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Evolutionary history and epigenetic regulation of the three paralogous *pax7* genes in rainbow trout

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Abstract

The extraordinary muscle growth potential of teleost fish, particular those of the *Salmoninae* clade, elicits questions about how the relatively highly conserved transcription factors of the myogenic program are regulated. In addition, the pseudotetraploid nature of the salmonid genome adds another layer of regulatory complexity, and this must be reconciled with epigenetic data to better understand how these fish achieve lifelong muscle growth. To this end, we identified three paralogous *pax7* genes (*pax7a1*, *pax7a2*, and *pax7b*) in the rainbow trout genome. During *in vitro* myogenesis, *pax7a1* transcripts remain stable, while *pax7a2* and *pax7b* mRNAs increase in abundance, similarly to *myogenin* mRNAs and in contrast to the expression pattern of the mammalian ortholog. In addition, we profiled the distribution of repressive H3K27me3 and H3K9me3 and permissive H3K4me3 marks during *in vitro* myogenesis across these loci, finding that *pax7a2* expression was associated with decreased H3K27 trimethylation, while *pax7b* expression was correlated with decreased H3K9me3 and –K27me3. Altogether, these data link the highly unique differential expression of *pax7* paralogs with epigenetic histone modifications in a vertebrate species displaying growth divergent from that of mammals and highlight an important divergence in the regulatory mechanisms of *pax7* expression among vertebrates. The system described here provides a more comprehensive picture of the combinatorial control mechanisms orchestrating skeletal muscle growth in a salmonid, leading to a better understanding of myogenesis in this species and across *Vertebrata* more generally.

Keywords

pax7; epigenetic regulation; histone methylation; ChIP; rainbow trout

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INTRODUCTION

Many teleost fish possess the extraordinary ability to augment their skeletal musculature throughout life (termed indeterminate muscle growth) as a result of the continuous formation of new fibers (myofiber hyperplasia) and increased size by hypertrophy of existing myofibers (Mommensen 2001, Johnston et al. 2011). The muscle stem cells (called myosatellite cells, MSCs) are thought to be essential for these two processes (Lepper et al. 2011, Sambasivan et al. 2011). Upon commitment, these cells engage into a myogenic program, allowing them to proliferate, differentiate, and fuse with existing myofibers (hypertrophy) or with other myoblasts to form new myofibers (hyperplasia). It is assumed that MSCs are controlled by highly conserved molecular networks that regulate myogenesis across growth paradigm and taxa (determinate and indeterminate growth; mammalian, fish, and even invertebrate clades; Konstantinides and Averof 2014), but that these networks vary enough in expression patterns to allow MSCs from indeterminate growing organisms to continuously contribute to either hyperplasia and/or hypertrophy in the adult animal.

In this regard, we have shown that MSCs isolated from rainbow trout (an indeterminate growing fish) enter into the differentiation program by very quickly expressing the myogenic regulatory factor (MRF) transcription factors myoD1 and myogenin (Gabillard et al. 2010). Interestingly, an “unexpected” increase of *pax7* expression (a well-accepted marker of quiescent MSCs in mammals) in culture of MSCs under differentiation into myotubes has been documented in a closely related species (*Salmo salar*), possibly reflecting the production of “quiescent reserve cells” (Bower and Johnston 2010). Likewise, we recently demonstrated that MSCs from the indeterminately growing giant danio (*Devario aequipinnatus*) in culture gradually increase *pax3* expression (another marker of quiescent MSCs in mammals) during a similar progression to myotubes (Froehlich et al. 2013). These data support the hypothesis that the induced expression of these specific paired-box transcription factors (*pax3/7*) during the culture of MSCs is a common feature of indeterminate growing species and could indicate a high self-renewal of MSCs, a process needed for these cells to continuously contribute to hyperplasia and/or hypertrophy during the lifespan of these fishes (Froehlich et al. 2013). However, very few data exist detailing how the expression of these Pax and MRF transcription factors is regulated in indeterminate growing organisms.

In mammals, the regulation of several genes involved in MSC activity has been shown to be under the strict control of epigenetic mechanisms through histone methylation (Perdiguerro et al. 2009, Saccone and Puri 2010). The consequences of histone methylation on transcriptional activation or repression depend on the site and degree of methylation. Histone lysine methylation can occur on histones H3 and H4 in monomethyl, dimethyl, or trimethyl states. Trimethylation of lysine 4, lysine 36, or lysine 79 of histone H3 (H3K4, H3K36, or H3K79) is generally associated with euchromatin and transcriptional activity (i.e., a ‘permissive’ state; Wang et al. 2001, Feng et al. 2002, Krogan et al. 2003, Krogan et al. 2003). Conversely, di- or trimethylation of lysine 9 or lysine 27 of histone H3 (H3K9 or H3K27) and trimethylation of lysine 20 on histone H4 (H4K20) are correlated with heterochromatin formation and transcriptional repression (Rea et al. 2000, Nakayama et al. 2001, Tachibana et al. 2001, Cao et al. 2002, Schultz et al. 2002, Schotta et al. 2004).

Histone methylation, however, is not a permanent state. Thus, genomic regions are subjected to the opposing actions of methylation and demethylation in a dynamic manner, thereby ensuring a transcriptional response to altered organismal and/or environmental conditions. In this regard, the coordination of methylation events at H3K4, H3K9, and H3K27 in the promoters of genes coding for various Pax and MRF transcription factors (*pax3/7*, *myf5*, *myoD1*, and *myogenin*) have been demonstrated to regulate many aspects of MSCs activation, proliferation, and differentiation in mammals (Sebastian et al. 2009, Palacios et al. 2010, Stojic et al. 2011, Taberlay et al. 2011, Tao et al. 2011, Diao et al. 2012, Kawabe et al. 2012, Ling et al. 2012, Sdek et al. 2013, Wang et al. 2013). However, no information detailing the epigenetic regulation of myogenesis in indeterminate growing organisms is available in the literature. In the present study, we therefore sought to profile the distribution of H3K27me3, H3K9me3 and H3K4me3 marks during the *in vitro* myogenesis in rainbow trout across loci with very different roles in the control of myogenesis: *pax7*, in the commitment and survival of MSCs; and *myogenin*, in the terminal differentiation of myotubes.

MATERIALS AND METHODS

In silico analysis of *pax7* genes

The presence of *pax7* genes was queried against the rainbow trout genome (Berthelot et al. 2014) with BLAST search in SIGENAE databases (<http://www.sigena.org/>). The new sequences of rainbow trout *pax7a1*, *pax7a2* and *pax7b* genes are available in Genoscope database (www.genoscope.cns.fr/trout) under the numbers GSONMG00081386001, GSONMG00061433001 and GSONMG00027288001, respectively. The phylogenetic analysis was carried out with Pax7 amino acid sequences available on Ensembl Genome database (*Homo sapiens*, Ensembl ID: ENSP00000403389; *Mus musculus*, Ensembl ID: ENSMUSP00000030508; *Danio rerio*, Ensembl ID: ENSDARP00000123695 and ENSDARP00000108935; *Astyanax mexicanus*, Ensembl ID: ENSAMXP00000003905 and ENSAMXP00000025517) and GenBank (*Branchiostoma floridae*, GenBank ID: AAF89581.1). Phylogenetic tree was constructed by the Neighbor-Joining (NJ) method using the Molecular Evolutionary Genetics Analysis (MEGA) software version 6.0 (Tamura et al. 2013). The reliability of the inferred tree was estimated using the bootstrap method with 500 replications. The amphioxus Pax7 sequence was used to root the tree. The Genomicus software program (<http://www.genomicus.biologie.ens.fr/genomicus-trout-01.01/cgi-bin/search.pl>) was used to identify syntenic genes that were located near the *pax7* genes of different vertebrate species.

Isolation of trout myosatellite cells

For all studies, MSCs were isolated from juvenile rainbow trout (*Oncorhynchus mykiss*) (15–45 g, depending on culture and availability) essentially as previously described (Froehlich et al. 2014). Isolated MSCs were plated on poly-L-lysine and laminin-coated plates at 200,000 cells per cm² in 6-well (chromatin immunoprecipitation, ChIP; or real-time quantitative PCR, RT-qPCR) or 24-well (immunocytochemistry) plates and cultured in 10% FBS-DMEM for 4 days (with cell fixation on days 2 and 4) and 2% FBS-DMEM for an additional 4 days (with cell fixation on day 8 post-isolation). Day 2 of culture represents

MSCs, day 4 myoblasts, and day 8 nascent myotubes as shown previously (Gabillard et al. 2010).

Analysis of cell proliferation

On days 2, 4, and 8 of culture, 10 μ M BrdU was added and cells were cultured for the following 24 h. The cells were then fixed with ethanol/glycine buffer (70% ethanol, 50 mM glycine, pH 2). The proliferation of the cells was measured by using a BrdU labeling and detection kit (Roche Diagnostics, no. 1 296 736). Briefly, the cells were incubated for 30 min at 37 °C with mouse anti-BrdU, washed, and then incubated with the secondary antibody anti-mouse FITC for 30 min. Cells were then mounted in Mowiol containing 0.5 μ g/ml DAPI.

Immunofluorescence analysis

Cells on glass coverslips were briefly washed twice with phosphate-buffered saline (PBS) and fixed for 10 min with 4% paraformaldehyde in PBS. For permeabilization, cells were incubated 3 min in 0.1% Triton X-100/PBS. After three washes, cells were saturated for 1 h with 3% BSA, 0.1% Tween-20 in PBS (PBST). Cells were incubated for 3 h with the primary antibodies anti-myosin heavy chain (MyHC) and anti-myogenin diluted in blocking buffer (Gabillard et al. 2010). The secondary antibodies were diluted in PBST and applied for 1 h. Cells were mounted with Mowiol 4-88 (#475904, Calbiochem) containing Hoescht (0.5 μ g/ml). Cells were photographed using a Canon digital camera coupled to a Canon 90i microscope.

Gene expression analysis

Total RNA samples were extracted from cells using TRIzol reagent (Invitrogen, Carlsbad, CA, USA), according to the manufacturer's recommendations. One microgram of the resulting total RNA was reverse transcribed into cDNA using the SuperScript III RNaseH-Reverse Transcriptase kit (Invitrogen, Carlsbad, CA, USA) and random primers (Promega, Charbonnières, France), according to the manufacturer's instructions. Target gene expression levels were determined by RT-qPCR using specific primers (Table 1). The primer sequences were chosen based on the rainbow trout genes sequence available in the expressed sequence transcript database (<http://www.sigenae.org/>). To avoid amplification of genomic DNA, when possible, primer pairs included one intron-spanning oligonucleotide. Real-time RT-PCR was carried out on an iCycler iQ™ real time PCR detection system (BIO-RAD, Hercules, CA, USA) using iQ™ SYBR® Green Supermix. Relative quantification of the target gene transcripts was made using *EF1a* gene expression as reference and following the Pfaffl method with the Relative Expression Software tool (REST©) (Pfaffl 2001, Pfaffl et al. 2002). PCR was performed using 10 μ l of the diluted cDNA mixed with five picomoles of each primer in a final volume of 25 μ l. The PCR protocol was initiated at 95°C for 3 min for initial denaturation of the cDNA and hot-start iTaq™ DNA polymerase activation followed by a two-step amplification program (20 sec at 95°C followed by 30 sec at specific primer hybridization temperature) repeated 40 times. Melting curves were systematically monitored (temperature gradient at 0.5°C/10 sec from 55 to 94°C) at the end of the last amplification cycle to confirm the specificity of the amplification reaction. The different

PCR products were initially sequenced to confirm the identities of the amplicons. Each PCR run included replicate samples (duplicate of reverse transcription and duplicate of PCR amplification) and negative controls (reverse transcriptase-free samples, NRT; RNA-free samples, NTC).

Chromatin Immunoprecipitation (ChIP)

On days 2, 4, and 8 of culture, MSCs, myoblasts, or nascent myotubes were fixed in 1% methanol-free formaldehyde (16% diluted in serum-free DMEM immediately prior to fixation) for 10 min at room temperature. Formaldehyde was neutralized by the addition of 2.5 M glycine for 5 min. Fixed cells were then washed twice in ice-cold 0.01 M PBS. Next, 1 mL of ice-cold 0.01 M PBS containing protease inhibitors (HALT™; Pierce) was added, and three wells were pooled for each ChIP sample (i.e., negative/mock ChIP control, total H3, H3K4me3, H3K9me3, and H3K27me3) by scraping cells into a microcentrifuge tube and pelleting cells at 3000×g for 5 min. Samples were stored at –80°C until preparation of chromatin and subsequent immunoprecipitation.

Chromatin preparation and subsequent immunoprecipitation were completed using a commercial kit (Pierce™ Agarose ChIP Kit) according to the manufacturer's instructions (Pierce) with modifications. Nuclei were extracted using a membrane extraction buffer spiked with HALT™ cocktail (Pierce kit) and centrifuged at 6000×g for 3 min. Intact nuclei were then resuspended in 10 mM Tris/1 mM EDTA/1% SDS sonication buffer and sonicated 13-15 times on ice (15 sec pulses followed by 2 min rests) until chromatin was 100-800 bps in size, with the center being ~300 bp. Prior to incubation with primary antibodies, chromatin was cleared of any cross-reactivity to agarose by incubating the isolated chromatin with agarose beads (5x input ChIP volume; Pierce Agarose Negative Control Resin) for 6 hours at 4°C before passing chromatin through columns provided with the kit. Chromatin was diluted in 1x dilution buffer (provided with Pierce kit) 1:2 before the addition of antibodies. Cross-reactive antibodies, previously identified as appropriate for use in zebrafish (Wardle et al. 2006), were obtained from Abcam (Cambridge, MA) and used at dilutions recommended by the manufacturer (10 µg normal rabbit IgG; 10 µg anti-H3; 10 µg anti-H3K4me3; 10 µg anti-H3K9me3; 10 µg anti-H3K27me3 per ChIP) with overnight (~18 h) incubations. DNA-protein complexes were immunoprecipitated using protein A/G-agarose beads, blocked in sonicated DNA/native DNA isolated from CHO cells (~3 million per 0.6 ml of beads) and bovine serum albumin (10 mg/ml) prior to use, for 1 hour, followed by three 5 min washes with each buffer (3 buffers = 9 washes; Pierce kit). Elution from agarose beads was performed as recommended by the manufacturer. Formaldehyde crosslinks of protein to DNA were released by overnight (~18 hour) incubation at 65°C in 5M NaCl/elution buffer using proteinase K (20 mg/ml; Pierce). Following DNA isolation using the provided columns, upstream regions of interest, identified using previous mammalian data (Palacios et al. 2010, Seenundun et al. 2010), as well as the ChIP-seq data from the Encyclopedia of DNA Elements (ENCODE) database available at the UCSC Genome Browser (<http://www.genome-euro.ucsc.edu/index.html>), were analyzed by qPCR using primer sets designed in Primer3 (Rozen and Skaletsky 2000) (see Table 2) using the following cycling parameters: 1 cycle, 95°C for 3 min; 40 cycles, 95°C for 15 sec, 60°C for 30 sec, 72°C for 15 sec. Samples were run in duplicate, with the appropriate 10%

input and DNA-free controls and dissociation curves following amplification. In advance of both types of PCR analysis, primer sets were validated using genomic DNA isolated from rainbow trout MSCs and found to amplify intended targets of interest. All primer sets were validated to recognize gene promoter regions enriched in H3 proteins using total H3 immunoprecipitation. Using C_q values, % input was calculated for each experimental and mock sample.

Statistical analysis

Results for proliferation, differentiation and RT-qPCR are expressed as means $\pm$ SEM and analyzed by one-way ANOVA followed by Student-Newman-Keuls test. For ChIP data, enrichment is expressed as percent-input (% input) after correction for dilution (1:10; $-3.32 C_q$). Data were analyzed by one-way ANOVA followed by Tukey's post-hoc test. For all statistical analyses, the level of significance was set at $P < 0.05$. All the experiments were performed twice. Within an experiment, results for each time point were obtained from three replicates (for the ChIP analysis), four replicates (for the proliferation and differentiation assays) and six replicates (for the RT-qPCR analysis).

RESULTS

Three orthologs of mammalian *pax7* exist in rainbow trout genome

Using the recent availability of the rainbow trout genome assembly (Berthelot et al. 2014), we identified three genes (Genoscope accession number: GSONMG00081386001, GSONMG00061433001 and GSONMG00027288001) displaying high homology (E-value $> 2e-40$, Sigenae tblasn <http://www.sigenae.org/>) with the two trout sequences of *pax7* previously available in GenBank (GenBank ID: NM_001258337 and NM_001258336). The phylogenetic analysis performed using the full-length Pax7 protein sequences of several vertebrate species and the amphioxus (*Branchiostoma floridae*) Pax3/7 sequence as an outgroup showed that the two trout sequences (GSONMG00081386001 and GSONMG00061433001), sharing the highest identity with the previously annotated NM_001258336 and NM_001258337 (99.7% and 100% identity, respectively), clustered with zebrafish (*Danio rerio*) and cave fish (*Astyanax mexicanus*) Pax7a, while the third sequence (GSONMG00027288001) was most similar to zebrafish and cave fish Pax7b (Fig. 1). These results suggested that the two former trout genes (GSONMG00081386001 and GSONMG00061433001) are paralogous genes and co-orthologous to the characid and cyprinid *pax7a* gene, while the last gene (GSONMG00027288001) was orthologous to the characid and cyprinid *pax7b* gene.

Our syntenic analysis showed that the tetrapod *pax7* gene was localized in the igsf21-tas1r2-kiaa0090-akr7a3 syntenic group (Fig. 2). Interestingly, a syntenic conservation of this region was found with two distinct chromosomal regions of the zebrafish genome, which included the duplicated copies of *pax7*, *tas1r3* and *akr7a3* (Fig. 2). In addition, duplicated copies of *errf11* (a and b) were found to be syntenic with *pax7a* and *pax7b*. The syntenic region around the *pax7a* and *pax7b* loci in zebrafish was conserved in other teleost species but in a lesser extent. Considering the newly sequenced rainbow trout genome, our syntenic analysis showed that GSONMG00081386001 (*pax7a1*) and GSONMG00061433001 (*pax7a2*) genes

were localized on two distinct scaffolds (scaffold_8 and scaffold_1156 carrying the genes GSONMG00081386001 and GSONMG00061433001, respectively). The synteny around *pax7a* locus in zebrafish and medaka (*akr7a3a-park7-errfi1a-tas1r3*) was well conserved around these two genes pointing out an additional duplication of *pax7a* in rainbow trout, thus giving birth to *pax7a1* and *pax7a2*. As regard the third identified trout *pax7* gene, the syntenic analysis showed that it was localized in the *zgc:154075-errfi1b-pax7b-akr7a3b* syntenic group conserved around the *pax7b* locus in zebrafish, confirming that this gene is related to *pax7b* of other teleosts. Interestingly, the synteny around the *pax7b* locus revealed a certain degree of chromosomal rearrangement in this genomic region between the zebrafish and the rainbow trout on one side and other teleost (e.g., medaka, stickleback and fugu) on the other, although the genetic environment is in overall conserved. Furthermore, we did not find any duplication of genes surrounding *pax7b* locus (*errfi1b-pax7b-akr7a3b*) in the trout genome, whereas our analysis revealed a duplication of the close locus *tas1r1b1-igsf21b-vps13d*. Taken together, these results revealed that the rainbow trout genome holds three *pax7* paralogs, namely *pax7a1* (GSONMG00081386001), *pax7a2* (GSONMG00061433001) and *pax7b* (GSONMG00027288001).

***pax7* paralogous genes are differently expressed during *in vitro* myogenesis**

MSCs isolated from white muscle of juvenile rainbow trout were then used to monitor the expression of *pax7a1*, *pax7a2* and *pax7b* during *in vitro* myogenesis. As shown in Fig. 3a, the cells (seeded at day 0) started to proliferate (as determined by measuring the incorporation of BrdU) at day 2 of culture and reached a peak at day 4. The proliferation rate subsequently decreased after 8 days of culture. Concurrently, the myogenic index of cultured cells (as determined by myogenin and myosin heavy-chain, MyHC, immunoreactivity) increases from day 2 to day 8 (Fig. 3b). These results demonstrated that isolated trout MSCs progress through the myogenic program to myotubes and validated their use as an *in vitro* model of myogenesis. In this regard, *myoD1* and *myogenin* transcripts increased at day 8 of culture and those of the S phase marker *PCNA* paralleled the incorporation of BrdU (Figs 4a-c). Interestingly, *pax7a1* transcripts remained stable during the culture period and those of *pax7a2* and *pax7b* increased from 2 to 8 days (Figs 4d-f). Together these results suggest that the control of the expression of these *pax7* paralogs diverged during evolution and make these genes relevant models for a better understanding of the evolutionary fate of duplicated genes.

H3K27me3, H3K9me3, and H3K4me3 profiling across the three *pax7* loci reveals a complex regulatory schema

The regulation of the expression of several genes involved in myogenesis has been shown to be under the strict control of epigenetic imprinting through histone methylation. Therefore, ChIP was used to profile the distribution of H3K27me3, H3K9me3 and H3K4me3 across *myogenin* and *pax7* loci (Figs 5a and 6a) during the *in vitro* myogenesis in rainbow trout. The results show an enrichment in repressive H3K27me3 and H3K9me3 marks throughout the upstream regions of *myogenin* in day 2, followed by an extensive loss of these histone modifications thereafter at day 4 and day 8 (Figs 5 b, c and c, i). In contrast, H3K4me3 levels remain relatively low at these loci in day 2 and display an increase at day 4 and 8 (Fig. 5d, g, j). Together these results fit with the increase of the mRNA levels of *myogenin*

from day 2 to day 8 (Fig. 4c) and suggest that the expression of *myogenin* is regulated through the antagonistic action of repressive H3K9me3/H3K27me3 and permissive H3K4me3 marks. In contrast, *pax7a1*, which presented a stable expression over the culture period (Fig. 4d), switched at the targeted locus from a transcriptionally repressive state of H3K27me3 and H3K9me3 marked chromatin to the transcriptionally permissive state of H3K4me3 marked chromatin, during the progression of MSCs to myotubes (Fig. 6b-d), suggesting that regulation may occur at a different target locus or through different means. Conversely, the increase in *pax7a2* and *pax7b* mRNA abundance (Fig. 4e and f) was associated to a statistical decrease in H3K27me3 levels for the former and in H3K27me3 and H3K9me3 levels for the last gene at the monitored locus (Figs 6e-j).

DISCUSSION

Since Neil Stickland (Stickland 1983) first documented both hyperplastic and hypertrophic skeletal muscle growth in the postlarval rainbow trout, there has been much interest in understanding how these fish are able to form such a large number of myofibers outside of the embryonic and larval periods of development. Of particular importance have been *in vitro* studies, where MSCs can be isolated to assess their potential contribution to the muscle growth of these fish (Gabillard et al. 2010, Froehlich et al. 2014). Using such an *in vitro* model for experimentation, Neil Bower and Ian Johnston (Bower and Johnston 2010) documented that *pax7* expression is up regulated during the differentiation of MSCs from Atlantic salmon (*Salmo salar*). *Pax7*, the canonical marker of MSCs in birds and mammals (Seale et al. 2000, Halevy et al. 2004), is rapidly down-regulated upon induction of MSC differentiation in rodents, birds, and *Xenopus* (Halevy et al. 2004, Zammit et al. 2004, Yamane and Nishikawa 2013), in agreement with its function in survival and quiescence of MSCs. Thus, the continued *pax7* transcript expression in salmonid fishes is highly unusual in light of the current literature and warrants further investigation.

In the present study, we identified three *pax7* paralogous genes in the recently available rainbow trout genome assembly (Berthelot et al. 2014). Based on phylogenetic analysis, we showed that two of these newly identified isoforms (*pax7a1*, *-a2*) belong to the *pax7a* clade while the third (*pax7b*) is orthologous to the characid and cyprinid *pax7b* gene. Previously, two *pax7* cDNA sequences had been available in GenBank: NM_001258337.1, initially named *pax7a* and now renamed *pax7a2*; and NM_001258336.1 initially renamed *pax7b* and now renamed *pax7a1*. The single promoter sequence originally deposited in GenBank by P.Y. Rescan and C. Ralliere (FJ713022.1) is that of our newly coined *pax7a2*. Together, the phylogenetic analysis and synteny conservation data strongly suggest that a single copy of *pax7* was present in the vertebrate ancestor and that a duplication of the *pax7* gene occurred with the teleost-specific whole-genome duplication (WGD) 225 to 333 million years ago (Amores et al. 1998, Jaillon et al. 2004, Hurley et al. 2007, Santini et al. 2009, Near et al. 2012), giving rise to *pax7a* and *pax7b* paralogs in the common teleost ancestor. Our results also showed that the two identified trout *pax7a1* and *pax7a2* genes are co-orthologous to the zebrafish *pax7a*, suggesting that they resulted from the additional and relatively recent salmonid-specific WGD event (the 4th salmonid-specific WGD or Ss4R) around 100 million years ago (Berthelot et al. 2014, Macqueen and Johnston 2014). However, we failed to identify any duplicated gene for the syntenic group containing *pax7b*, possibly due to the

recently described loss of half of the duplicated protein-coding genes in the rainbow trout genome (Berthelot et al. 2014). Taken together, these results demonstrate that the rainbow trout offers, with its three *pax7* paralog genes, a unique opportunity to better understand the divergence in function of the best marker of MSCs currently known.

Real-time qPCR analysis revealed that these three *pax7* paralogs are differentially expressed during the differentiation of isolated trout MSCs, in line with the recently described substantial number of Ss4R paralogs that has strongly diverged in terms of expression profiles and/or expression levels (Berthelot et al. 2014). While *pax7a1* transcript levels remained stable over the culture period, *pax7a2* and *pax7b* transcripts significantly increased on day 8 of culture, a time point in this cell culture model where myotubes have formed. Such an increase in *pax7* expression during the myogenic progression to myotubes has been documented in the closely related species *Salmo salar* (Bower and Johnston 2010). Similarly, we recently demonstrated that expression of *pax3* (another marker of MSCs in mammals) increases during a similar myogenic progression to myotubes in the indeterminately growing giant danio (*Devario aequipinnatus*) (Froehlich et al. 2013). In mammalian myogenic cell cultures, *pax7* is not expressed in differentiated myotubes, but is maintained in a smaller accompanying population of undifferentiated cells that stops proliferating and returns to a non-proliferating state reminiscent of the quiescent reserve MSCs, which can then reenter the cell cycle, proliferate, and differentiate (Kitzmann et al. 1998, Yoshida et al. 1998). By analogy with these data, our results may indicate that the increase in *pax3/7* expression during the myogenic progression to myotubes in indeterminate growing organisms may be involved with the continuous formation of new fibers through the enhanced production of reserve MSCs. In this regard, analyzing the mechanisms involved in the regulation of the expression of these genes is paramount to the understanding of how these fish are able to form such a large number of myofibers outside of the embryonic and larval stages of development and growth. An understanding of the mechanisms underlying differences in *pax7* paralog expression is also of particular importance from an evolutionary point of view. Following WGD, it is well accepted that the remaining paralogs can acquire new expression patterns, potentially leading to neo- or subfunctionalization (Force et al. 1999). This suggests that evolution after WGD may also act on the regulatory regions of paralogous genes, leading to subsequent silencing, sub-, or neofunctionalization. Thus, the identification of the regulatory mechanisms affected by and/or modified between paralogous genes may lead to a better understanding of the evolutionary fate of genes within a duplicated vertebrate genome.

In mammals, the transcriptional state of several muscle-specific genes such as *myogenin* or *pax7* has been shown to be associated with repressive H3K27me3 and H3K9me3 and permissive H3K4me3 chromatin marks (Mal 2006, Rampalli et al. 2007, Palacios et al. 2010, Mozzetta et al. 2011, Stojic et al. 2011). In the present study, we therefore sought to profile the distribution of H3K27me3, H3K9me3, and H3K4me3 marks across the three *pax7* and *myogenin* loci during *in vitro* myogenesis in the rainbow trout. The obtained results show that, during the myogenic progression to myotubes, the *myogenin* locus appears to be modified from a transcriptionally repressive state (i.e., H3K27me3 and H3K9me3 enriched chromatin) to a transcriptionally permissive state (i.e., H3K4me3 enriched

chromatin), in close agreement with previous findings obtained in C2C12 cell lines (Mal 2006, Rampalli et al. 2007, Seenundun et al. 2010). According to these studies, the proper temporal and spatial pattern of *myogenin* expression is established through the control of these repressive and permissive histone modifications mediated by the antagonistic functions of Polycomb group (PcG) and Trithorax group (TrxG) proteins (Mal 2006, Rampalli et al. 2007) as well as the action of histone lysine demethylases (Seenundun et al. 2010, Verrier et al. 2011). Although all of these factors remain to be characterized in fish, the parallel between the expression of *myogenin* and the monitored epigenetic marks in our cell culture model suggest a good conservation of these regulatory mechanisms among vertebrates. However, results obtained for the three trout *pax7* genes appear to indicate a more complex regulatory schema. Indeed, *pax7a1*, with stable expression over the culture period, switched from a transcriptionally repressive state (i.e., H3K27me3 and H3K9me3 enriched chromatin) to a transcriptionally permissive state (i.e., H3K4me3 enriched chromatin) during *in vitro* myogenesis. Such a discrepancy between mRNA expression and histone methylation may be the result of timing, and an increase in the expression of *pax7a1* may not become detectable until later time points. However, we cannot rule out the possibility that other post-translational modifications to exposed histone amino acid residues (acetylation, phosphorylation, mono-/di-/tri-methylation) may be at play in the control of the expression of this gene. Conversely, the increase in *pax7a2* and *pax7b* mRNA abundance was clearly associated with a statistical decrease in H3K27me3 levels for the *pax7a2* paralog and in H3K27me3 and H3K9me3 levels for the *pax7b* paralog. Interestingly, previous findings in other systems have demonstrated increasing levels of in the promoter region of *pax7* during the myotube differentiation of murine MSCs (Palacios et al. 2010). These contrasting results highlight a possible divergence in the regulatory mechanisms of *pax7* expression among vertebrates. Whether this difference in histone modifications and chromatin accessibility contributes to the increase in *pax7a2* and *pax7b* expression during the *in vitro* myogenic program is worth investigating. Further studies are also warranted to describe a role for the “atypical” expression of *pax7* paralogs in the extraordinary ability of trout to augment their skeletal musculature throughout life.

This report, linking the highly unique differential expression of *pax7* paralogs with epigenetic histone modifications, provides a platform for future studies for a better understanding of myogenesis in this indeterminate growing species and other teleost fishes. Information from these studies will be critical not only to the mechanistic understanding of the indeterminate growth paradigm, but also to a more complete understanding of the evolutionary fate of duplicated genes (and genomes) in *Vertebrata*.

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Figure 1. *pax7* phylogenetic analysis

The phylogenetic analysis was carried out using the Molecular Evolutionary Genetics Analysis (MEGA) software version 6.0. Phylogenetic tree was constructed using the Neighbor-Joining (NJ) method. The reliability of the inferred tree was estimated using the bootstrap method with 500 replications. The amphioxus *pax7* sequence (Branchiostoma floridae, GenBank ID: AAF89581.1) was used to root the tree. Proteins accession numbers are listed into the brackets.

Figure 2. *pax7* syntenic analysis

Conserved synteny around the *pax7* loci in tetrapods and teleosts. The Genomicus software program (<http://www.genomicus.biologie.ens.fr/genomicus-trout-01.01/cgi-bin/search.pl>) was used to identify syntenic genes that were located near the *pax7* genes of different vertebrate species.

Figure 3. Proliferation and differentiation of isolated trout MSCs cultured in DMEM with 10% fetal bovine serum (FBS) for 4 days and with 2% FBS for an additional 4 days
 (a), Proliferation of mononuclear myogenic cells at days 2, 4, and 8. Proliferation was determined by measuring the number of nuclei that incorporated BrdU during a period of 24 h. (b), Differentiation of myogenic cells at days 2, 4, and 8. Percentage of nuclei was determined by immunofluorescence analysis of myogenin (dark bars) and MyHC (clear bars). The different letters indicate significantly ($P < 0.05$) different means. Capital letters represent pairwise comparisons between myogenin across days, while lowercase letters represent pairwise comparisons between myosin. Significant differences were determined by

one-way ANOVA followed by Student-Newman-Keuls test for multiple comparisons. The presented experiment is representative of two experiments, and results for each time point were obtained from four replicates.

Figure 4. mRNA levels of *myoD1* (a), *PCNA* (b), *myogenin* (c), *pax7a1* (d), *pax7a2* (e) and *pax7b* (f) in isolated trout MSCs cultured in DMEM with 10% fetal bovine serum (FBS) for 4 days and with 2% FBS for an additional 4 days

mRNA levels of studied genes were estimated using real-time RT-qPCR at days 2, 4 and 8.

Values are means $\pm$ SE, n = 6. They were analyzed by one-way ANOVA followed by Student-Newman-Keuls test for multiple comparison ($P < 0.05$). Different letters represent significantly different values. The presented experiment is representative of two experiments.

Figure 5. Methylation of H3 at the *myogenin* gene in isolated trout MSCs cultured in DMEM with 10% fetal bovine serum (FBS) for 4 days and with 2% FBS for an additional 4 days (a), Schematic representation of the *myogenin* gene. ATG represents the start codon and the gray box indicates the open reading frame. FAR (far upstream region), DIS (distal upstream region) and PRO (proximal upstream region) indicate the location on the gene where each of the different primer sets localizes for analysis of ChIP studies. (b - j), ChIP analyses of the temporal (day 2, 4 and 8) changes in H3K27me3 (b, e, h), H3K9me3 (c, f, i), and H3K4me3 (d, g, j) enrichment across the targeted *myogenin* regions FAR (b-d), DISTAL (e-g), and PROXIMAL (h-j). Chromatin isolated from trout myogenic cells at various stages of proliferation and differentiation (Day 2, 4 and 8) was subjected to immunoprecipitation using either an anti-H3K27me3, anti-H3K9me3, or anti-H3K4me3 antibody. Immunopurified DNA was then quantified by qPCR using primer sets that recognize the different regions indicated in the schematic in Figure 5a. Values are means $\pm$ SE, $n = 3$ (mean of 2 replications). They were analyzed by one-way ANOVA followed by Tukey's

post-hoc test for multiple comparisons ($P < 0.05$). Different letters represent significantly different values. nd = no difference detected.

Figure 6. Methylation of H3 at the *pax7a1*, *pax7a2* and *pax7b* genes in isolated trout MSCs cultured in DMEM with 10% fetal bovine serum (FBS) for 4 days and with 2% FBS for an additional 4 days

(a), Schematic representation of the *pax7* paralogous genes. ATG represents the start codon and the gray box indicates the first exon. PRO (proximal upstream region) indicates the location on the *pax7* genes where the used primer sets localize for analysis of ChIP studies. (b-j), ChIP analyses of the temporal (day 2, 4 and 8) changes in H3K27me3 (b, e, h), H3K9me3 (c, f, i), and H3K4me3 (d, g, j) enrichment across the *pax7a1* (b-d), *pax7a2* (e-g), and *pax7b* (h-j) genes. Chromatin isolated from trout myogenic cells at various stages of proliferation and differentiation (Day 2, 4 and 8) was subjected to immunoprecipitation using either an anti-H3K27me3, anti-H3K9me3, or anti-H3K4me3 antibody. Immunopurified DNA was then quantified by qPCR using the primer set that recognize specifically the targeted *pax7* gene (*a1*, *a2* or *b*) at the region indicated in the schematic in Figure 6a. Values are means $\pm$ SE, n = 3 (mean of 2 replications). They were analyzed by

one-way ANOVA followed by Tukey's post-hoc test for multiple comparisons ($P < 0.05$). Different letters represent significantly different values. nd = no difference detected.

Table 1

Sequences of primer pairs used for real-time quantitative RT-PCR

Target	Forward primer (5'–3')	Reverse primer (5'–3')	Accession # ^a
<i>pax7a1</i>	TGGGACTACGATTTATTGTCTCC	TCGTGCAAAGTCCAGACAAG	GSONMG00081386001
<i>pax7a2</i>	TGGGACTACGATTATAGTTCGATT	TTCTTACTCGCGCAAAGTCC	GSONMG00061433001
<i>pax7b</i>	CGTCAAAACAATTACCACAAACA	AAAGACGACTGCATTCTACAGC	GSONMG00027288001
<i>myoD1</i>	CTGGACCCGAGACTCGTTCA	TTCCTGCGATCAGCATTGGT	X75798
<i>PCNA</i>	CCACGTCTCCCTAGTTCAGC	TCCTGGTTGAGTGTCTCGAA	BT046966
<i>myogenin</i>	CACTACATTG AGAGGCTGCAGGCA	CTCACTCGACGACGACACGCTG	Z46912
<i>EF1α</i>	TCCTCTTGGTCTGTTTCGCTG	ACCCGAGGGACATCCTGTG	AF498320.1

^a Accession numbers are from www.genoscope.cns.fr/trout and <http://www.ncbi.nlm.nih.gov/>

Table 2

Sequences of primer pairs used for ChIP analysts

Target	Forward primer (5'–3')	Reverse primer (5'–3')	Location ^a
<i>myogenin gene^b</i>			
FAR	CTCCTGCTTGTCATTGAGGA	TGGTTGGACATTTTGACAGG	–2749/–2587 ^c
DIS	CTTGAGGGATTAGCGACAGG	GGACACCCCTGAGTGACTGT	–1161/–989
PRO	TCTCTGGGCTCGTCTAAAT	GAAGTTTAGGCGGCGAAGAC	–258/–13
<i>pax7 genes^b</i>			
Pax7a1	AGAGCAAAGAAGGAGTCGTG	TGTCGACCAATCAGACAACC	–652/–496
Pax7a2	TTTTTCCCAGGTACAAGGA	TAGGACGTAAAGCACACATTTC	–633/–519
Pax7b	TGAGTTTACTTCTACGTCCAAAACA	GCATGCGACGTAAAACACAT	–684/–532

^aThe values indicate the location on the gene where each primer set localizes from the ATQ start codon.