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Journal

Human Molecular Genetics, 24(14)

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Publication Date

2015-07-15

DOI

10.1093/hmg/ddv047

Peer reviewed

ORIGINAL ARTICLE

A single nucleotide polymorphism associated with isolated cleft lip and palate, thyroid cancer and hypothyroidism alters the activity of an oral epithelium and thyroid enhancer near *FOXE1*

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Abstract

Three common diseases, isolated cleft lip and cleft palate (CLP), hypothyroidism and thyroid cancer all map to the *FOXE1* locus, but causative variants have yet to be identified. In patients with CLP, the frequency of coding mutations in *FOXE1* fails to account for the risk attributable to this locus, suggesting that the common risk alleles reside in nearby regulatory elements. Using a combination of zebrafish and mouse transgenesis, we screened 15 conserved non-coding sequences for enhancer activity, identifying three that regulate expression in a tissue specific pattern consistent with endogenous *foxe1* expression. These three, located –82.4, –67.7 and +22.6 kb from the *FOXE1* start codon, are all active in the oral epithelium or branchial arches. The –67.7 and +22.6 kb elements are also active in the developing heart, and the –67.7 kb element uniquely directs expression in the developing thyroid. Within the –67.7 kb element is the SNP rs7850258 that is associated with all three diseases. Quantitative reporter assays in oral epithelial and thyroid cell lines show that the rs7850258 allele (G) associated with CLP and hypothyroidism has significantly greater enhancer activity than the allele associated with thyroid cancer (A). Moreover, consistent with predicted transcription factor binding differences, the –67.7 kb element containing rs7850258 allele G is significantly more responsive to both MYC and ARNT than allele A. By demonstrating that this common non-coding variant alters *FOXE1* expression, we have identified at least in part the functional basis for the genetic risk of these seemingly disparate disorders.

Introduction

Isolated cleft lip with or without cleft palate (CL/P) and cleft palate (CP) are common structural birth defects with a strong genetic component that remains incompletely understood. While periconceptional and gestational environmental influences contribute (smoking, alcohol and nutrition), twin, family and population-based studies have clearly established a genetic basis for CL/P etiology. For instance, there is a 40-fold increased prevalence among first-degree relatives compared with within the general population (1). Moreover, genome-wide association studies (GWAS) have identified at least 18 loci in which certain alleles of specific single nucleotide polymorphisms (SNPs) are significantly associated with isolated CL/P (2–5). However, coding mutations have been identified in <10% of patients. This follows the trend of GWAS of other diseases, where a meta-analysis of over 150 GWAS studies revealed only 12% of associated SNPs were in a haplotype block containing coding exons (6). Mutations landing in non-coding DNA are presumed, in many or most cases, to disrupt cis-regulatory elements (CREs) (i.e. promoters and enhancers). However, our understanding of how sequence variation alters the function of regulatory elements remains in its infancy. Consequently, very few mutations that directly elevate risk for CL/P have been identified, hindering progress in the investigation of disease mechanisms (7–8).

We have recently identified a strong peak of association for both CL/P and CP at 9q22, encompassing the gene *FOXE1* (9). This association has been replicated in a variety of Caucasian populations (10,11) and Hispanics from Honduras (12). Importantly, *FOXE1* is likely to be the relevant gene, because homozygous *FOXE1* mutations cause Bamforth–Lazarus syndrome which is characterized by cleft palate, choanal atresia, bifid epiglottis, thyroid agenesis or dysgenesis, hypothyroidism and spikey hair (13–15). Inactivation of *FOXE1* in mice causes a similar phenotype (16). We have found *FOXE1* coding mutations in only 1% of subjects, despite exhaustive mutation screens (9). This is incongruent with estimates that *FOXE1* has a 25–38% attributable risk, suggesting that causal variants occur in nearby cis-regulatory elements.

Consistent with thyroid defects in patients with Bamforth–Lazarus syndrome, a variety of common thyroid diseases also map to *FOXE1*, but causal variants for these disorders at this locus remain largely unknown. Thyroid disorders mapping to 9q22 include congenital hypothyroidism due to thyroid dysgenesis (17–19), hypothyroidism (20,21), goiters, non-medullary thyroid cancer (22,23), papillary thyroid carcinoma (24–27), radiation induced papillary thyroid carcinoma as a result of the Chernobyl accident (28,29), and thyroid cancer (30–32). Also, a number of biomarkers of thyroid metabolism are also associated at this locus (27,30,33–37). These observations are consistent with evidence that *FOXE1* is involved in thyroid homeostasis, negatively regulating thyroglobulin and thyroperoxidase expression (38). Interestingly, a SNP associated with papillary thyroid cancer, rs1867277, is located in the *FOXE1* promoter and alters binding of the *USF1/USF2* transcription factors (26). However, rs1867277 is unlikely to explain all of the risk for thyroid cancer at this locus, because other SNPs show more significant association, and because the haplotype block structure in the region suggests more than one risk allele is present for these diseases. Although *FOXE1* coding mutations may be present in a subset of patients with thyroid conditions mentioned earlier, it is probable that additional causative variants in non-coding DNA remain to be identified.

We hypothesized that causal mutations for orofacial clefting and thyroid diseases occur within cis-regulatory elements at the human *FOXE1* locus. To that end, we screened evolutionarily conserved sequences of human DNA at the *FOXE1* locus (Fig. 1) for craniofacial and thyroid enhancers using both transgenic zebrafish and mouse strategies that have been shown to effectively identify cis-regulatory elements (39,40). We identified three such elements, including one containing a SNP that is associated with CL/P, hyperthyroidism and thyroid cancer. *In vitro* quantitative reporter assays in embryonic oral epithelial and thyroid cells demonstrated that different alleles of this SNP altered the level of enhancer activity of the element.

Results

Endogenous *foxe1* expression during zebrafish development

We reasoned that oral and thyroid enhancers near human *FOXE1* would be more likely to be functional in zebrafish if the fish ortholog, *foxe1*, were expressed in oral tissues and thyroid gland. Indeed, RNA *in situ* hybridization revealed *foxe1* expression in the central nervous system at 11.4 h post fertilization (hpf) (Fig. 2A), in oral epithelium at 36 hpf (Fig. 2B, C, E and F, data not shown) and thyroid gland (Fig. 2D and G) at 36 hpf and later stages. We detected *foxe1* expression in pharyngeal arch epithelium at 72 and 96 hpf (Fig. 2H and I). Furthermore, we observed *foxe1* expression in the developing heart at 48 hpf, the pectoral fin starting at 72 hpf and the gill rakes (data not shown). These observations confirm and extend a previous expression analysis (41).

Enhancer screen results

We amplified from human DNA, 15 human conserved non-coding elements (hsCNEs) in a 152 kb region spanning from 101 kb upstream and 51 kb downstream of *FOXE1* and tested them for enhancer activity in transient transgenic zebrafish (Fig. 1; Supplementary Table 1). The hsCNEs are named by their position relative to the *FOXE1* start codon, negative being upstream and positive downstream. Our zebrafish enhancer screen was based on an eGFP reporter, pT2cfosGW eGFP (39). This reporter construct is designed to clone the human test DNA upstream of the mouse minimal *cfos* promoter that is unable to independently initiate mRNA transcription. To confirm this, we injected >200 single cell embryos with the reporter lacking any human sequences and did not detect any GFP expression. In contrast, of 15 hsCNEs, three had enhancer activities capable of directing transcription in a pattern consistent with endogenous *foxe1* expression (hsCNE+22.6, hsCNE-67.7 and hsCNE-82.4; Supplementary Material, Table S1), as summarized below. These positive hsCNEs acted independently and without the *FOXE1* promoter.

FOXE1 hsCNE+22.6 is active in the developing pharyngeal arches and heart

In embryos injected with *FOXE1* hsCNE+22.6 kb element (hg19 chr9: 100 638 752–100 639 947), we detected a reproducible pattern of eGFP expression in a mosaic pattern within pharyngeal muscles, a subset of cranial muscles (see Fig. 3 legend), heart and pharyngeal arch epithelium ($n > 800$ embryos scored) (Fig. 3). We established five independent, stable transgenic lines and all exhibited non-mosaic GFP expression in the same structures that

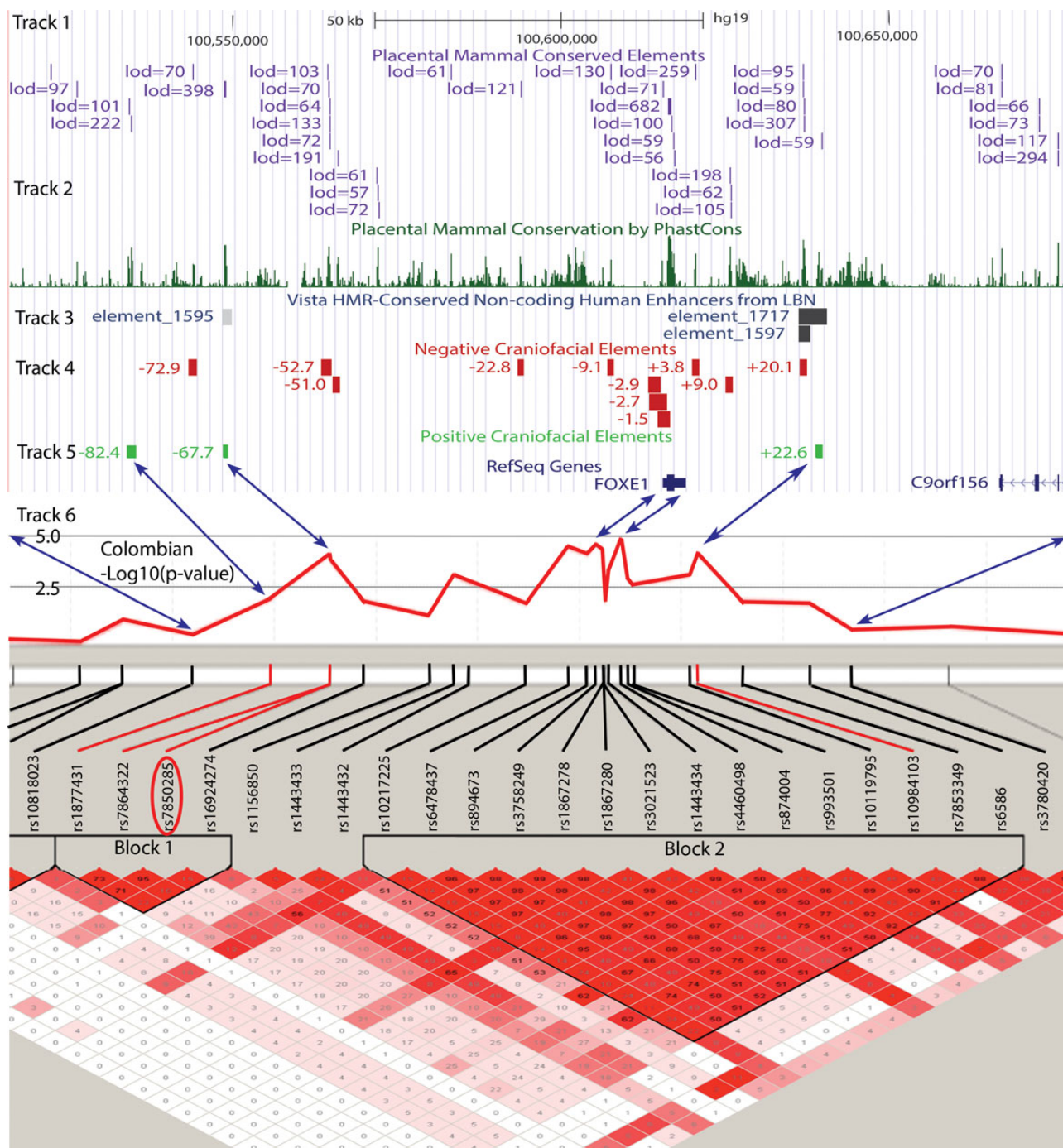


Figure 1. Genomic overview of the FOXE1 locus. Shown is a view from the UCSC Genome Browser of a 152 kb region bracketed by the SNPs rs10818023 and rs3780420. Track 1 shows genomic regions with a mammalian PhastCons score of >400. Track 2 is a histogram plot of mammalian conservation. Track 3 is from the Vista Enhancer indicating tissue specific enhancers (<http://enhancer.lbl.gov/>). Tracks 4 and 5 show the location of the tested conserved elements relative to the FOXE1 start codon and depicted by green and red boxes, representing negative and positive enhancer activity, respectively. Track 6 shows association with CL/P in the Colombian population. The linkage disequilibrium (LD) plot shows the D' correlation between any given SNP pair, red indicating a maximum D' of 1, pink shades ranging through values $D' < 1$ with white indicating no correlation. Blue arrows show the correspondence between the physical genomic and recombinant genetic distances. Circled in red is SNP rs7850285 that in addition to being the most significantly associated SNP with CL/P in block 1 is also the most significantly associated SNP in genome-wide association studies of hypothyroidism or thyroid cancer.

expressed GFP mosaically in transient transgenic G_0 embryos. In stable transgenic embryos, GFP was evident in the developing mouth by 24 hpf (data not shown) and by 72 hpf the muscles of the second and branchial arch were labeled along with the branchial arch cartilages including the gill rakes (Fig 3A–C). Moreover,

this element also directed GFP expression in the developing heart and pectoral fins (Fig. 3A–C). This pattern was confirmed by *in situ* hybridization with an eGFP probe. While aspects of this pattern matched that of *foxe1* expression, for instance in epithelium of ceratobranchial arches (e.g. compare Fig. 3D to Fig. 1H), however

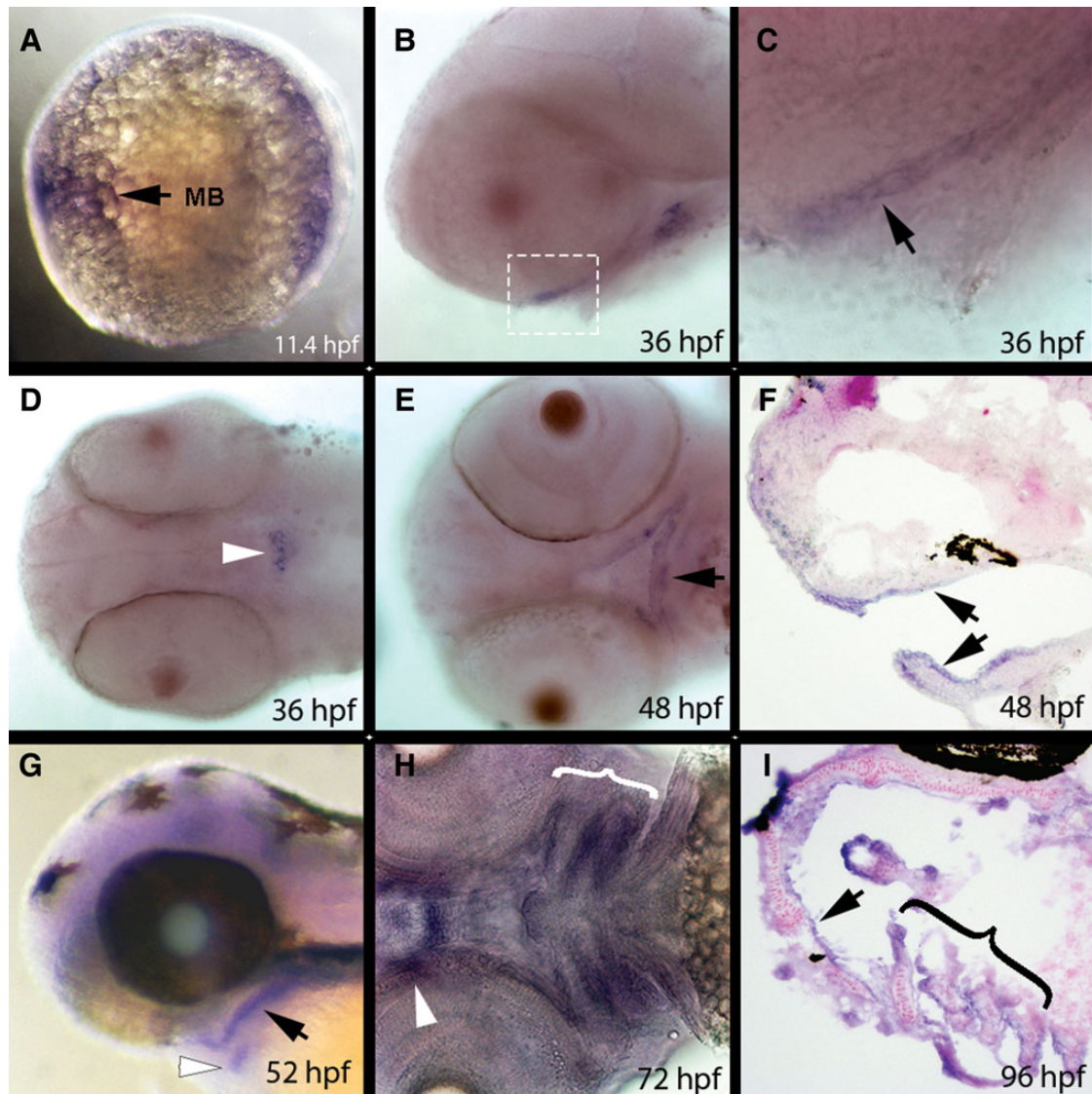


Figure 2. Endogenous *foxe1* expression during zebrafish development. *In situ* hybridization of *foxe1* on embryos fixed at the indicated stages. All embryos are presented with anterior to the left. (A) Lateral view of an 11.4 hpf embryo. Expression is evident in the central nervous system at approximately the level of the midbrain (mb). (B and C) Lateral views of a 36 hpf embryo. Expression is evident in oral epithelium (arrow in C). White box in B indicated region shown in C. (D) Ventral view of a 36 hpf embryo. Expression is visible in the developing thyroid (arrowhead). (E) Ventral view of a 48 hpf embryo. Expression is evident in oral epithelium (arrow). (F) Sagittal section of a 48 hpf embryo. Expression is seen in oral epithelium (arrowheads). (G) Lateral view of a 52 hpf embryo. Expression is seen in oral epithelium (black arrow), and in thyroid (white arrowhead). (H) Ventral view of a 72 hpf embryo. Expression is seen in epithelium of the pharyngeal arches (bracket) and in oral epithelium around mouth (arrowhead). (I) Transverse section of a 96 hpf embryo. Expression is seen in the oral epithelium (black arrowhead) and pharyngeal arches (bracket). MB, midbrain. hpf, hours post fertilization.

expression of *foxe1* mRNA was not detected in pharyngeal muscles. If human *FOXE1* is also absent from pharyngeal muscles, we infer that additional regulatory elements must normally prevent the enhancer at +22.6 kb from driving *FOXE1* expression in this tissue. Alternatively, through divergence in tissue specific expression of upstream regulatory factors, this enhancer may be active in pharyngeal muscles only in fish, although ENCODE data show this region has chromatin marks of enhancer activity in human skeletal muscle myoblasts.

While *hsCNE+22.6* was able to activate *lacZ* expression in human embryonic oral epithelial cell lines, no difference was observed between the two rs10984103 alleles (Supplementary Material, Fig. S1).

FOXE1 hscNE-67.7 drives expression in the developing oral epithelium, hyoid arch, branchial arches, thyroid and heart

In transient transgenic embryos injected with the *FOXE1* -67.7 element (hg19 chr9: 100 548 460–100 549 280), we reproducibly detected GFP expression in the jaw, hyoid and pharyngeal arches, thyroid and heart (data not shown). The same pattern was observed in five independent transgenic lines and in these, expression in the oral epithelium was more evident than in transient transgenics (Fig. 4A and B). Expression in oral epithelium and thyroid precursors was detectable by 24 hpf, while heart expression became evident at 48 hpf (data not shown). Staining with the thyroid specific T4 antibody demonstrated coincident signal with

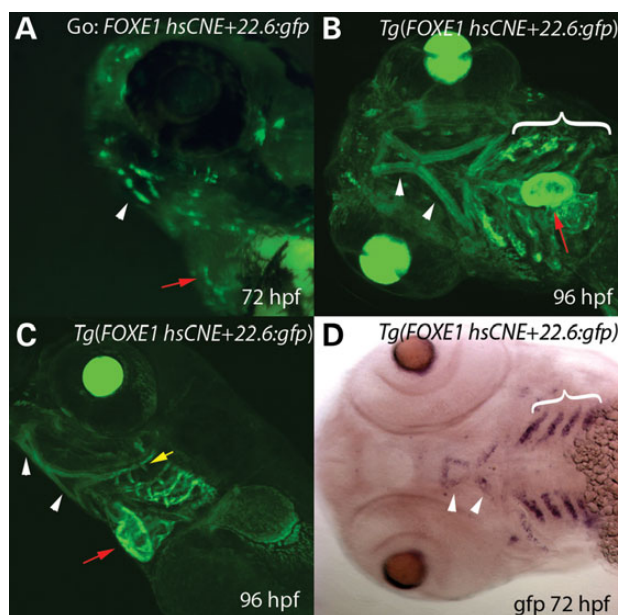


Figure 3. FOXE1 hsCNE+22.6 kb element drives GFP expression in pharyngeal muscles, the heart and pharyngeal epithelium. (A–C) Compressed Z-stacks from confocal imaging of live (A) transient (G_0), and (B,C) stable (F1) transgenic $Tg(foxe1+22.6:gfp)$ animals. (A) Oblique lateral view of a G_0 72 hpf embryo mosaic for $Tg(foxe1+22.6:gfp)$. GFP is visible in cells in the region of the developing mouth (white arrow) and in the heart (red arrow). (B) Ventral and (C) lateral views of a 96 hpf larvae showing eGFP expression in the pharyngeal arches (bracket), heart (red arrow) and the ventral pharyngeal muscles (arrowheads) including the hyohyoideus muscle, interhyoideus, intermandibularis anterior and intermandibularis posterior muscles. Also labeled are the gill rakes (yellow arrow). (D) Ventral view of 72 hpf $Tg(foxe1+22.6:gfp)$ embryo processed to reveal *gfp* expression by RNA in situ hybridization, which is evident in the pharyngeal arches (bracket) and ventral pharyngeal muscles (arrowheads). The heart, which was positive for *gfp* expression, was removed in the specimen shown here.

eGFP within the thyroid in the hsCNE-67.7 transgenic fish (data not shown).

We next tested an overlapping region, (hg19 chr9: 100 548 557–100 549 640) in transient transgenic mouse embryos using a β -galactosidase reporter with an Hsp68 minimal promoter. This region, Vista Enhancer *hs1595*, drove reporter expression in a pattern matching endogenous *foxe1* expression in the oral epithelium and developing thyroid in three of four mouse embryos (Fig. 4C–F, Supplementary Material, Fig. S2). In summary, the FOXE1 hsCNE-67.7 element has enhancer activity in multiple oral tissues, in the heart and the thyroid gland during development in zebrafish, and an overlapping human DNA element has similar activity in mouse embryos.

FOXE1 hsCNE-82.4 directs expression in the oral epithelium in zebrafish embryos

For the two hsCNEs just described, the expression patterns in stable transgenic lines could be accurately predicted from summing the expression patterns in G_0 embryos. Therefore, we screened additional elements solely in G_0 embryos. G_0 embryos injected with the FOXE1 hsCNE-82.4 element (hg19 chr9: 100 533 759–100 535 187) showed eGFP expression initially in the presumptive oral epithelium at 24 hpf until at least 96 hpf, with additional signal in the pharyngeal region (Fig. 5) ($n > 370$ embryos injected). We observed such expression in 7–19% of embryos in seven different injection batches, a rate comparable to those observed for the elements above (Table 1).

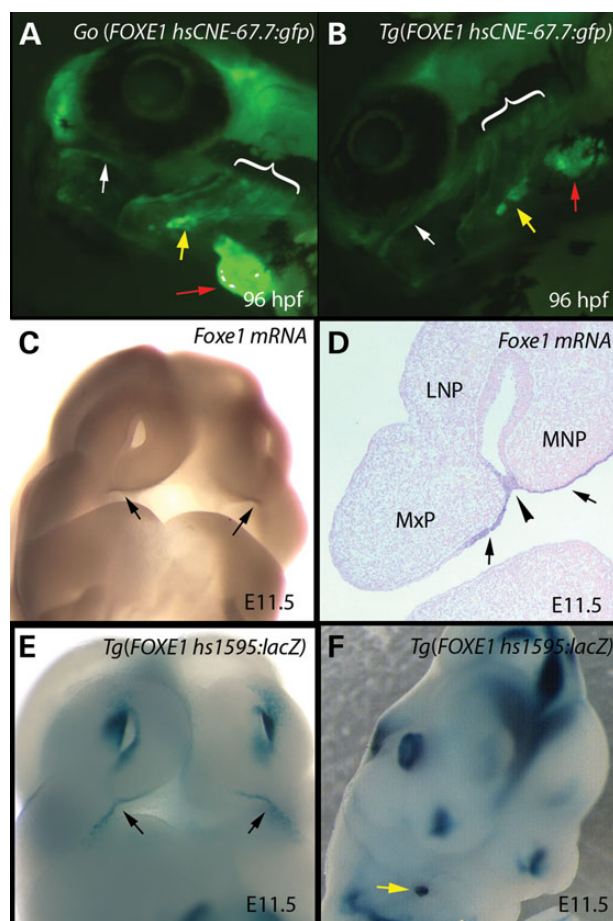


Figure 4. FOXE1 hsCNE-67.7 kb element drives GFP expression in the oral epithelium, pharyngeal arches, thyroid and heart tissues. (A and B) Compressed Z-stacks from confocal imaging of live stable transgenic $Tg(foxe1-67.7:gfp)$ animals from two independent lines. (A and B) Oblique lateral views of 96 hpf larvae showing GFP expression in the oral epithelium (white arrows); pharyngeal arches (brackets), thyroid (yellow arrows) and heart (red arrows). (C) Oblique frontal view of E11.5 mouse embryo showing endogenous *Foxe1* expression in the oral epithelium at the juncture of the medial nasal and maxillary processes (arrows). (D) Frontal section of E11.5 mouse embryo showing endogenous *Foxe1* expression in the oral epithelium of the medial nasal and maxillary processes (arrows), including the point of fusion (arrowhead) between them that results in the formation of the upper lip. (E and F) Mouse embryos transiently transgenic $Tg(hs1595:lacZ)$ stained for β -galactosidase. *hs1595* corresponds to hg19 chr9: 100 548 557–100 549 640 overlapping the $Tg(foxe1-67.7:gfp)$ tested in zebrafish. (E) Oblique frontal view of E11.5 mouse embryo showing β -galactosidase staining in the oral epithelium at the juncture of the medial nasal and maxillary processes (arrows). (F) Oblique view of E11.5 mouse embryo showing β -galactosidase staining in the developing thyroid (yellow arrow).

Differential enhancer activity for rs7850258 alleles in FOXE1 hsCNE-67.7

Interestingly, rs7850258, for which one allele (G) is associated with hypothyroidism and CLP, and another (A), with thyroid cancer, is present within the FOXE1 hsCNE-67 element. We engineered variants of this element, containing one or the other alleles of SNP rs7850258, into a vector containing a minimal promoter and the gene encoding firefly luciferase. Since *Foxe1* is expressed in the oral epithelium during facial development (9), an embryonic oral epithelial cell line that endogenously expresses FOXE1, GMSM-K (human fetal oral epithelial cells), was used in these assays. For assessment in the context of thyroid function, the rat FRTL epithelial



Figure 5. FOXE1 hsCNE-82.4 directs GFP expression in the oral epithelium. (A and B) Lateral view line drawings representing a composite of transient G_0 embryos at 48 and 72 hpf, showing eGFP labeled cells highlighted in pink. GFP expression is primarily in the oral epithelium. (C) Lateral view of a representative G_0 larvae at 96 hpf showing expression in the oral epithelium (white arrowheads). In all over 370 embryos were screened from seven different injection batches.

Table 1. Scoring of expression in G_0 embryos

FOXE1 hsCNE	Total live embryos	Expressers (%)	Expression in oral or pharyngeal regions ^a				Number transgenic lines
			24 hpf (%)	48 hpf (%)	72 hpf (%)	96 hpf (%)	
+22.6	822	41	29	18	13	11	5
-67.7	262	30	ND	ND	ND	ND	5
-82.4	453	20	19	17	13	7	0

^aPercent of live embryos.

thyroid cell line that expresses *Foxe1* under hormonal control was used (42). In both cell types, the enhancer activity of the hsCNE-67 was confirmed with significant increases (1.5–5.5-fold) above the empty vector (Fig. 6 A and B). Moreover, in both cell types, the rs7850258 G allele had statistically significant greater enhancer activity, ranging from 1.2- to 1.7-fold more than the rs7850258 A allele (Fig. 6A and B). rs7864322 is another SNP within the hsCNE-67.7 enhancer that is also associated with CL/P (9), raising the possibility that it has functional effects. Yet testing all allelic combinations with rs7850258 in both GSM-K and FRTL cells revealed no difference between the two rs7864322 alleles (C/T) whether they were on either the rs7850258 A or G alleles (Fig. 6A and B). These results suggest that rs7850258 is the functional variant for both oral epithelial and thyroid diseases. Transient transgenic zebrafish assays, scored by two individuals blinded to the rs7850258 genotype, did not reveal any statistically significant differences in GFP expression patterns, supporting conclusion that variation at rs7850258 alters the level of expression rather than the tissue specificity of expression (data not shown).

Using JASPAR (43), it was predicted that there was a gain of E-box transcription factor binding site to the rs7850258 G allele that could be bound by the basic-helix-loop-helix transcription factors MYC and ARNT (Supplementary Material, Table S2). Co-transfection of either human MYC or ARNT along with the reporter construct of hsCNE-67.7 showed differential allelic activation with the G allele being more strongly activated by both transcription factors in both human embryonic oral epithelial and rat thyroid cell lines (Fig. 6C and D).

Discussion

Zebrafish and mouse orthologs of FOXE1 are expressed in the oral epithelium and in the thyroid gland consistent with role for FOXE1 in diseases of both tissues

We have confirmed in zebrafish that *foxe1* is expressed in the developing oral and pharyngeal epithelium, thyroid gland and

pectoral fins. Furthermore, *foxe1* is expressed in the heart, and in the developing midbrain to hindbrain. While *foxe1* expression in central nervous system has not been explicitly mentioned previously, it is evident in an earlier study in zebrafish [see Fig. 1 of (41)]. Similarly, in mice *Foxe1* expression in the developing central nervous system and heart has not been previously reported, but a review of the literature shows expression in the midbrain and hindbrain on embryonic Day 10.5 (44). Furthermore, in the public domain via EuroExpress, *Foxe1* mRNA expression is evident in the midbrain, cerebellum and heart outflow tracts in E14.5 mouse embryos (Supplementary Material, Fig. S3) (45). This is consistent with RT-PCR data showing in zebrafish adults very strong expression in the heart and brain (41).

Identification of human tissue specific enhancers at the FOXE1 locus

Using an strategy based primarily on sequence conservation, we tested 15 regions of human DNA totaling 14.9 kb, encompassing 9.8% of the human FOXE1 locus as defined by the CL/P critical region (9). Of these, three showed positive tissue specific enhancer activity in patterns consistent with endogenous *foxe1* expression. All three enhancers, hsCNE+22.6, hsCNE-67.7 and hsCNE-82.4 shared the same capacity to independently regulate expression in the oral and pharyngeal epithelium, and pharyngeal arch structures. Two of these, hsCNE+22.6 and hsCNE-67.7 are also heart and pectoral fin enhancers. Unique is the ability of hsCNE-67.7 to direct expression in the developing thyroid. The oral epithelial and thyroid activity of the -67.7 element was also confirmed in transient transgenic mice. In contrast, the enhancer activity of hsCNE-82.4 is primarily confined to the oral epithelium.

In several previous studies, human DNA, which is not detectably conserved to zebrafish, has been shown to nonetheless possess enhancer activity in zebrafish. Examples include enhancers for RET, a gene for Hirschsprung disease (46), for PHOX2B, which is associated with neuroblastoma and Crohn's disease (47–49), and

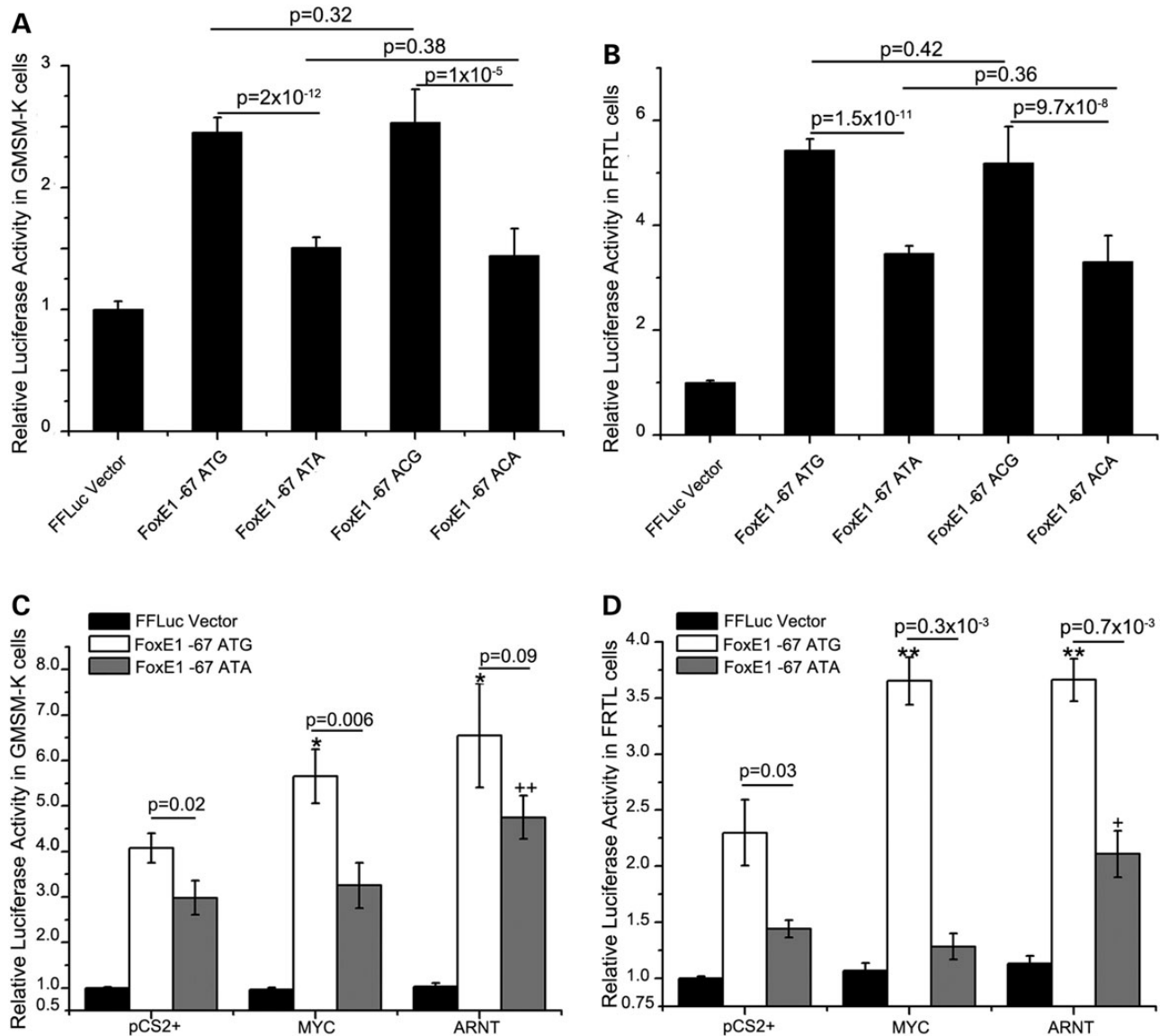


Figure 6. rs7850258 alleles differentially affect FOXE1 hsCNE-67 enhancer activity in human embryonic oral epithelial and rat thyroid cell lines. Different alleles of rs7864322 (C/T) and rs7850258 (A/G) in the FOXE1 hsCNE-67.7 element were cloned pTol2-cFos-FLuc (firefly luciferase vector with a c-fos minimal promoter). The same genotype for the other SNP in hsCNE-67, rs1006125 (A), was maintained in all constructs. The clones are named ATG, ATA, ACG and ACA designating the sequential genomic order of rs1006125, rs7864322 and rs7850258 within the constructs. The ATA and ATG constructs along with Renilla luciferase (pTol2-cFos-RLuc) were separately transfected into (A and C) GMSM-K human oral epithelia cells and (B and D) FRTL rat thyroid cells. Luciferase measurements were normalized across transfections using Renilla luciferase readings and then the firefly luciferase readings were normalized to the empty Firefly luciferase vector. (A and B) In each cell line, the rs7850258 G allele had increased enhancer activity compared with the A allele. (C and D) rs7850258 alleles differentially respond to MYC and ARNT. Since there was no difference between the ATG versus ACG or ATA versus ATA, only the ATG and ATA clones described earlier were tested for differential response to cotransfected human MYC and ARNT. In both cell lines, the rs7850258 G allele had increased enhancer activity compared with the A allele in the presence of either MYC or ARNT. */+ P-value <0.05 and **/+ P-value <0.01. The data in each column represent the mean \pm SEM of three measurements each of four biological replicates.

one for NOS1AP that contains a functional SNP associated with cardiac QT interval (50). Similarly, the FOXE1 enhancers that we have identified here lack detectable conservation to zebrafish, the longest stretch of 100% identity being 26 bp and for any identity >85% being 47 bp of which none map to the zebrafish *foxe1* locus (Supplementary Material, Table S1). While the interpretation of CNE-82.4 as a facial enhancer is based entirely on transient transgenic zebrafish embryos, our experience and published work has shown expression patterns in transient transgenic zebrafish embryos accurately reflect that found in founder lines (51,52). Nonetheless, there is no guarantee that every element

conserved in mammals will have enhancer activity in zebrafish, and therefore negative results, as we observed for 12 of 15 tested elements, cannot be strongly interpreted. Alternative explanations for the negative results could be those elements act as silencers or insulators, which would not have been detected using the reporter construct in this study. Additional craniofacial enhancers for FOXE1 are likely to exist; criteria other than conservation, for instance chromatin marks in appropriate cell lines, may be of use to identify them.

In retrospect to see if we could improve upon our 20% success rate when choosing human candidate elements based on

conservation, we compared our results to available ENCODE data. Only the hscCNE+22.6 enhancer had chromatin marks and an abundance of ChIP-Seq peaks, clearly indicating regulatory activity. Beyond that, we could not identify a matching pattern for either positive or negative elements with chromatin marks for regulatory function in any cell type. This underscores the need to have such data available on the relevant cell type. This is especially true for studying embryonic structural birth defects for which there is a paucity of data resources.

Relevance to human diseases

The expression of *FOXE1* in the developing heart suggests it may be associated with cardiac defects. To date, variations at the *FOXE1* locus have not been associated with cardiac anomalies. Yet tangential data exist in that among infants with congenital hypothyroidism there is a 3–5-fold over representation of heart defects (53,54), suggesting they may have shared developmental processes.

At the *FOXE1* locus, based on association peaks and LD blocks in different populations, there is evidence for three CL/P risk haplotypes relative to the start codon: –89 to –59 kb; –19.2 to +0.4 kb; and +7.0 to +81 kb (9). We postulate that rs7850258 within hscCNE-67.7 explains entirely or in part the risk for CL/P associated with the most upstream risk haplotype. The facial enhancer, hscCNE+22.6, containing rs10984103, the SNP most strongly associated with CL/P in the Filipino population, likely has a role in the most downstream risk haplotype. However, there was no statistical difference between the two rs10984103 alleles.

The discovery of the hscCNE-67.7 thyroid enhancer properties is noteworthy given the number of thyroid diseases ascribed to the *FOXE1* locus. It contains rs7850258, that has the most significant association in GWAS studies of hypothyroidism [$P = 2.5 \times 10^{-19}$ (20); $P = 3.96 \times 10^{-9}$ (21)] and papillary thyroid cancer ($P = 1.7 \times 10^{-9}$) (21,29). We show here that rs7850258 alters enhancer function in both oral epithelial and thyroid cell lines. Of interest is that the G allele is associated with increased risk for CLP and hypothyroidism, while the A allele is associated with thyroid cancer. Relative to the pathogenesis of CLP, that the G allele has more enhancer activity is consistent the gain of function mutation characterized in one patient with Bamforth–Lazarus syndrome (55). Additionally, over expression of *Foxe1* in mice also leads to cleft palate (56). One possible mechanism proposed for the association with hypothyroidism is that ectopic *FOXE1* induces apoptosis in the developing thyroid, leading to agenesis or dysgenesis (55).

The decreased enhancer activity of the A (cancer associated) allele of rs7850258 also implies that *FOXE1* is a tumor suppressor. Evidence for this exists in that the *RET/PTC3* proto-oncogene caused by chromosomal rearrangements leading to thyroid papillary carcinoma is known to decrease *FOXE1* expression (57). Furthermore, somatic loss of the 9q22 region containing *FOXE1* is frequent in squamous cell carcinoma (58). Moreover, hypermethylation of the *FOXE1* promoter and/or decreased *FOXE1* expression have been observed in a number of cancers including squamous cell carcinoma, (59), pancreatic cancer (60), adenoid cystic carcinoma of salivary gland (61), breast cancer (62), colorectal cancer (63) and anaplastic carcinomas (64). Additionally, another risk allele at rs965513 for papillary thyroid carcinoma is associated with decreased *FOXE1* expression in thyroid tissue (65). In total, these reports are also consistent with *FOXE1* being pro-differentiation in the developing thyroid and differentiation maintenance in the adult thyroid (44,65).

rs7850258 is predicted to disrupt ARNT and MYC transcription factor binding and consistent with this our data shows a significant response to either factor when the binding site is present in contrast to no response when absent. rs7850258 affects an E-box binding site, so it is possible that other E-box binding transcription factors may be involved as well. That being said there is evidence for both ARNT and MYC being involved in orofacial clefting and thyroid diseases.

ARNT and ARNT2 are basic helix-loop-helix transcription factors that are expressed in the palatal shelf epithelium (66–68). Furthermore, ARNT2 is expressed in the developing thyroid (66). ARNT and ARNT2 regulate gene expression through dimerization with the aryl hydrocarbon nuclear receptor (AHR). AHR responds to 2,3,7,8-tetrachlorodibenzo-p-dioxin (TCDD), a well-known teratogen that causes cleft palate in mice (69) and is associated with clefting in humans through agricultural chemical exposures (70). TCDD is also found in tobacco smoke that through maternal gestational exposure attributes to 11% of orofacial clefts (71). Moreover, a genetic variation at the ARNT locus is associated with CL/P in humans (72). While there is convincing data in animal models demonstrating TCDD exposure interferes with fetal and newborn thyroid function, epidemiological studies with equivocal results have not shown a consistent correlation in humans (73). There have not been any reports of association between ARNT, ARNT2 or AHR with hypothyroidism or thyroid cancer in any human studies.

c-MYC is also a b-HLH protein that is expressed in both the epithelia and mesenchyme of the facial processes (74,75). Conditional inactivation in cranial neural crest cells results in craniofacial anomalies (76) and deletion of a facial enhancer at the *Myc* locus results in cleft lip in conjunction with decreased facial *Myc* expression. Increased MYC expression is associated with undifferentiated thyroid cancer and the loss of thyroid differentiation markers, PAX8 and TTF1 (77). So in theory, the rs7850258 cancer associated allele A would be less responsive to MYC, resulting in relative decrease of *FOXE1* expression compared with the non-risk cancer allele G. This model is consistent with *FOXE1* being a tumor suppressor.

A final caveat in this discussion is that until there is association data based on comprehensive sequencing across the region in individuals with CLP and thyroid diseases, it is not possible to state whether rs7850258 is the main contributing factor or if there are additional functional variants on a shared haplotype that in total comprise the genetic risk for these disorders at this locus.

In summary, using zebrafish and mouse transgenesis, we have identified craniofacial and thyroid enhancers at the *FOXE1*, including one that contains functional risk alleles for both CLP and thyroid diseases. These results reveal the pathogenesis of these disorders associated with the *FOXE1* locus. We anticipate that there are additional genetic and biological mechanisms at this locus yet to be discovered that contribute to risk for these disorders. Given that thyroid diseases and orofacial clefting converge on the same SNP within a thyroid and facial enhancer, there may a shared risk for both diseases within families or a population.

Materials and Methods

Zebrafish husbandry

Parental fish were housed at room temperature overnight prior to breeding. Embryos were raised at 28.5°C. The ethical use of animals for research was approved by the University of Iowa The Institutional Animal Care and Use Committee.

In situ hybridization

A 870 bp *Foxe1* probe was cloned using primers [forward—ATGCCTGTGGTTAAAGTGGAGAGT (chr1: 25 668 976–25 668 999; Zv9/danRer7) and reverse—GCACTCAGCATTATGGGCCA (chr1: 25 668 130–25 668 149; Zv9/danRer7)] to amplify from a zebrafish cDNA library. The probe was sequenced and by BlastN search of the Zebrafish genome, 100% identity over the entire 870 bp probe was observed for only *foxe1* (ref|XM_690973.3|) at chr1: 25 668 130–25 668 999 in Zv9/danRer7. The next level of BlastN identity was 79% with *Foxe3* for only 328 base pairs. Based on these results and those published by Nakada *et al.* (41), there is only one zebrafish *foxe1* gene and that this gene has been given the gene ID number ENSDARG0000079266 in Ensembl. *In situ* hybridization was performed as previously described (78). Given the stringency conditions used, it is highly unlikely that hybridization of the full length probe would occur with anything other than the endogenous *foxe1* mRNA.

Selection of potential regulatory elements

To be conservative relative to boundaries established by our CL/P association data, we have targeted a 152 kb interval between hg19 Chr9: 100 514 919–100 666 931 (rs10818023 to rs3780420) (Fig. 1). We chose 15 conserved non-coding human sequences (hsCNE) using publicly available sequence data (UCSC) and the Phastcons algorithm (79) with a mammalian Phastcons score of >400 as inclusion criteria (Supplementary Material, Table S1). Fourteen of our chosen elements met these criteria. One additional element, hsCNE-2.9, was added since it contained a region overlapping with the rat *Foxe1* promoter (80). For all 15 regions there was no identity within the Zebrafish genome for more than 30 base pairs.

Zebrafish enhancer screen

Potential regulatory elements were PCR amplified from human BAC clones RP11-10012 or RP11-151120, cloned into the Gateway® (Invitrogen) pENTR/D-TOPO vector and transferred to the pT2cfosGW eGFP reporter construct that is constructed on the Gateway® system (39). The pT2cfosGW eGFP reporter is designed to clone the human test DNA upstream of the mouse minimal cfos promoter that is unable to independently initiate mRNA transcription. Also, built into the pT2cfosGW vector are Tol2 recombination sites bracketing the entire construct that facilitate integration of the expression construct into the zebrafish genome thus reducing cell mosaicism within the injected embryos. All PCR products and plasmid constructs were sequenced and the sequence results were aligned onto the human genome using the UCSC tool BLAT (81) to ensure fidelity of these steps. Tol2 mRNA was transcribed from the plasmid pCS-TP (82).

Constructs were injected along with the tol2 mRNA into 100–200 zebrafish embryos at the 1 cell stage. The developing embryos (generation zero; G₀) were screened at 24, 48, 72 and 96 hpf for eGFP expression patterns consistent with endogenous *foxe1* expression. A consistent pattern of expression in a minimum of 10% of injected fish was the criteria for tissue specific enhancer activity (83). The use of recombinant DNA was approved by University of Iowa Institutional Biosafety Committee.

Mouse enhancer assay

The methods for evaluating conserved non-coding sequences in transgenic mice have been previously described as part of the Vista Enhancer project (84). Briefly, target regions were amplified

from human genomic DNA, cloned into an HSP68-LacZ reporter vector, and sequence validated. The linearized vector was injected into the pronucleus of FVB oocytes to generate transgenic embryos. LacZ activity was assessed at E11.5 for craniofacial staining. The results for Vista Enhancer hs1595 have not been previously reported in detail.

Luciferase reporter constructs, transfections and luciferase assays

To assess differential allelic activity for the rs10984103 CL/P risk SNPs within hsCNE+22.6, the alleles (A/C) two were introduced by PCR mediated mutagenesis (85) into the hsCNE+22.6 enhancer and subsequently subcloned into pTol2-cFos-FLuc (firefly luciferase vector with a c-fos minimal promoter) using LR reaction according to the manufacturer's protocol (Life Technologies). In a similar fashion different combinations of the alleles for rs7850258 (A/G) and rs7864322 (C/T) were introduced into hsCNE-67.7, keeping the same allele (A) for the other SNP, rs1006125 that also resides in this enhancer. Sequencing was used to ensure clone integrity. Transient transfections were performed using X-tremeGENE HP (Roche, Germany) into GMSM-K or FRTL cells with every firefly luciferase vector. For each construct, three independent transfections were performed with Renilla luciferase (pTol2-cFos-RLuc) co-transfection, as an internal control for transfection efficiency. The Dual-Luciferase Reporter Assay System (Promega, Madison, USA) and a luminometer were used to measure luciferase activity in cell lysates. All quantified results are presented as mean ± SEM. Three luciferase measurements were made on each of three or four independent biological replicates. A two-tailed unpaired Student's t-test was used to determine statistical significance.

Predictions of differential allelic transcription factor binding for rs7850258 were performed using JASPAR (43). Ten base pairs surrounding rs7850258 GCCAGa/gTGTCT was evaluated and relative score of >0.85 was used as a threshold for significance.

The human ARNT expression vector is from GeneCopoeia (Rockville, MD, USA) plasmid #EX-C0312-M68 and the human MYC expression vector is from Addgene (Cambridge, MA, USA) plasmid #18773: MSCV MYC T58A puro (86).

Cell culture

GMSM-K a human fetal oral epithelial cell line (a kind gift from Dr Daniel Grenier) (87) was maintained in keratinocyte serum-free medium (Life Technologies) supplemented with EGF 1–53 and bovine pituitary extract (Life Technologies) in a 5% CO₂ atmosphere at 37°C. GMSM-K cells constitutively express FOXE1. FRTL a rat thyroid cell line was purchased from ATCC (CRL-1468; 42) and cultured in Ham's F-12K (Kaighn's, Life Technologies) supplemented with 10 mU/ml TSH (Sigma, St Louis, USA), 0.01 mg/ml insulin (Sigma), 10 nM hydrocortisone (Sigma), 0.005 mg/ml transferrin (Sigma), 10 ng/ml somatostatin (Sigma), 10 ng/ml glycyl-L-histidyl-L-lysine acetate (Sigma), and 0.5% FBS. In order to induce the expression of *Foxe1*, the concentration of TSH was increased to 30 mU/ml.

Supplementary Material

Supplementary Material is available at HMG online.

Acknowledgements

We thank Dr Shannon Fisher for sharing the pT2cfosGW eGFP reporter construct. Also, Dr Koichi Kawakami of Division of

Molecular and Developmental Biology, National Institute of Genetics and Department of Genetics, Graduate University of Advanced Studies, Mishima, Shizuoka, Japan generously provided the Tol2 plasmid pCS-TP. Dr Daniel Grenier at the Université Laval, Québec graciously provided the GSM-K human fetal oral epithelial cell line. Youssef Kousa aided with Figure construction. We greatly appreciate Drs Rick Domann and Adam Dupuy for sharing the ARNT and MYC expression plasmids.

Conflict of Interest statement. None declared.

Funding

This publication was supported by the National Institutes of Health (RO1 DE014667), the University of Iowa Department of Orthodontics and the National Center for Advancing Translational Sciences through Grant 2 UL1 TR000442-06 (A.C.L.). RO1 DE021071 and the National Science Foundation IOS-1147221 (R. A.C.). U01DE020060 (A.V.). The content is solely the responsibility of the authors and does not necessarily represent the official views of the NIH.

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