-fold increase in affinity, even those that result in relatively low-affinity sites, have the potential to contribute to migration defects during heart development. The strong impact of the optimized ETS3 site on migration could be due to nearby E-box and ATTA sites<sup>21,31</sup>, suggesting that enhancer grammar<sup>41</sup> may influence the functional consequences of affinity-optimizing SNVs within genomes.

To determine the impact of migration defects on heart development, we followed 52 FoxF-ETS1-T-to-C embryos with migration defects from the larval to juvenile stages. Strikingly, 79% of embryos with migration defects have abnormally developed hearts (Figures 3F,H). Phenotypes range from enlarged hearts to two distinct beating hearts within the same pericardium (Movie S2). As the *Ciona* age, hearts from animals with the FoxF-ETS1-T-to-C SNV could not pump blood and these animals did not survive beyond twelve days post metamorphosis.

### **Low-affinity ETS sites are prevalent in putative developmental heart enhancers across chordates and these enhancers are vulnerable to affinity-optimizing SNVs.**

The role of FGF signaling in specification and migration of heart cells is conserved from flies to vertebrates<sup>10,11</sup>. To determine if low-affinity sites are common within developmental heart enhancers, we used epigenomic datasets to define putative developmental heart enhancers (DHEs) in both *Ciona* and human. Using ATAC-seq data<sup>42</sup> for *Ciona* heart cells at 6.5hr post fertilization, a time at which TVCs receive FGF signaling, we identify 15,174 putative *Ciona* DHEs. Using ATAC-seq and ChIP-seq for p300 and H3K27Ac from primary fetal human hearts and iPSC-derived embryonic-like cardiomyocytes at time points when FGF signaling is important for heart development, we find 252,931 putative human DHEs<sup>43–46</sup>. In both *Ciona* and human DHEs, the median affinity of putative ETS sites is 0.12, suggesting that low-affinity sites are prevalent within chordate putative DHEs. Within 6,618 (44%) of *Ciona* DHEs and 154,685 (61%) of human putative DHEs, a SNV can increase the affinity of ETS sites by 3-fold with the average affinity after optimization being 0.52 (Figure 4A). Thus, DHEs are vulnerable to affinity-optimizing SNVs, some of which may lead to GOF gene activity and changes in cellular identity.

### **Human and mouse *Gata4* heart enhancers contain low-affinity ETS sites**

GATA4 is a key TF within the developing heart. One of the human DHEs is the ortholog of the mouse enhancer for *Gata4* (Figure 4B). The mouse *Gata4* G9 enhancer is ETS-dependent and drives expression in the endocardial and myocardial heart layers<sup>35</sup>. Ablating four ETS sites within the mouse *Gata4* G9 enhancer leads to loss of expression, suggesting that these sites are functional and required for enhancer activity<sup>35</sup>. However, the affinity of these ETS sites has not been studied. We assign the affinity of all ETS sites within the



*Gata4* G9 enhancer using mouse ETS1 PBM data. All of the ETS sites within the mouse *Gata4* G9 enhancer have affinities ranging from 0.09 and 0.12. The four sites previously shown to contribute to enhancer activity and bind ETS-1 have affinities of 0.11, 0.12, 0.10, and 0.12, respectively. These findings demonstrate that low-affinity ETS sites are necessary for heart-specific expression. The human *GATA4* G9 enhancer contains low-affinity binding sites that are almost identical to those within the mouse *Gata4* G9 enhancer. A reporter construct containing the human *GATA4* G9 enhancer, basal SCP promoter and reporter (GFP and barcode) drives expression in human iPSC-derived fetal-like cardiomyocytes, which are the primary cell type found within the myocardial layers of the heart (Figure 4C).

### **Affinity-optimizing SNVs within the human *GATA4* G9 enhancer cause gain-of-function expression in human iPSC-derived cardiomyocytes**

To determine if optimizing the affinity of the ETS sites within the *GATA4* G9 enhancer leads to GOF expression, in this case an increase in expression within cardiomyocytes, we conducted a reporter assay in human iPSC-CMs to compare the expression of the reference human *GATA4* G9 enhancer (Hs *GATA4*) to the same enhancer with a single (Hs *GATA4* SNVopt) or four affinity-optimizing mutations (Hs *GATA4* 4Opt)<sup>47,48</sup>. Increasing the affinity of a single ETS site from 0.11 to 0.73 (Hs *GATA4* SNVopt) leads to a significant increase in expression. Increasing the affinity of four ETS sites (Hs *GATA4* 4opt) leads to an even larger increase in expression (Figure 4C). Thus, in the human *GATA4* G9 heart enhancer, affinity-optimizing variants drive gain-of-function gene expression, consistent with our findings in *Ciona*.

## **Discussion**

### **Low-affinity ETS sites are a prevalent feature of developmental heart enhancers**

Our genome-wide analysis finds low-affinity sites are a common feature of developmental heart enhancers across *Ciona*, mouse and human. Ablation of low-affinity sites both within the *Ciona* FoxF enhancer and mouse *Gata4* enhancer cause a significant reduction in enhancer activity, demonstrating that these low-affinity sites are functional. Low-affinity sites are also important within enhancers activated by other TFs downstream of signaling pathways and pleiotropic factors<sup>23,25,49–51</sup>. Our findings highlight the importance and vastly underappreciated contribution of very low-affinity binding sites to regulation of gene expression and the advantages of scoring the affinity of sites.

### **Affinity-optimizing SNVs cause gain-of-function enhancer activity, which can contribute to phenotypes**

In *Ciona*, affinity-optimizing SNVs lead to loss of tissue-specific expression and migration defects that alter heart development, in the most extreme cases we see two beating hearts. We propose that ETS affinity-optimizing SNVs likely contribute to congenital heart disease and cardiac traits. The ubiquitous role of FGF in other developmental programs<sup>52,53</sup> and cancer<sup>54</sup> indicates that ETS affinity-optimizing SNVs are likely involved in other enhanceropathies. Searching for affinity increases as a method to filter for causal enhancer variants is not common, yet from our work and a few other studies it appears this could be a powerful approach. To our knowledge, only three studies search for an affinity increase



and measure the impact on gene expression, yet do not analyze phenotype<sup>55–57</sup>. In a handful of examples, causal enhancer variants have been functionally validated via reporter assays. After the validation of the impact of these variants on gene expression, these sequence changes have been identified as increasing affinity. Although using affinity to filter for causal variants is in its infancy, this approach appears to generalize beyond ETS as these other aforementioned studies have found sequence changes that increase affinity in TFs as diverse as Wnt effectors and Hox members that can alter gene expression and are associated with phenotypes<sup>58–61</sup>.

### Affinity-optimizing SNVs may contribute to evolution of novel traits

There is a fine line between creation of novel phenotypes that are beneficial or detrimental<sup>5</sup>. The *Ciona* heart, like all invertebrate chordates, is single-chambered, while all extant vertebrates have at least two chambers. The evolution of a dual-chambered heart in vertebrates is thought to involve recruitment of additional precursor cells to the ventral midline<sup>18</sup>. Our study indicates that mutations within enhancers that increase binding affinity for ETS result in more cells migrating to the ventral midline and creation of another compartment within the pericardium. Although the animals in our study with two hearts did not survive, it is possible that some animals with multi-chambered hearts could survive and have a selective advantage. More generally, affinity-optimizing SNVs could contribute to the evolution of traits by affecting gene expression patterns and cell behavior.

### Violations in regulatory principles governing enhancers can pinpoint causal enhancer variants

How enhancer variants contribute to phenotypes is a pressing challenge we need to solve if we want to causally link genomic variation to phenotypic diversity in health and disease. To date, only a handful of enhancer variants have been functionally validated as altering gene expression and phenotype. Yet thousands of non-coding variants are associated with changes in gene expression and phenotypic variation. We cannot possibly test all non-coding variants in the relevant cell types and time points to pinpoint causal enhancer variants. Finding rules that generalize across different enhancers relies on understanding the regulatory principles that govern enhancer function. One such regulatory principle is the use of low-affinity binding sites to encode tissue-specific expression and levels of expression. Here we show that SNVs that violate this regulatory principle can cause gain-of-function gene expression and organismal-level phenotypic changes. Identification of other regulatory principles and subsequently variants that violate these principles could provide further mechanistic approaches to pinpoint causal enhancer variants at scale.

### Limitations of the Study

While many studies use position weight matrices (PWM) to find matches to binding sites, here we use PBM data, an *in vitro* measurement of relative binding affinity that allows us to give a relative affinity score to every 8mer sequence. *In vitro* PBM measurements do not consider chromatin structure, protein-protein interactions, cooperative binding between multiple sites and phase separation. While this is a limitation, it also has some advantages as it enables us to have a fixed measurement of relative binding affinity that can be used across genomes, cell types and species to annotate putative binding sites. As illustrated in our

study, this approach can predict binding and the impact of SNVs in different cell types and organisms (e.g FoxF *Ciona* TVC enhancer and human *GATA4* cardiomyocyte enhancer). As such, it is a score that can be used to analyze genomic sequences and predict the impact of enhancer SNVs without the need to measure specific binding affinity of every site at every time point and cell type of an organism.

Our studies take advantage of disparate biological systems *Ciona* and human cardiomyocytes to detect gain-of-function gene expression. In *Ciona*, we are able to assay expression in all cells of the embryo, but it is difficult to quantify expression levels within these cell types. While in human cells we see that the affinity-optimizing SNV increase levels of expression, however, we cannot assay the impact of this SNV on specificity of gene expression within the entire human heart. Together, these complementary assays illustrate that affinity-optimizing SNVs can increase the levels of expression within the endogenous location and also lead to expression in other cell types, both types of GOF expression could contribute to phenotypes.

Another limitation relates to how we study the role of enhancer variants on cell behavior. We characterize *Ciona* heart phenotypes by electroporation of a plasmid carrying a large 3kb regulatory region driving *FoxF* cDNA into fertilized eggs. While the WT 3kb region driving *FoxF* mRNA does not alter migration, the same regulatory element with only a single SNV leads to migration defects. Indeed, we see consistent results for three different SNVs tested in this way. Ideally, we would show that a SNV within the endogenous locus has the same effect. However, targeted genome editing with HDR, or base editing is not possible within *Ciona*. Once these techniques are established, testing the impact of the FoxF SNVs within the genome will be important. While we have not tested the FoxF SNVs in the endogenous locus, we have done similar studies in mice and find that affinity-optimizing SNVs within the endogenous locus in mice lead to GOF expression and phenotypes<sup>62</sup>.

## STAR Methods

### RESOURCE AVAILABILITY

**Lead contact**—Further information and requests for resources and reagents should be directed to and will be fulfilled by the lead contact, Emma Farley (efarley@ucsd.edu).

**Materials availability**—Plasmids generated in this study will be made available from lead contact upon request.

#### Data and code availability

- Sequencing data have been deposited at SRA and are publicly available as of the date of publication. Accession numbers are listed in the key resources table.
- Microscopy and scoring data reported in this paper will be shared by the lead contact upon request.
- All original code has been deposited at GitHub and is publicly available as of the date of publication. DOI is listed in the key resources table.

- Bed files of putative *Ciona* developmental heart enhancers and putative human developmental heart enhancers have been deposited at GitHub and DOI is listed in the key resources table
- Any additional information required to reanalyze the data reported in this paper is available from the lead contact upon request.

## EXPERIMENTAL MODEL AND STUDY PARTICIPANT DETAILS

**Tunicates**—Adult *C. intestinalis* type A aka *Ciona robusta* (obtained from M-Rep) were maintained under constant illumination in seawater (obtained from Reliant Aquariums) at 18°C. *Ciona* filter feed on material in seawater and we do not add any food for the week we keep them for. *Ciona* are hermaphroditic, therefore there is only one possible sex for individuals. Age or developmental stage of the embryos studied are indicated in the main text.

**iPSC-derived cardiomyocytes**—iPSC-CMs from line UDID\_109 and from line UDID\_139<sup>47</sup> were thawed and seeded onto gelatin-coated 48-well plates at a density of  $1.3 \times 10^5/\text{cm}^2$ . UDID\_109 cell line is from an Asian male and UDID\_139 cell line is from an Asian female. Cells were cultured using previously described culture conditions<sup>63</sup>.

## METHOD DETAILS

**Mutagenesis and cloning**—The FoxF enhancer construct was a gift from the Levine and Christiaen labs. The construct consists of the following elements in order: AscI, FoxF enhancer, XbaI, bp10g promoter, NotI, H2B-mCherry, EcoRI. Point mutations were introduced into the construct using mutagenesis with partially overlapping primers with 3'-overhangs. To clone the construct with all 6 ETS sites ablated from GGAW to GCAW, we ordered the FoxF enhancer synthesized from IDT. For experiments using the 3kb FoxF enhancer and *FoxF*cDNA, the 3kb FoxF regulatory region was amplified from the genome and cloned into our electroporation vector, the *FoxF*cDNA was also cloned into this vector. The *FoxF*cDNA sequence can be found in Aniseed with the Gene Model ID KH2012:KH.C3.170. The *GATA4* G9 reporter plasmid for testing in iPSC-CMs consists of the following elements in order: *GATA4* fragment, supercore promoter, unc76-EGFP, 30-nucleotide barcode and polyA.

**Electroporation**—Constructs to be compared to each other are midiprepmed in the same batch using the NucleoBond Xtra Midi kit (Macherey-Nagel). For each construct, 70 µg of DNA in 100 µL is added to 400 µL of 0.96 M D-mannitol. Dechoriation, *in vitro* fertilization and electroporation were performed as described previously<sup>64</sup>. Typically for each electroporation, eggs and sperm were collected from 6 adults. Embryos were fixed at the appropriate developmental stage for 15 minutes in 3.7% formaldehyde. The tissue was then cleared in a series of washes of 0.3% Triton-X in PBS and then of 0.01% Triton-X in PBS. Samples were mounted in Prolong Gold. Differential interference contrast microscopy was used to obtain transmitted light micrographs with an Olympus FV3000, using the 40X objective. The same microscope was used to obtain all fluorescent images. Three biological replicates were analyzed for each construct unless otherwise noted.

**Counting embryos**—In *Ciona*, as there are defined cell lineages, we can determine cell identity based on visually inspecting the nuclear signal and the region of the embryo in which the signal occurs. Strong, moderate, and weak expression levels through the eyepiece were determined as strong being visible at 25x of light source, moderate being visible at 50x of light source, and weak being visible at 100x of light source. Fifty embryos were counted for each biological replicate, unless otherwise noted. For each experiment, slides were counted blind.

**Acquisition of Images**—For enhancers being compared, images were taken from electroporations performed on the same day using identical settings. For representative images, embryos were chosen that represented the average from counting data. Representative embryos on the slide were chosen and imaged prior to unblinding the slides. Two exposure times were taken for each construct. In each figure, the same exposure time for each image is shown to allow direct comparison, which occasionally leads to overexposed images being used for stronger constructs (e.g., Fig 1C).

**Electrophoretic mobility shift assay**—EMSAs were performed using the LightShift™ Chemiluminescent EMSA Kit (ThermoFisher Scientific) with biotinylated and non-biotinylated double-stranded oligonucleotides corresponding to the ETS1 region of the FoxF enhancer and mutations that alter affinity to 0.97 and 1.00. Oligonucleotides were annealed according to the advanced protocol at this link (<https://tools.thermofisher.com/content/sfs/brochures/TR0045-Anneal-oligos.pdf>). *Ciona* ETS1 DBD protein was synthesized using the TNT Quick Coupled Transcription/Translation System (Promega) from the pTNT plasmid containing *Ciona* ETS1 DBD. *Ciona* ETS1 DBD was synthesized with flanking XhoI and NotI sites using Twist Bioscience and cloned into the pTNT-B18R vector (Addgene #58978). The *Ciona* ETS1 DBD sequence was determined as the sequence of *Ciona* ETS1 that aligns to Mouse ETS1 DBD<sup>65,66</sup>. The binding reaction was carried out in a 20 µL volume containing 2 µL of 10X Binding Buffer (100 mM Tris, 500 mM KCl, 10 mM DTT; pH 7.5), 50 ng Poly(dI:dC), 4 µL of protein extract and 20 femtomol biotin-labeled probe. For competition experiments, a 200-fold molar excess of unlabeled probe was added. Binding reactions were pre-incubated for 10 minutes before adding the biotin-labeled probe. Binding reactions were then incubated at room temperature for 20 min and loaded onto a DNA retardation gel (6%). After sample electrophoresis with 0.5X TBE on ice and transfer to a 0.45-µm Biotinylated B Nylon membrane (ThermoFisher Scientific) in the cold room, DNA was crosslinked for 15 minutes using 312 nm light, and the membrane was put between blank sheets of paper overnight. The next day, the biotinylated probes were detected using the Chemiluminescent Nucleic Acid Detection Module (ThermoFisher Scientific). Images of the resulting membrane were acquired using a Chemidoc MP imaging system (Bio-Rad).

**Counting Larvae for Migration Defects**—Embryos were transferred from gelatin-coated electroporation plates to 15cm uncoated plastic plates filled with filtered seawater at 7 hpf. All well-developed larvae on the plate were counted from 16–20 hpf and scored for presence or absence of normal ATM development. All larvae with migration defects were marked for heart morphology analysis in the juvenile stage. For each replicate, 10–20 larva

with normal ATM development were selected for heart morphology analysis in the juvenile stage.

**Juvenile Heart Morphology Analysis and Imaging**—Juvenile hearts were analyzed for morphological defects starting at 48hrs after metamorphosis. Hearts with abnormal structure or function were scored as “deformed.” Images and video of hearts were taken on an Axiozoom microscope with a mounted iPhone 7. Video editing was done in Adobe Premiere Pro. Heartbeat animations were made in Adobe Animate.

**TVC Migration Time-lapse Imaging**—Embryos were electroporated with MesP 2kb>GFP as previously described and checked for GFP fluorescence at 6 hpf. Well-developed embryos with strong GFP expression were transferred to a glass-bottom microscopy plate in seawater. Z-stacks were taken of each embryo every 30 minutes for 8 hours with an Olympus FV3000 microscope with a 20x objective lens. Resulting time-lapses were analyzed in Fiji. A max projection was taken of the GFP channel and merged with the DIC. Time-lapses were cropped and rotated in Adobe Photoshop.

**Scoring Relative Affinities**—Relative affinities were calculated using high-throughput binding datasets<sup>67</sup> ([thebrain.bwh.harvard.edu/uniprobe/index.php](http://thebrain.bwh.harvard.edu/uniprobe/index.php)). They were calculated using median signal intensities of mouse ETS1 universal protein binding microarray data from the UniProbe database<sup>33</sup>. It has previously been shown that the binding specificity of ETS1 across 600 million years from flies to humans is conserved and thus the use of mouse ETS1 to calculate ETS affinity is a valid approach<sup>34</sup>. The relative affinity represents the ratio of the median signal intensity of the native 8-mer motif to the optimal 8-mer motif for ETS.

**Computational Search for Motifs Altered by each SNV**—For each SNV, a 30 base-pair region centered on the ETS site was put into JASPAR and scanned for all 1205 vertebrate profiles with 80% cutoff. Then, the same 30 base-pair region centered on the ETS site with the SNV was put into JASPAR<sup>68</sup> and scanned for all 1205 vertebrate profiles with 80% cutoff. The list of sites destroyed and created by the SNV were found, we further filtered this list to focus on individual transcription factor sites with mutations that affect nucleotides critical for transcription factor binding. Of the ETS-family JASPAR profiles, we used MA0098.1.ETS1. For the GGAW to GCAW ablations, no TFBS for repressors expressed in the TVCs (ANISEED database) were created and no TFBS for activators expressed in the TVCs (ANISEED database) were destroyed. For the SNV opt mutations, no TFBS for repressors expressed in the ATMs (ANISEED database) were destroyed and no TFBS for activators expressed in the ATMs (ANISEED database) were created.

## ATAC-seq Data Analysis

**Alignment of ATAC-seq reads.** ATAC-seq alignment was performed following the methods in<sup>42</sup>. Raw reads from 6 hours post fertilization (hpf) (GEO accession GSE126691, LacZ\_6hpf\_1–3, LacZ\_10hpf\_1–4) were first preprocessed by FastQC (version 0.11.2, <http://www.bioinformatics.babraham.ac.uk/projects/fastqc>). Adaptors were trimmed using Trim Galore (version 0.4.4, [http://www.bioinformatics.babraham.ac.uk/projects/trim\\_galore](http://www.bioinformatics.babraham.ac.uk/projects/trim_galore))

and trimmed reads were aligned to the *Ciona robusta* genome<sup>69</sup> using Bowtie2 (version 2.3.2,<sup>70</sup>) with the parameters `--very-sensitive -p 16 -X 1000`. Reads with mapping quality score > 30 were kept for downstream analysis using SAMtools (version 1.2,<sup>71</sup>). Mitochondrial reads were removed using bash command `egrep -v`. Following examination of read quality of the 6hpf data, replicate 2 was removed. Read pileup (BAM) files for replicates 1 and 3 at 6hpf were merged using the SAMtools merge function and peaks were called using MACS2 (version 2.7.9)<sup>72</sup> (`--nomodel --bdg --g 99000000 -f BAMPE -q 0.01`). To correct for nonspecific sequencing biases, we subtracted gDNA from these libraries<sup>73</sup>.

**Calculating top ATAC peaks.:** To find the highest confidence peaks in our dataset, we calculated the area under the curve (AUC) as the sum of the read depth at each position in the peak. Bedtools genomecov (-bga) and intersect functions were used to calculate the read depth at every genomic position within all the called peaks<sup>74</sup>. Peaks with AUCs in the top 90% were kept and the rest were discarded.

**Downloading putative enhancers.:** Human fetal heart putative enhancers were identified from epigenomic datasets from the key resource table. Any coordinates for hg18 or hg19 were lifted over to hg38 using pyliftover 0.4 (<https://pypi.org/project/pyliftover>).

**Identifying affinity optimizing SNVs in putative developmental heart enhancers.:** We first determined the distribution of ETS affinities in putative developmental heart enhancers. We collected affinities for all ETS sites defined as NNGGAWNN. Once all of the ETS binding site affinities were determined, we then performed all possible point mutations to each binding site and plotted those which resulted in 3-fold change. We reported the number of putative heart enhancers which contained 1 ETS which could be affinity-optimized 3 fold by a SNV.

**Library Construction—**A library containing Hs *GATA4*, Hs *GATA4* SNVopt and Hs *GATA4* 4opt enhancers upstream of the supercore promoter, GFP and barcode, was constructed as previously described in<sup>75</sup>.

**iPSC-CM transfection and collection—**iPSC-CMs from line UDID\_109 and from line UDID\_139<sup>47</sup> were thawed and seeded onto gelatin-coated 48-well plates at a density of  $1.3 \times 10^5/\text{cm}^2$ . The iPSC-CMs were transfected with 325nM/well of plasmid containing the enhancer library driving GFP using ViaFect<sup>TM</sup> Transfection Reagent (Promega) at a 6:1 ratio in a medium containing 10% FBS and 5uM (Y27632) ROCK Inhibitor (Sigma). 16hrs post transfection and every other day, medium was changed. Cells were then collected 96hrs post transfection and stored in TriZol reagent frozen at  $-80^\circ\text{C}$ . Library transfection was performed in duplicate for line 109 and triplicate for line 139.

**RNA extraction, cDNA synthesis, DNA extraction, barcode extraction, sequencing—**Total RNA was extracted using TriZol extraction and treated to remove contaminating DNA (TURBO DNA-free DNase digestion kit, Ambion). Subsequent mRNA was isolated (Dynabeads mRNA Isolation kit, Invitrogen) and was used for cDNA synthesis (Transcriptor High Fidelity cDNA Synthesis kit, Sigma-Aldrich). Subsequent



DNA extraction, barcode extraction, and sequencing was performed as previously described<sup>75</sup>.

### **Enhancer to Barcode Assignment, Dictionary Analysis, and SEL-Seq Data**

**Analysis**—Assignment of barcodes to enhancers, dictionary setup and SEL-Seq data analysis was performed as previously described<sup>75</sup>, with the only difference being that assignment of barcodes to enhancers was done using PacBio sequencing. Active enhancers will transcribe the mRNA barcode, thus the mRNA barcodes were measured to calculate enhancer activity. The library was transfected into human iPSC-CMs, samples were collected 96 hours after transfection, and we isolated barcode mRNA and DNA. To normalize the enhancer activity to differences in the amount of plasmid, we calculated  $\log_2(\text{mRNA RPM} / \text{DNA RPM})$ . This was done in 6 biological replicates across 2 cell lines, with an average Pearson correlation coefficient between replicates of 0.86.

## **QUANTIFICATION AND STATISTICAL ANALYSIS**

For comparisons of counting data, the chi-squared test was used in Excel with the CHISQ.TEST function. For the post hoc test to avoid false positives, Bonferroni correction was used to adjust thresholds for significance. With N comparisons, the Bonferroni-adjusted P value needed for significance is  $0.05/N$ . Enhancer activity scores in Figure 4 were statistically analyzed using a standard two-sided T-test with Benjamini-Hochberg adjustment.

## **Supplementary Material**

Refer to Web version on PubMed Central for supplementary material.

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## **Inclusion and Diversity:**

One or more of the authors of this paper self-identifies as an underrepresented ethnic minority in their field of research or within their geographical location. One or more of the authors of this paper self-identifies as living with a disability.



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**Highlights:**

Low-affinity binding sites are prevalent within developmental heart enhancers

Single nucleotide variants that increase ETS affinity drive excess enhancer activity

Affinity-optimizing enhancer variants disrupt heart development.

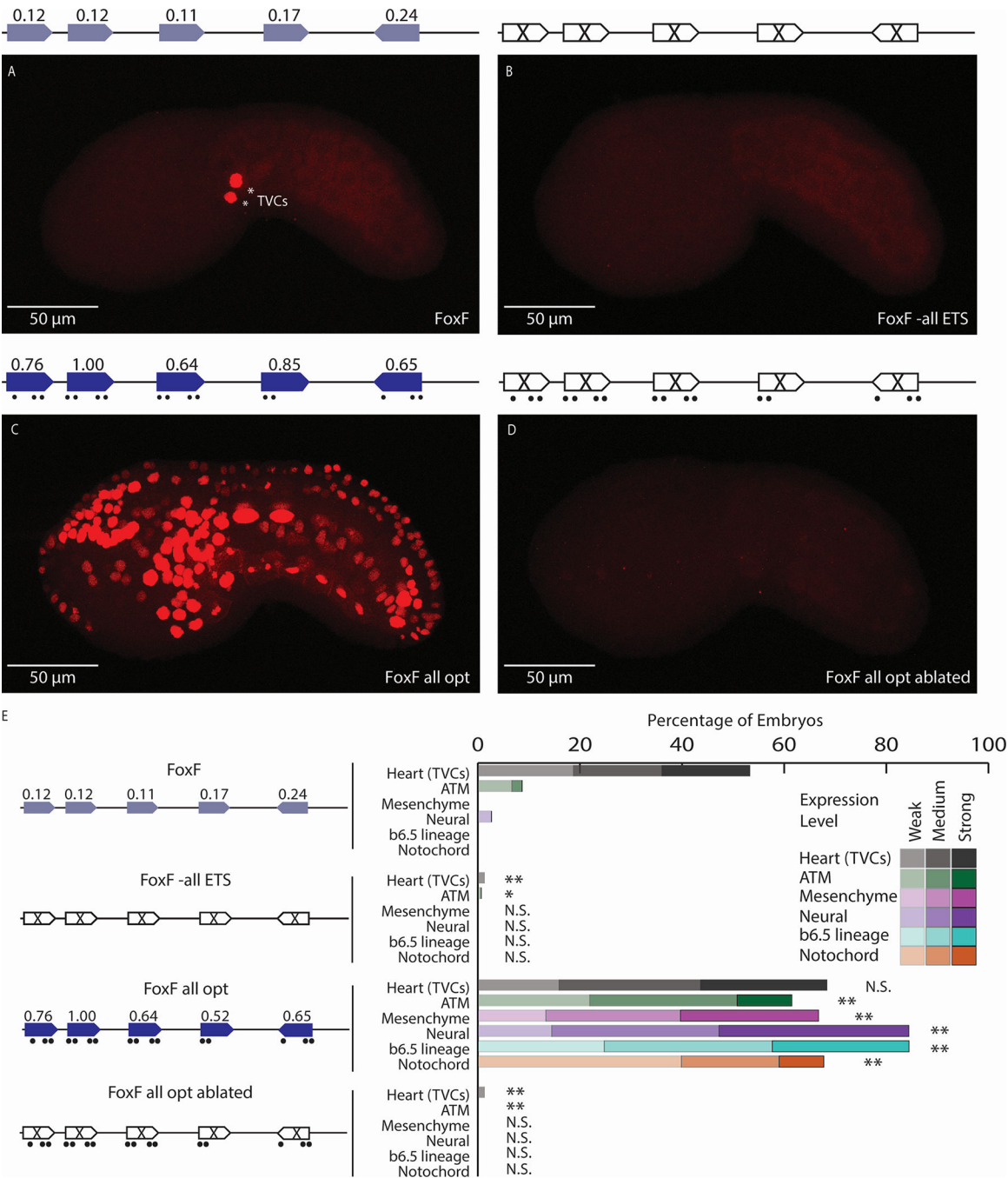
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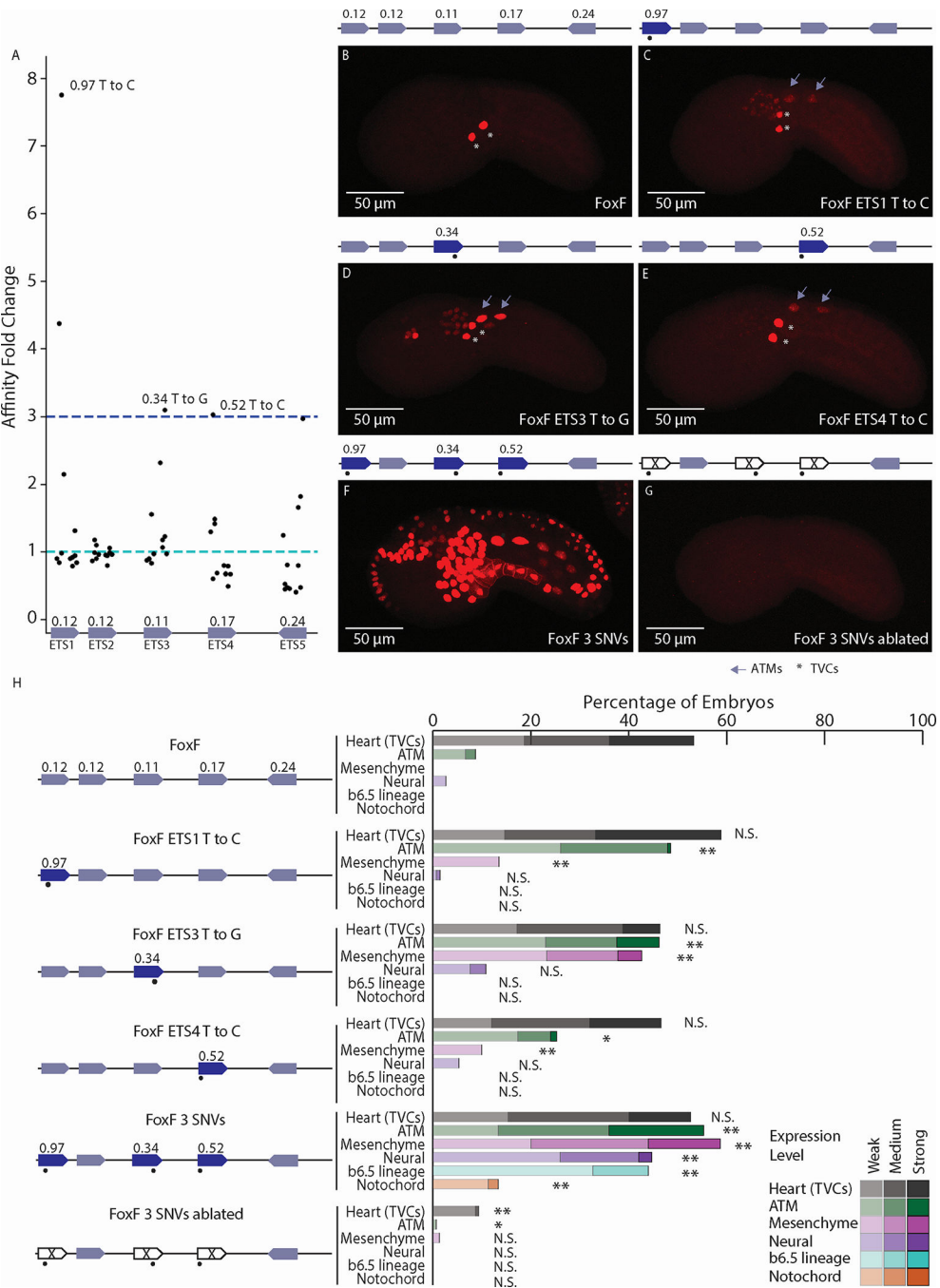




**Figure 1. Low-affinity ETS sites are necessary for tissue-specific expression.**  
**A.** The FoxF enhancer drives expression in the TVCs, marked by asterisks. **B.** An enhancer with all six ETS sites ablated (by changing each site from GGAW to GCAW) drives no enhancer activity. **C.** Optimizing the affinity of all ETS sites leads to ectopic expression in ATMs, mesenchyme, nervous system, and notochord. **D.** Ablating the ETS sites within the context of the FoxF all opt (FoxF all opt ablated) drives no enhancer activity. **E.** Embryo scoring for above constructs (3 replicates, n = 32 embryos per replicate). P-values are obtained using Chi-square test comparing each condition to WT with Bonferroni correction:

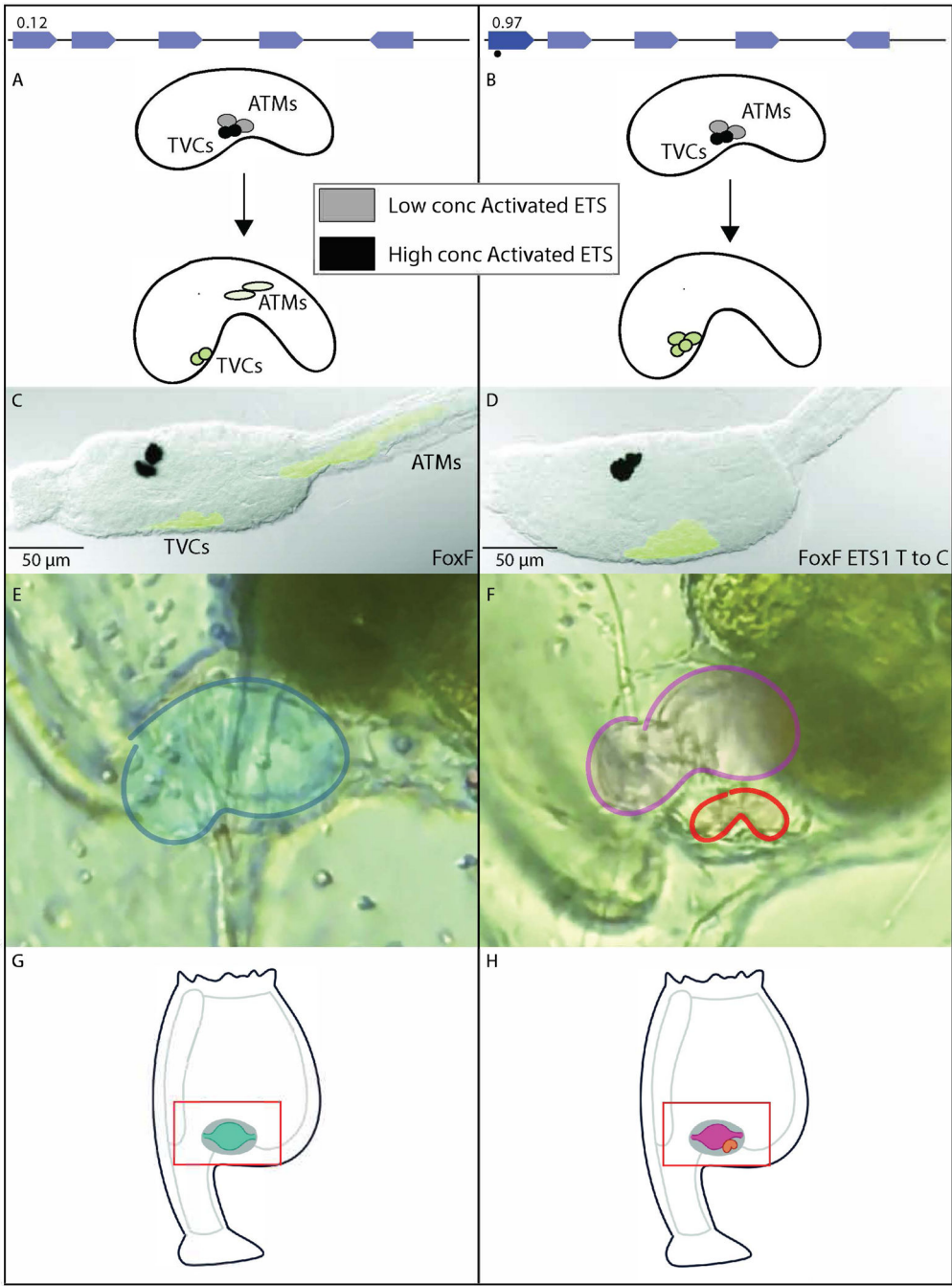


\*\*  $P < 0.0017$ , \*  $P < 0.017$ , N.S. not significant. Schematics show electroporated enhancers, values are affinities, dots show nucleotide changes. ETS6 is not shown in the schematic as ablation of this site did not impact expression (see Figure S1). Scale bars, 50  $\mu\text{m}$ . Data in stacked bar chart are represented as the mean. See also Figures S1,S2 and Tables S1,S2.



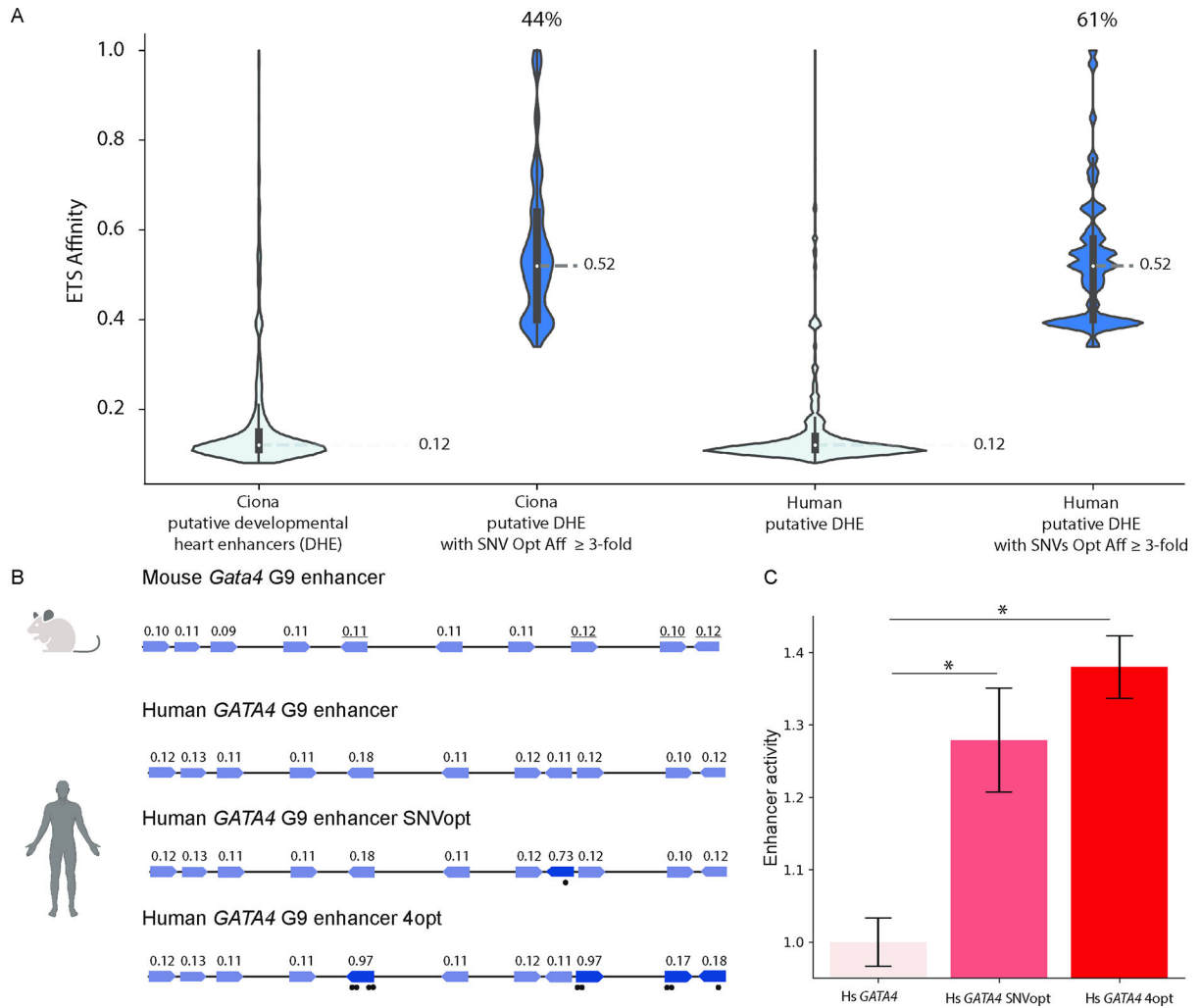
**Figure 2. Affinity-optimizing SNVs lead to loss of tissue-specific expression.**  
**A.** Affinity fold change for SNVs within functional ETS sites of the FoxF enhancer.  
**B.** The WT FoxF enhancer drives expression in the TVCs. **C.** The FoxF-ETS1-T-to-C enhancer drives expression in the TVCs, ATMs, and mesenchyme. **D.** The FoxF-ETS3-T-to-G enhancer drives expression in the TVCs, ATMs, mesenchyme, and neural tissues. **E.** The FoxF-ETS4-T-to-C enhancer drives expression in the TVCs and ATMs. **F.** Optimizing the affinity of three ETS sites with SNVs leads to ectopic expression in ATMs, endoderm, mesenchyme, nervous system, and notochord. **G.** Ablating the ETS sites (GGAW to GCAW)

within the context of the 3 SNVs no longer drives ectopic expression, only very weak TVC enhancer activity. **H.** Embryo scoring for above constructs (3 replicates, n = 17 embryos per replicate). P-values are obtained by using Chi-square test comparing each condition to WT with Bonferroni correction: \*\*  $P < 0.001$ , \*  $P < 0.01$ , N.S. not significant. Asterisks mark TVCs and arrows mark ATMs. Schematics show electroporated enhancers, values are affinities, dots mark SNVs. Scale bars, 50  $\mu\text{m}$ . Data in stacked bar chart are represented as the mean. See also Figures S2,S3 and Tables S1,S2.



**Figure 3. Affinity-optimizing SNVs cause migration defects and disrupt heart development.**  
**A.** Schematic showing a tailbud wild-type embryo, the concentration of activated ETS in the TVCs and ATMs and the role of FoxF in migration of the TVCs to the ventral midline. The TVCs have high levels of activated ETS while the ATMs have lower levels of activated ETS. FoxF enhancer drives expression of *FoxF* in the TVCs, causing them to migrate. **B.** Schematic showing a tailbud embryo containing the FoxF-ETS1-T-to-C enhancer driving *FoxF*. TVC cells have high levels of activated ETS, while ATMs have low levels of activated ETS. Embryos with the FoxF-ETS1-T-to-C enhancer drive expression of *FoxF* in the TVCs

and ATMs causing the ATMs to express *FoxF* and migrate with the TVCs. **C.** Embryo with WT *FoxF* enhancer driving *FoxF*, co-electroporated with a *MesP* enhancer driving GFP marking the ATMs and TVCs at 16 hpf. **D.** Embryo with *FoxF*-ETS1-T-to-C enhancer driving *FoxF*, co-electroporated with a *MesP* enhancer driving GFP in the TVCs and ATMs at 16 hpf; all cells migrate to the ventral midline. Black cells in **C** and **D** are the otolith and ocellus. **E.** Juvenile with WT *FoxF* enhancer driving *FoxF* with heart colored in blue; normal heart morphology is seen. **F.** Juvenile with *FoxF*-ETS1-T-to-C enhancer driving *FoxF* has two hearts with main heart colored in pink, auxiliary heart colored in orange; abnormal heart morphology is seen in 79% (41/52) of embryos with migration defects. Hearts were false-colored digitally. **G.** Schematic of juvenile *Ciona* with normal heart morphology. **H.** Schematic of juvenile *Ciona* with auxiliary heart. Scale bars, 50  $\mu$ m. See also Figure S4 and Movies S1,S2.



**Figure 4. Low-affinity ETS binding sites are prevalent within developmental heart enhancers and affinity-optimizing SNVs lead to gain-of-function expression in human iPSC-CMs.**

**A.** Violin plots in light blue of ETS site affinities within *Ciona* and human putative DHEs. 44% of *Ciona* DHEs and 61% of human DHEs contain an ETS site in which a SNV can increase ETS affinity 3-fold. Dark blue violin plots show distributions of the affinity-optimizing SNVs. **B.** Schematics of human and mouse *Gata4* enhancers show high conservation of low-affinity ETS sites. Sites with underlined affinities mark functionally validated ETS sites in mouse<sup>35</sup>. *GATA4* G9 enhancer SNVopt harbors a SNV that increases relative affinity from 0.11 to 0.73. *GATA4* G9 enhancer 4opt has four ETS sites with increased affinity. **C.** Hs *GATA4*, Hs *GATA4* SNVopt and Hs *GATA4* 4opt enhancer activity as measured by reporter assay in human iPSC-CMs. Data are represented as mean  $\pm$  SEM. \*  $p < 1e-5$ . See also Table S2.

## Key resource table

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Deposited Data		
mouse ETS1 universal PBM data	Wei et al. <sup>33</sup>	<a href="http://thebrain.bwh.harvard.edu/uniprobe/index.php">thebrain.bwh.harvard.edu/uniprobe/index.php</a>
<i>Ciona robusta</i> genome	Satou et al. <sup>69</sup>	N/A
LacZ_6hpf_1–3	Racioppi et al. <sup>42</sup>	GEO: GSE126691
Human reference genome NCBI build 38, GRCh38	Genome Reference Consortium	<a href="http://www.ncbi.nlm.nih.gov/projects/genome/assembly/grc/human">http://www.ncbi.nlm.nih.gov/projects/genome/assembly/grc/human</a>
Human fetal heart putative enhancers predicted from H3K27ac ChIP and ATAC-seq	Richter et al. <sup>43</sup>	Supplementary Table 11 in source
iPSC-cardiomyocyte putative enhancers predicted from ATAC-seq and H3K27ac ChIP	Benaglio et al. <sup>44</sup>	GEO: GSE133833
Human fetal heart putative enhancers predicted from p300/CBP	May et al. <sup>45</sup>	GEO: GSE32587
Sequencing data for MPRA	This paper	SRA: PRJNA999706
Experimental Models: Organisms/Strains		
<i>Ciona intestinalis</i> type A ( <i>Ciona robusta</i> )	M-REP	N/A
iPSC-CMs	Frazer Lab	UDID_109 <sup>48</sup>
iPSC-CMs	Frazer Lab	UDID_139 <sup>48</sup>
Oligonucleotides		
Oligonucleotides for mutagenesis, see Table S2	This paper	N/A
Recombinant DNA		
Plasmid: foxf295-bpfog>H2B-mCherry	Levine & Christiaen Labs	N/A
Plasmid: foxf295-ets1mut-bpfog>H2B-mCherry	This paper	N/A
Plasmid: foxf295-ets2mut-bpfog>H2B-mCherry	This paper	N/A
Plasmid: foxf295-ets3mut-bpfog>H2B-mCherry	This paper	N/A
Plasmid: foxf295-ets4mut-bpfog>H2B-mCherry	This paper	N/A
Plasmid: foxf295-ets5mut-bpfog>H2B-mCherry	This paper	N/A
Plasmid: foxf295-ets6mut-bpfog>H2B-mCherry	This paper	N/A
Plasmid: foxf295-ets1optT2C-bpfog>H2B-mCherry	This paper	N/A
Plasmid: foxf295-ets3optT2G-bpfog>H2B-mCherry	This paper	N/A
Plasmid: foxf295-ets4optT2C-bpfog>H2B-mCherry	This paper	N/A
Plasmid: foxf295-ets134opt-bpfog>H2B-mCherry	This paper	N/A
Plasmid: foxf295-ets134optAbl-bpfog>H2B-mCherry	This paper	N/A
Plasmid: foxf295-ets12345opt-bpfog>H2B-mCherry	This paper	N/A
Plasmid: foxf295-ets12345optAbl-bpfog>H2B-mCherry	This paper	N/A
Plasmid: foxf295-ets123456mut-bpfog>H2B-mCherry	This paper	N/A
Plasmid: foxf3kb>foxfcDNA	This paper	N/A
Plasmid: foxf3kb-ets1OptT2C>foxfcDNA	This paper	N/A
Plasmid: foxf3kb-ets3OptT2G>foxfcDNA	This paper	N/A
Plasmid: foxf3kb-ets4OptT2C>foxfcDNA	This paper	N/A



REAGENT or RESOURCE	SOURCE	IDENTIFIER
Plasmid: gata4-g9-scp>unc76-GFP	This paper	N/A
Plasmid: gata4-g9-SNVopt-scp>unc76-GFP	This paper	N/A
Plasmid: gata4-g9-4opt-scp>unc76-GFP	This paper	N/A
Plasmid: pTNT-ciETS1	This paper	N/A
Software and Algorithms		
ImageJ	Schneider et al. <sup>76</sup>	<a href="https://imagej.nih.gov/ij/">https://imagej.nih.gov/ij/</a>
FastQC (version 0.11.2)	N/A	<a href="http://www.bioinformatics.babraham.ac.uk/projects/fastqc">http://www.bioinformatics.babraham.ac.uk/projects/fastqc</a>
Trim Galore (version 0.4.4)	N/A	<a href="http://www.bioinformatics.babraham.ac.uk/projects/trim_galore">http://www.bioinformatics.babraham.ac.uk/projects/trim_galore</a>
Bowtie2 (version 2.3.2)	Langmead et al. <sup>70</sup>	<a href="http://bowtie-bio.sourceforge.net/bowtie2/index.shtml">http://bowtie-bio.sourceforge.net/bowtie2/index.shtml</a>
Samtools (version 1.2)	Li et al. <sup>71</sup>	<a href="https://samtools.sourceforge.net">https://samtools.sourceforge.net</a>
MACS2 (version 2.7.9)	Zhang et al. <sup>72</sup>	N/A
BEDTools	Quinlan et al. <sup>74</sup>	N/A
pylifter 0.4	N/A	<a href="https://pypi.org/project/pylifter">https://pypi.org/project/pylifter</a>
Adobe Premiere Pro	Adobe	N/A
Adobe Animate	Adobe	N/A
Code to generate Fig 2A, 4A, 4C and putative <i>Ciona</i> developmental heart enhancers and putative human developmental heart enhancers	This paper	<a href="https://doi.org/10.5281/zenodo.8215976">https://doi.org/10.5281/zenodo.8215976</a>