Minor change, major difference: divergent functions of highly conserved cis-regulatory elements subsequent to whole genome duplication events

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SUMMARY
Within the vertebrate lineage, a high proportion of duplicate genes have been retained after whole genome duplication (WGD) events. It has been proposed that many of these duplicate genes became indispensable because the ancestral gene function was divided between them. In addition, novel functions may have evolved, owing to changes in cis-regulatory elements. Functional analysis of the PAX2/5/8 gene subfamily appears to support at least the first part of this hypothesis. The collective role of these genes has been widely retained, but sub-functions have been differentially partitioned between the genes in different vertebrates. Conserved non-coding elements (CNEs) represent an interesting and readily identifiable class of putative cis-regulatory elements that have been conserved from fish to mammals, an evolutionary distance of 450 million years. Within the PAX2/5/8 gene subfamily, PAX2 is associated with the highest number of CNEs. An additional WGD experienced in the teleost lineage led to two copies of pax2, each of which retained a large proportion of these CNEs. Using a reporter gene assay in zebrafish embryos, we have exploited this rich collection of regulatory elements in order to determine whether duplicate CNEs have evolved different functions. Remarkably, we find that even highly conserved sequences exhibit more functional differences than similarities. We also discover that short flanking sequences can have a profound impact on CNE function. Therefore, if CNEs are to be used as candidate enhancers for transgenic studies or for multi-species comparative analyses, it is paramount that the CNEs are accurately delineated.

KEY WORDS: Cis-regulatory elements, DDC model, Enhancer analyses, Evolution, Transcriptional regulation, Whole genome duplication events

INTRODUCTION
It is widely accepted that during vertebrate evolution two whole genome duplication (WGD) events occurred, followed by another in the lineage leading to teleosts (Amores et al., 1998; Dehal and Boore, 2005; Holland et al., 1994; Ohno et al., 1968; Taylor et al., 2003; Wittbrodt and Scharl, 1998). These events coincided with a rapid expansion in organisational complexity particularly in teleosts, a lineage that constitutes half of extant vertebrates. The persistence of many duplicate genes after these WGD events forms the basis of the duplication-degeneration-complementation (DDC) model of gene evolution (Force et al., 1999). The DDC model predicts that at least some aspects of ancestral gene function are sub-partitioned between duplicate genes in a complementary manner, such that each copy remains indispensable. At the same time, redundancy of associated cis-regulatory elements may increase the ‘evolvability’ of these sequences (Jimenez-Delgado et al., 2009) and their potential to direct novel gene sub-functions. These ideas have led to the hypothesis that, rather than changes to protein-coding sequences, divergence in transcriptional regulation is the main driving force behind innovations in the vertebrate body plan (Aburomia et al., 2003; Levine and Tjian, 2003). Here, we explore these concepts using the PAX2/5/8 gene subfamily as a model and, in the process, discover key considerations for CNE and enhancer studies.

The highly related vertebrate PAX2, PAX5 and PAX8 genes derive from the two pan-vertebrate WGD events while the more recent teleost specific WGD resulted in two co-orthologous pax2 genes: pax2a and pax2b in zebrafish, and pax2.1 and pax2.2 in other teleosts (Pfeffer et al., 1998; Wada et al., 1998). The current data suggest that these genes have evolved in a manner consistent with the DDC model. In all vertebrates examined so far, PAX2, PAX5 and PAX8 collectively have important functions in the development of the CNS, eye, ear, kidney and thyroid, but the roles of individual genes have diverged both within the subfamily and across species (reviewed by Goode and Elgar, 2009). Bouchard and colleagues provided tangible evidence that cis-regulatory elements may be responsible for at least some of the functional divergence of these genes. They demonstrated that the insertion of Pax5 cDNA into the Pax2 locus is able to rescue Pax2-mutant phenotypes in mouse, even in domains where Pax5 is not normally expressed (Bouchard et al., 2000). Therefore, given the correct regulatory environment, mouse Pax5 at least is capable of substituting for Pax2.

In this paper, we investigate an interesting class of putative cis-regulatory elements consisting of non-coding sequences that are highly conserved between Fugu and humans [an evolutionary distance of around 450 million years (Sandelin et al., 2004; Woolfe et al., 2005)]. These conserved non-coding elements (CNEs) cluster around genes that are involved in transcriptional and developmental regulation and many exhibit in vivo enhancer activity in model...
organisms, including zebrafish (de la Calle-Mustienes et al., 2005; Shin et al., 2005; Woolfe et al., 2005), mouse (Pennacchio et al., 2006), frog (de la Calle-Mustienes et al., 2005) and chick (Sabherwal et al., 2007). Additional evidence that at least some of these CNEs regulate gene expression has been provided by the identification of point mutations (Lettice et al., 2003; Benko et al., 2009), deletions (D’Haene et al., 2009; Sabherwal et al., 2007) and translocations (Gill et al., 2009) of individual CNE sequences that produce mutant phenotypes owing to dysregulation of the associated protein-coding region.

Previous analyses (Woolfe and Elgar, 2007) (http://condor.nimr.mrc.ac.uk/) have shown that the vertebrate PAX2 gene is associated with a large number of CNEs (around 60). Interestingly, many tetrapod PAX2 CNEs have sequence homology to both teleost pax2 loci, suggesting that a large proportion of CNEs have been retained in duplicate subsequent to the WGD event that occurred in the teleost lineage. Here, we have exploited the wealth of these CNE duplicates and analysed their sequences in relation to the single tetrapod CNE copies. Coupling this with intra-species comparative functional analyses has enabled us to assess their function with regard to the DDC model. Strikingly, our results show that most duplicate CNEs have differences in their enhancer activities and that even highly similar sequences can direct very different patterns of reporter gene expression.

MATERIALS AND METHODS

Bioinformatic analyses

CNEs associated with PAX2/5/8 gene loci were originally identified from the CONDOR database (Woolfe et al., 2007) (http://condor.nimr.mrc.ac.uk/). Subsequently, sequences from multiple species were extracted from Ensembl (Hubbard et al., 2009) (http://www.ensembl.org/index.html). These were aligned using MLAGAN (http://lagan.stanford.edu/lagan_web/index.shtml) (Brudno et al., 2003), with a Vista graphical output (Mayor et al., 2000). At the time that this analysis was performed, zebrafish pax2a/b loci had assembly errors, so Fugu was used as the model organism for comparative genomics and functional analyses. ClustalW (Thompson et al., 1994) was used for the alignment of individual CNEs.

Sequence conservation indices were calculated as a product of the proportion of sequence overlap between human and Fugu CNEs, and the proportion of identical bases, i.e. (length of overlapping Fugu sequence/length of human CNE) × number of identical bases/length of human CNE. These are reported in the text as n±s.d.

PCR design

Nineteen pairs of CNEs that are retained in duplicate (are associated with both Fugu pax2 co-orthologues) were selected from intergenic and intronic regions of the loci. These range in size from 57 to 432 bp and their percentage of shared sequence identity ranges from 77-97%. Where possible, oligonucleotides were designed using Primer 3 software (Rozen and Skaltsky, 2000). Otherwise, in order to be as close as possible to the CNE sequence, they were designed by eye, maximising the criteria for optimal primer design (as stipulated in Primer 3). CNEs were amplified and purified as described previously (Woolfe et al., 2005). CNE and oligonucleotide sequences are provided in the supplementary information (see Table S1 in the supplementary material).

Functional assay in zebrafish embryos

Purified CNEs were co-injected together with a GFP reporter gene under control of a human β-globin minimal promoter as previously described (Woolfe et al., 2005). Micro-injections were performed in one- to four-cell zebrafish embryos (day 1). Embryos were screened for GFP-positive cells and scored on day 2 and day 3 as described previously (Woolfe et al., 2005). Schematic diagrams and numbers of embryos with GFP expression in each domain have been deposited in our online database (http://condor.nimr.mrc.ac.uk/). At least 25 embryos were scored for each CNE assayed.

We also performed control experiments to demonstrate that our results using this co-injection assay are comparable with a more conventional cloning strategy using Gateway Tol2 cloning vectors. We show these results in Fig. S5 and Table S2 in the supplementary material. Briefly, elements were either blunt-end cloned into a Smal-digested 228 p5E-MCS vector (kindly provided by the Chien laboratory) (Kwan et al., 2007) or cloned into pENTR™-TOPO TA (Invitrogen #K591-20), after adding adenine overhangs (10-minute extension at 72°C using NEB Taq DNA polymerase, #M0273). A one-way LR reaction using LR Clonase II Plus (Invitrogen #12538-120) was then used to clone into R4-L1 basEGFPpA To12, a vector containing a carp β-actin minimal promoter and modified for efficient single 5′ entry. (This vector was created using sequences from 353-pENTRbasEGFP and 426-pDext Tol2PA, kindly provided by the Lawson lab) (Villefranc et al., 2007). Injection mix (5 pl of 50 ng/μl DNA, 25 ng/μl of transposase mRNA) was injected into one-cell embryos. The Transposase mRNA was synthesised using Ambion mMESSAGE mMACHINE (Invitrogen #AM1340M).

RESULTS AND DISCUSSION

The distribution of CNEs around the PAX2/5/8 genes

Following WGD events, CNEs are often asymmetrically partitioned around gene duplicates (Woolfe and Elgar, 2007). The PAX2/5/8 genes are no exception as they are, respectively, associated with around 60, 16 and two CNEs across all representative gnathostome groups (http://condor.nimr.mrc.ac.uk/) (Fig. 1; see Figs S1 and S2 in the supplementary material). By contrast, the teleost pax2 co-orthologues have retained a similar number of CNEs, about two-thirds of which exist in duplicate (Woolfe and Elgar, 2007) (Fig. 1). Of these duplicate CNEs, four share sequence homology with pax5 CNEs and another shares sequence homology with a pax8 CNE (Fig. 1; see Fig. S3 in the supplementary material). In all cases, duplicate CNEs are in a similar position relative to the protein-coding region of the gene. There are no CNEs retained between all three paralogues and none are retained between pax5 and pax8 (Fig. 1; see Figs S1 and S2 in the supplementary material). Owing to the abundance of CNE duplicates retained between the teleost pax2 co-orthologues and to the fact that we can compare them to the single copy tetrapod CNEs, we have used this dataset in order to explore CNE functionality subsequent to a WGD event.

A majority of pax2 CNEs exhibit enhancer activity

Using an in vivo co-injection reporter gene assay in zebrafish embryos, we tested 19 pairs of pax2 elements that are retained in duplicate between Fugu pax2 co-orthologues. Remarkably, over 80% (31/38) of the elements are able to drive reporter gene expression in a tissue-specific and reproducible manner (Fig. 2), and in the majority of cases this strongly overlaps with endogenous gene expression. For example, most elements (27/38) drive expression in the hindbrain and spinal cord, and expression in thyroid and pronephros regions is also frequently observed (24 and 21 elements, respectively).

Duplicate pax2 CNEs have diverged in function

Strikingly, we observed more differences than similarities between the expression profiles of duplicate CNEs. Most obviously, there are four pairs of CNEs (4, 10, 11 and 17) in which one CNE is able to drive reporter gene expression (interestingly always the pax2.2 CNE), while the other shows little or no GFP expression (Figs 2 and 3).
The CNE pair with the most similar expression profile (15) drives expression predominantly in muscle but even here there are noticeable differences between individual CNEs. As well as spatial differences (the pax2.1 element also drives expression in the CNS), there are temporal differences. Compared with day 2, there is a fourfold increase (39% versus 9%) in the percentage of GFP-positive embryos on day 3 for the pax2.1 CNE, whereas this is less pronounced for the pax2.2 CNE (45% versus 34%; Fig. 2).

In terms of spatial expression, the most dramatic difference is observed with CNE pair 1. In this case, the pax2.1 CNE activates GFP in only a few regions, with most expression in the notochord and some expression in muscle and fin (Figs 2 and 3). By contrast, the pax2.2 CNE drives GFP expression in a highly complex pattern in virtually all of the domains scored in this assay, except the notochord and the fin. This is an extraordinary result given the high similarity in sequence of these CNEs, both in terms of overlap (see Fig. S3 in the supplementary material) and sequence identity (~90%).

Comparison of duplicate CNE sequences
These striking differences in expression profiles led us to compare carefully the two sequences that we were using to test each pair of duplicate CNEs. In some cases (e.g. 8) the teleost CNEs overlap asymmetrically with the human sequence, and others (e.g. 2 and 7) are of unequal length (see Fig. S3 in the supplementary material). However, over one-third (37%) differ in length by less than 10 bp and align to the equivalent region of the single human PAX2 locus (see Fig. S1 in the supplementary material). Although almost half (47%) of the pax2.2 CNEs are shorter than the pax2.1 counterparts, their average conservation indices are identical [respectively, 0.805±0.077 (s.d.) and 0.795±0.068 (s.d.)] suggesting a similar rate of divergence. In two cases (pairs 5 and 9) one CNE duplicate (pax2.1 and pax2.2, respectively) contains an internal gap in conservation, suggesting a fragmentation of the CNE.

Non-conserved flanking sequences influence enhancer activity
Intriguingly, as mentioned above, the pair 1 elements have highly similar overlapping sequences, and yet, their expression profiles are dramatically different. The PCR products for this CNE pair included very short additional sequences (50 bp 5′/H11032 and 3′/H11032, respectively, for the pax2.1 and pax2.2 elements). In order to rule out any potential influence from these sequences, we repeated our assay using PCRs derived from sub-optimal primers located at the boundaries of the CNE sequence.

Interestingly, we see a striking difference in the expression profiles of these more tightly defined CNE sequences, compared with the original elements (Fig. 3). Expression driven by the new pax2.1 element is more complex and includes domains (e.g. cardiovascular) that overlap with those produced by the pax2.2 element. In contrast to the original element, the new pax2.2 element drives expression in fin, more expression in muscle, less expression in CNS and no expression in sensory organs. However, although the new pax2.1/pax2.2 expression profiles are more similar than the original ones, they are still not identical. Unlike the pax2.2 element, the pax2.1 element does not drive expression in CNS but does drive expression in the otic vesicle. Likewise, the pax2.2 expression profile has lower expression in skin and fin, higher cardiovascular expression and lacks notochord expression. Therefore, even highly similar sequences (with 88% sequence identity) can be functionally divergent.
These remarkable results led us to re-analyse another pair of elements, pair 4, which have substantial differences in their expression profiles (Figs 2 and 3). Whereas the pax2.2 element shows strong enhancer activity (over 30% of embryos have GFP expression), the pax2.1 element drives very little expression (0.8% and 2.7% of embryos express GFP on day 2 and day 3, respectively). In this case, the pax2.2 element extends beyond the defined CNE region but shows good overlap and alignment with the pax2.1 element (see Fig. S4 in the supplementary material). However, the pax2.1 sequence incorporates an additional 50 bp 5′ of this alignment (see Fig. S3 in the supplementary material). Therefore, we re-assayed this element eliminating this extra sequence.

This new element has slightly more enhancer activity than the original one (21/512 embryos (4%) expressed GFP). This expression occurred in similar domains to the pax2.2 element (in the hindbrain and spinal cord), but also elsewhere (the tectum and forebrain; Fig. 3). However, on day 3 we could detect GFP in only five out of 421 embryos (1.2%). Therefore, even this more tightly defined CNE sequence only has very weak enhancer activity, which is still significantly different from its duplicate pax2.2 CNE.

Isolated cis-regulatory sequences drive expression in both endogenous and ectopic domains

Our co-injection assay consistently shows that the majority of CNEs are able to drive reporter gene expression in a tissue-specific and reproducible manner. Our results are consistent with results that we obtained using more conventional Tol2 cloning strategies and they are apparently independent of the basal promoter used (see Fig. S5 and Table S2 in the supplementary material). In most cases, GFP expression recapitulates endogenous gene expression, with most CNEs driving expression in hindbrain and spinal cord, many driving expression in thyroid and pronephros regions, and about half driving expression in sensory organs. This strongly suggests that these sequences normally regulate Pax2 expression. However, many of the CNE expression profiles are complex, and frequently include ectopic domains. This may be because we are testing these elements as isolated sequences outside the influence of the genomic environment.

Ectopic expression is a common, but little discussed, occurrence in assays using isolated cis-regulatory sequences and transgenic lines constructed using subpopulations of cis-regulatory elements (see Goode and Elgar, 2009). For example, mouse (Ohyama and Groves, 2004; Rowitch et al., 1999) and zebrafish (Picker et al., 2002) pax2 transgenic lines recapitulate many aspects of endogenous Pax2 expression, but they also have ectopic expression and expression within endogenous domains is not always temporally correct. Strikingly, none of these transgenic lines exhibits expression in the eye, even though an optic stalk enhancer (Schwarz et al., 2000) is embedded within the upstream sequences used to generate these lines. The potential activity of an element can, therefore, be missed if it is not analysed in isolation. However, our results show that CNEs often have the ability to drive reporter gene expression that both overlaps with and occurs outside endogenous domains. Notably, regardless of methodology (see Fig. S5 and Table S2 in the supplementary material) even the ‘ectopic’ expression is reproducible across a large number of embryos. This indicates the potential enhancer properties of CNEs, which can be seen when they are tested in isolation but which are presumably latent (or repressed in some way) when these sequences are in their normal context within the genome. Cumulative data such as these are invaluable for determining the underlying sequence language that can regulate gene expression and also for identifying potential sequences that might be responsible for anomalous gene expression when the gene regulatory landscape is perturbed by mutations or translocations.

Highly similar sequences can have divergent functions

Strikingly, our results show that even highly conserved duplicate CNE pairs can have different expression profiles. This illustrates the relative ease with which functionality can evolve with little sequence change. Like the single tetrapod Pax2 gene, the teleost co-orthologues have complex expression patterns in the CNS, eye and ear. However, although their expression overlaps in these domains, there are temporal-spatial differences. In addition, only pax2a is expressed in the kidney and thyroid (Goode and Elgar, 2009; Pfeffer et al., 1998). This functional partitioning is consistent with the DDC model and experimental evidence shows that the co-
orthologues are able to functionally substitute for one another in overlapping expression domains (reviewed by Goode and Elgar, 2009). The retention of both duplicates therefore adds a robustness to some Pax2 functions while allowing others to diverge between the duplicated genes. From our analyses of Pax2 CNEs, we can readily appreciate how subtle sequence changes within these elements could rapidly expand the already complex range of pax2 functions, thus influencing the evolving animal body plan. Given our results, it is crucial that assays of CNEs should use sequences that are as accurately delineated as possible. Even our initial stringent primer design strategy included short additional sequences, and yet as we have shown, these can dramatically influence the activity of CNEs. This is a crucially important finding, given the prevalence with which CNEs are now used to try and construct transgenic animals and/or drive expression in specific cells and tissues.

![Fig. 3. Schematic diagrams of expression profiles derived from assaying pax2 element pairs 1 and 4. Expression in each domain is colour coded according to the key in Fig. 2 and mapped onto camera lucida drawings of day 2 and day 3 zebrafish embryos (day 2 is shown here). Results are overlaid from multiple embryos. *n* indicates the number of embryos analyzed. The percentage of GFP-positive embryos with expression in each domain (y-axis) is indicated in the respective bar charts. For pair 1, ‘old’ indicates results from our original assay and ‘new’ indicates results from our newer assay without the flanking sequences. Live images of day 3 embryos are also shown for these assays, with expression in the notochord (pax2.1 old), eye (pax2.2 old), otic vesicle (pax2.1 new) and fin (pax2.2 new). Expression in the eye is shown as a fluorescent image, whereas the rest are shown as merged fluorescent and bright-field images. b, blood; e, eye; f, fin; fb, forebrain; h, heart; hb, hindbrain; m, muscle; mb, midbrain; n, notochord; o, otic vesicle; p, pronephric region; s, skin; sc, spinal cord; t, thyroid region; x, other, unclassified; y, yolk. Scale bars: 50 μm.](image-url)
Importance of our results for cross-species comparisons of CNEs

A powerful way of assessing sub-functions embedded within duplicate CNEs is to use phylogenetic and functional analyses of multiple vertebrate sequences. Here, we have shown the potential behind comparing duplicate teleost CNEs with single-copy tetrapod sequences. Extending such analyses should enable us to identify which base changes are permissive during the evolution of these sequences and how subtle sequence differences affect the regulatory ability of CNEs. Multi-species comparisons also allow us to delineate putative cis-regulatory elements in terms of the boundary of sequence conservation. Given that as little as 50 bp of non-conserved flanking sequence can dramatically influence CNE enhancer activity, it is obviously important to identify the limits of CNE sequences as accurately as possible. Our results strongly suggest that once a consensus has been reached, these boundaries need to be strictly adhered to when analysing CNEs if we are to be able to compare functional data from related genes and different species. Only then will we be able to confidently interpret functional data in order to generate an evolutionary profile of these extraordinary sequences.

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Competing interests statement

The authors declare no competing financial interests.

Supplementary material

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