Photoperiod Influences Growth and mll (Mixed-Lineage Leukaemia) Expression in Atlantic Cod

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Abstract

Photoperiod is associated to phenotypic plasticity of somatic growth in several teleost species. However, the molecular mechanisms underlying this phenomenon are currently unknown but it is likely that epigenetic regulation by methyltransferases is involved. The MLL (mixed-lineage leukaemia) family comprises histone methyltransferases that play a critical role in regulating gene expression during early development in mammals. So far, these genes have received scant attention in teleost fish. In the present study, the mean weight of Atlantic cod juveniles reared under continuous illumination was found to be 13% greater than those kept under natural photoperiod conditions for 120 days. We newly determined cDNA sequences of five mll (mll1, mll2, mll3a, mll4b and mll5) and two setd1 (setd1a and setd1b) paralogues from Atlantic cod. Phylogenetic analysis revealed that the cod genes clustered within the appropriate mll clade and comparative mapping of mll paralogues showed that these genes lie within a region of conserved synteny among teleosts. All mll and setd1 genes were highly expressed in gonads and fast muscle of adult cod, albeit at different levels, and they were differentially regulated with photoperiod in muscle of juvenile fish. Following only one day of exposure to constant light, mll1, mll4b and setd1a were up to 57% lower in these fish compared to the natural photoperiod group. In addition, mRNA expression of myogenic regulatory factors (myog and myf-5) and pax7 in fast muscle was also affected by different photoperiod conditions. Notably, myog was significantly elevated in the continuous illumination group throughout the time course of the experiment. The absence of a day/night cycle is associated with a generalised decrease in mll expression concomitant with an increase in myog transcript levels in fast muscle of Atlantic cod, which may be involved in the observed epigenetic regulation of growth by photoperiod in this species.

Introduction

Histone modifications, including acetylation, methylation, phosphorylation, and ubiquitination, have emerged as key mechanisms of transcriptional regulation and may serve as an epigenetic regulation marking system that is responsible for maintaining heritable programs of gene expression during development [1,2]. In particular, histone methylation plays a critical role in gene expression and epigenetic regulation [3,4]. Mixed-lineage leukaemias (MLLs) are histone methyltransferases (HMTs) that specifically methylate histone H3 at lysine 4 (H3K4) and are linked to gene activation [5,6,7]. In yeast, Set1 exists as a multi-protein complex (known as COMPASS), which is the only H3K4-specific HMT [8,9]. In contrast, the human genome encodes seven Set1 homologues: MLL1 [10], MLL2 [11], MLL3 [12], MLL4 [13], MLL5 [14], SETD1A and SETD1B [15]. Each of these protein acts as a multi-protein complex sharing several common subunits [2].

MLLs are widely expressed during development and in most adult tissues, including myeloid and lymphoid cells [16]. Moreover, they are well known as master regulators of homeobox-containing (Hox) genes that are critical for cell differentiation and development [6,17,18]. Heterozygous Mll1-knockout mice show posterior shifts in Hox gene expression, and homozygous Mll1-knockout mice are embryonic lethals in which the patterns of Hox expression initiate normally but are not maintained past embryonic day 9.5 [19]. The involvement of MLL2 in mammalian myogenesis has been demonstrated by McKinnell et al. [20], who reported that an HMT complex containing MLL2 interacted with paired box protein 7 (Pax7) to directly regulate the expression of myogenic factors, particularly myogenic factor 5 (Myf-5). MLL2 also regulates the cell cycle in cultured myoblasts and is required for the expression of transcription factors that regulate the myogenic programme, including Myf-5 and myogenin (MyoG) [21]. The full repertoire of mll paralogues has never been determined in fish species so far and the few reports available are restricted to model fish species such as zebrafish, (Danio rerio) [5,22], except for a single mll gene that was cloned in tiger pufferfish (Takifugu rubripes) [23].

Atlantic cod (Gadus morhua) is one of the most economically important fish species worldwide. Nevertheless, the profitability of the cod farming industry is severely restricted by precocious sexual maturation of the fish in captivity, which reach puberty prior to attaining commercial size [24]. Photoperiod manipulation, typified by continuous illumination, has been successfully used to delay sexual maturation to some extent in Atlantic cod [24], similarly to
what has been observed in other farmed fishes, including Atlantic
salmon (Salmo salar) [25] and European sea bass (Dicentrarchus labrax)
[26]. While most studies involving photoperiod manipulation in
Atlantic cod have been conducted on two-year old fish [24], it has
been reported that short-term photoperiod manipulation during
early juvenile stages has a significant positive effect on somatic
growth, which is dependent on genetic background and environ-
mental temperature [27]. Remarkably, juvenile cod kept under
continuous light for three months had a significantly higher weight
than the simulated natural photoperiod group and this difference
remained even after 30 months of sea-pen rearing under identical
ambient conditions until harvest. By this point, fish subjected to the
initial continuous light treatment were up to 9% larger than their
counterparts reared under natural photoperiod [28]. The present
study was designed to further our limited understanding about the
epigenetic regulation of somatic growth in Atlantic cod, with
particular focus on the mll family, since these genes are known to play
a crucial role in myogenesis. We have cloned all representatives of
the mll gene family in Atlantic cod and examined their expression
levels in fast muscle of juvenile fish kept under continuous illumination or simulated natural photoperiod.

Results

Influence of photoperiod on growth performance

The initial mean weight did not differ between fish in the
continuous light and natural photoperiod groups (Fig 1, P>0.05, n = 123). Significant differences in mean weight between them
were only observed at days 120 and 180 (P<0.0001 and
P<0.0002, respectively, n = 123). Fish kept under continuous
illumination or simulated natural photoperiod groups (Fig 1, P
0.0001, respectively). Constant light was also

The mll gene family in Atlantic cod

Using degenerate PCR primers, we have successfully obtained
partial cDNA sequences for five mll and two setd1 paralogues in
Atlantic cod: mll1 (GU441836), mll2 (GU441837), mll3a
(GU441838), mll4b (GU441839), mll5 (GU441840), setd1a
(GU441841) and setd1ba (HQ315825) (Table 1). Comparative
mapping of genes surrounding each mll and setd1 parologue

Tissue distribution of mll paralogues

With few exceptions, mll paralogues were ubiquitously expressed
in all tissues examined, albeit at different levels (Fig 4). Mll3a
transcripts were abundant in all tissues, except gas bladder, whereas mll4b was present in smaller amounts in brain, heart and
head kidney. Mll1, mll2, mll5, setd1a and setd1b paralogues were
expressed at lower levels in the digestive tract (stomach and mid
gut) and gas bladder. It is noteworthy that all seven mll and setd1 paralogues were highly expressed in testes, ovaries, blood and fast
skeletal muscle of adult cod.

Differential expression of mll and key myogenic genes with photoperiod manipulation

Relative mll expression in fast muscle of juvenile cod subjected
to different photoperiod conditions was determined by qPCR,
using the geometric average of arp and ubi reference genes to
normalise the data. In general, all mll and setd1 genes were
significantly down-regulated at most time points in fast muscle
of fish from the continuous light group (P<0.05, Fig 5). Mll1, mll2,
mll5b and setd1a expression were significantly repressed in the
continuous illumination group throughout the time course of the
experiment until 120 days (P<0.05). Constant light was also

Figure 1. Growth history of Atlantic cod juveniles reared under
continuous light (LD 24:0, red bars) or natural photoperiod
conditions (LDN, blue bars) for six months. Details of the light
regimes are shown by red diamonds or green triangles for LD 24:0
and LDN groups, respectively. Sea water temperature is also indicated by blue circles. Significant differences in mean weight between the two
light groups at a particular time point (two-tailed t-test, n = 123) are
highlighted by an asterisk.
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significantly associated with a decrease in mll3a, mll5 and setd1ba expression only at some time points from one to 60 days (P<0.05). Remarkably, there was a rapid change in mll1, mll2, mll4b, mll5 and setd1ba expression with light regime, since their transcript levels were significantly lower in the continuous illumination group just 12 hours through the experiment. At day one, transcript levels

Table 1. Gene name, GenBank accession number, primer sequences (5’ to 3’), amplicon sizes (bp) and PCR efficiency (%) of mll, setd1 and myogenic genes cloned in Atlantic cod.

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<th>Name</th>
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<th>qPCR</th>
<th>Size</th>
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Reference genes used are also indicated. doi:10.1371/journal.pone.0036908.t001

Figure 2. Partial synteny map of the genomic region surrounding mll2. Synteny was disrupted between teleosts and tetrapods. Orthologous genes in Gadus morhua, Oryzias latipes, Gasterosteus aculeatus, Takifugu rubripes, Tetraodon nigroviridis and Danio rerio are colour coded and represented by block arrows that show their orientation in the genome. mll2 paralogues are indicated by the arrow. Additional synteny results for other mll paralogues can be found in Figures S1–9. doi:10.1371/journal.pone.0036908.g002
of mll1, mll3a, mll4b and setd1a in fast muscle of fish from the continuous light group were reduced between 42 and 57% compared to the natural photoperiod group (Fig 5). Mll expression differences faded after 180 days, as expected since the light regime was identical for both groups (Figs 1 and 5). In addition, relative expression of myogenic regulatory factors (MRFs: myog and myf5) and pax7 in fast muscle was examined in relation to photoperiod (Fig 5). Constant illumination was generally significantly associated with an increase in myf5 and pax7 expression, and the difference amongst light groups was significant at 12 hours (P<0.05). Myog transcript levels were significantly elevated with continuous illumination compared to the natural photoperiod group throughout the time course of the experiment from 12 hours until 120 days (P<0.05).

Discussion

In the present study we have cloned mll1, mll2, mll3a, mll4b, mll5, setd1a and setd1ba orthologues in Atlantic cod and correlated their expression with differences in growth between fish reared under two photoperiod regimes. At days 120 and 180, age 1 Atlantic cod juveniles kept under continuous illumination were 13.3% and 10.5% larger than the ones from the natural photoperiod group, respectively. A similar effect has been previously shown in Atlantic cod juveniles [28,29] but a recent report described a negative influence of photoperiod on growth rate [30]. This apparent discrepancy is most likely due to differences in the fish genetic background. There is a significant interaction between genotype and the response to photoperiod treatment. For example, specific growth rates of cod juveniles with the haemoglobin genotype Hb-I(2/2) increase from 1.8% day$^{-1}$ under natural photoperiod to 2.3% day$^{-1}$ under constant illumination at 13°C, whereas the

Figure 3. Phylogenetic tree of the seven mll and setd1 paralogues found in vertebrates. Numbers at the nodes indicate posterior probability and approximate likelihood-ratio values obtained from Bayesian (left) and maximum likelihood (right) methods, respectively. Species abbreviations are as follows: Bt, Bos Taurus; Ce, Caenorhabditis elegans; Cl, Canis lupus familiaris; Dr, Danio rerio; Gg, Gallus gallus; Gm, Gadus morhua; Hs, Homo sapiens; Mm, Mus musculus; On, Oreochromis niloticus; Pt, Pan troglodytes; Rn, Rattus norvegicus; Xt, Xenopus tropicalis. GenBank accession numbers for mll sequences are listed in Table S1.

doi:10.1371/journal.pone.0036908.g003
average specific growth rate of $H_b-I(1/1)$ fish remains almost unchanged with light regime [27]. It is not entirely clear how light stimulates somatic growth but it is not due to a simple extension of foraging activity and corresponding feed intake. In fact, in their natural environment age 1 Atlantic cod, like the ones used in our study, preysed preferentially on benthos at night time [31]. Moreover, our experimental fish were fed equal amounts daily and there were no apparent differences in feed consumption between the two light groups. It is likely that the short-term photoperiod treatment induces metabolic changes that promote higher growth rates, probably due to more efficient nutrient utilisation. Day length in Bodo, Norway, had reached 24 hours by 120 days into the experiment and, therefore, all fish were kept under identical conditions from this point onwards. Nevertheless, there was still an average 10.5% weight difference between the natural photoperiod and continuous illumination groups at 180 days. This indicates that short-term light manipulation may have a persistent effect on muscle growth and corroborates a previous report, which showed that juvenile cod reared under continuous light for three months and then transferred to sea pens became up to 9% larger than their counterparts initially kept under simulated natural photoperiod conditions [28].

Photoperiod has long been known as a factor affecting somatic growth in teleosts and it has been used in aquaculture to control growth and maturation of several commercial fish species [24,25,26]. However, the molecular mechanisms underlying growth plasticity induced by light are still unknown. Four basic helix-loop-helix (bHLH) transcription factors (myoD, myog, myf5 and myf6) known as MRFs have received considerable attention as key players involved in determination and differentiation of skeletal muscle. Recent evidence supports the existence of interactions between MRFs and chromatin modifying complexes, including HMTs [20,32]. Therefore we hypothesised that HMTs may be involved in this epigenetic regulation of growth in teleosts, since histone methylation is acknowledged as one of the most important systems to regulate chromatin status in mammals. In particular, MLL proteins are major H3K4-specific HMTs that regulate expression of Hox [18] and MRFs [21] during early development. MLL1 is known to catalyse H3K4 methylation of $HoxA7$, $HoxA9$ and $HoxC8$ [18,33] and heterozygous $Mll1$-knockout mice ($Mll1^{+/−}$) show impaired development due to insufficient Hox protein concentrations [19]. Microarray hybridisation studies have revealed that $Mll1$ affects the expression of 197 potential target genes in mice, namely cathepsin C and the CD34 stem cell antigen [34]. The role of HMTs, including MLLs, is still largely unclear in teleosts. To investigate their potential involvement in epigenetic regulation of muscle growth, we have cloned seven $mll$ and $setd1$ orthologues in Atlantic cod. Phylogenetic and synteny analyses revealed that unlike tetrapods most fish species contained two copies of $mll3$, $mll4$ and $setd1$. Their chromosomal localisation shows that these paralogues arose from the teleost-specific genome duplication prior to divergence of the teleost/tetrapod lineage [35]. Interestingly, $mll3a$ is further duplicated in green-spotted pufferfish.

Cod $mll$ paralogues had a broad tissue distribution in adult fish, albeit at various levels. Some $mll$ genes (e.g., $mll1$ and $mll5$) were highly expressed in blood, which might have biased the results observed in extensively vascularised tissues. $mll2$ and $mll3a$ tissue distributions were similar to their human counterparts [11,12]. In mice, $mll2$ is required for development and spermatogenesis, and conditional knock-out male mice lacking $mll2$ are infertile [36]. The high transcript levels of $mll2$ and all other $mll$ and $setd1$ orthologues in Atlantic cod gonads indicate that they may play an important role in gametogenesis.

Differential expression in fast muscle with photoperiod was observed to some extent in all cod $mll$ and $setd1$ paralogues. The influence of light on $mll$ transcript levels was noticeable as early as 12 hours following photoperiod manipulation, suggesting that $mll$ genes may be associated with physiological adaptation to light and perhaps even involved in circadian rhythmicity. The largest differences in $mll$ mRNA levels between photoperiod groups were detected at day one. By this point $mll1$, $mll3a$, $mll4b$ and $setd1a$ expression in fast muscle of cod from the continuous light group were reduced by up to 57% compared to the natural photoperiod group. $Mll2$ was found to be down-regulated with continuous illumination at various time points from 12 hours to 60 days. There are no published functional or expression studies of $mll2$ in teleosts but it is known to influence expression of key myogenic genes in mammals. In mice, overexpression of Pax7 in satellite cells is known to result in elevated levels of $Myf5$ expression [20]. The Wdr5-Ash2L-MLL2 HMT complex interacts directly with Pax7. This MLL2-Pax7 complex then binds to $Myf5$, resulting in H3K4 tri-methylation of surrounding chromatin [20]. Cod $pax7$ was found to be significantly up-regulated in fast muscle with continuous illumination at 12 hours, even though $mll2$ expression was slightly reduced. $Myf5$ expression in fast muscle, which might be induced by Pax7, was also elevated in the continuous light group at 12 hours and 30 days. Also, hysul oxidase-like 1 ($hox1l1$) is down-regulated 23-fold in $mll2$-knockdown HeLa cells [37]. LOX

Figure 4. Representative tissue distribution of $mll$ paralogues in adult Atlantic cod. cDNAs from various tissues (brain, gill, heart, head kidney, kidney, liver, spleen, stomach, mid gut, gas bladder, testis, ovary, fast skeletal muscle, skin and blood) were used for semi-quantitative RT-PCR. Actb and eef1a were used as endogenous references. Expression patterns were determined using three biological replicates. doi:10.1371/journal.pone.0036908.g004
proteins are involved in collagen cross-linking and, therefore, play an important role in the structural integrity of muscle fibres [38]. In mll5-knockdown mice myoblast cell lines, expression of key players in myogenesis such as Pax7, Myf5 and Myog is impaired and these cells have limited ability to differentiate [21]. It seems that mll5 controls the inappropriate expression of proliferation genes and maintains expression competence of key genes associated with myogenic differentiation in quiescent myoblasts.

Figure 5. Quantification of mll paralogues and key myogenic genes (myog, myf5 and pax7) in fast muscle of Atlantic cod juveniles reared under continuous light (red bars, LD 24:0) or natural photoperiod conditions (blue bars, LDN) for 6 months. In general, mll genes were differentially expressed between the two light groups and there was a decrease in mll transcript levels with continuous illumination as early as 12 hours, compared to the natural photoperiod group. Myog transcript levels were consistently higher in the constant light group compared to natural photoperiod. Asterisks * and ** indicate significant differences at p<0.05 and p<0.01, respectively (n=6). doi:10.1371/journal.pone.0036908.g005
Throughout the time course of the our trial, \textit{mll5} transcript levels were 20, 38, 40 and 31\% lower in the continuous light group at 12 hours, one, 30 and 60 days, respectively. Down-regulation of \textit{mll2} and \textit{mll5} in cod exposed to continuous illumination may result in a higher number of proliferating myoblasts, which would increase growth potential and explain at least in part the higher growth rate observed in these fish group compared to the natural photoperiod group. These results are consistent with the observed increase in transcript levels of \textit{pax7} and \textit{myf5} in fish kept under continuous illumination, since \textit{Pax7} is a known marker of myosatellite cells that is crucial for cell proliferation and My\textit{f5} is involved in commitment of myoblasts to the myogenic programme [20]. Moreover, \textit{myog} expression was consistently higher in the continuous light group compared to natural photoperiod throughout from 12 hours until 120 days. Myog plays a major role in myoblast differentiation and is known to be involved in thermally-induced phenotypic plasticity of muscle growth in fish [39].

We have characterized all representatives of the \textit{mll} gene family in Atlantic cod and found that continuous illumination led to growth enhancement, which was accompanied by an increase in \textit{pax7}, \textit{myf5} and \textit{myog} expression but associated with transcriptional repression of \textit{mll} and \textit{setd1} genes in fast muscle. To the best of our knowledge, this is the first study that investigated the molecular mechanisms of photic-induced plasticity of muscle growth in teleosts. MLL proteins are deemed global activators of multiple transcription factors and their reduced expression with light may be involved in epigenetic regulation of growth. For example, a decrease in activation of genes that inhibit myoblast differentiation into mature muscle fibres, such as \textit{myostatin}, may induce enhanced growth of cod juveniles reared under continuous illumination. In zebrafish, knock-down of \textit{myostatin-1} during embryonic somitogenesis results in up-regulation of muscle-specific transcription factors, including \textit{myog} [40]. During the last two months of our photoperiod manipulation experiment light conditions were identical for both fish groups but the growth effect persisted, even if not accompanied by differential \textit{mll} expression. Hence, epigenetic transcriptional memory may be due to chromatin remodelling that occurred during the first four months in response to photoperiod changes.

Materials and Methods

Photoperiod experiment and sample collection

Atlantic cod juveniles with an initial mass of 2.7±0.8 g (mean ± standard deviation [SD], \(n=123\)) were kept at Mørkvedbukta Research Station (University of Nordland, Norway) in two groups of three 250 m\(^2\) tanks at an initial density of 130 individuals per tank and acclimated under continuous light until the start of the treatment. Sea water was pumped from 200 m depth and supplied at 7.4±0.4°C (mean ± SD). A commercial diet (Amber Neptun, Skretting AS, Stavanger, Norway) corresponding to 5\% (w/w) of the fish body weight was provided daily by automatic belt feeders. Fluorescent white light tubes (Aura Light International AB, Karlskrona, Sweden) were used to illuminate the tanks evenly. Light intensity was monitored regularly with a Hanna Hai 97500 Luxmeter (Hanna Instruments, Kungsbacka, Sweden) and it was approximately 120 Lux near the water surface in the centre of the tanks. During the photoperiod experiment one group of three tanks was kept under continuous light whereas the other was kept under normal light regime that corresponded to natural environmental photoperiod conditions in Bodo (67°N), Norway from January until July 2010. \textit{Cápa} 120 fish from each group were weighed at the start of the experiment and then 0.5, 1, 7, 30, 60, 120 and 180 days thereafter. Statistical differences in mean weights were determined by Student’s t-test using GraphPad Prism (GraphPad software, San Diego, USA). At each time point, 9 fish were humanely killed by immersion in seawater containing 1 g·L\(^{-1}\) tricaine methanesulfonate (Sigma, Oslo, Norway). Fast muscle was carefully dissected below the second dorsal fin from these specimens, taking special care to avoid skin and red muscle, and samples were snap-frozen in liquid nitrogen and stored at −80°C until RNA extraction.

Two-year old Atlantic cod were maintained in land-based tanks at Mørkvedbukta Research Station. Six fish with 50.8±2.8 cm fork length and 1.52±0.33 kg body weight were humanely killed as above. Brain, blood, gill, gas bladder, heart, liver, head kidney, kidney, stomach, mid gut, spleen, testis, ovary, muscle and skin were collected, snap-frozen in liquid nitrogen and stored at −80°C for subsequent RNA extraction. All procedures were conducted in accordance to the guidelines set by the National Animal Research Authority (Forsoksdyrvalget, Norway) and approved by the Faculty of Biosciences (University of Nordland, Norway) ethics committee.

Cloning \textit{mll} genes in Atlantic cod

Total RNA was extracted from the above adult cod tissues and used to synthesise cDNA with the QuantiTect kit (Qiagen, Nydalen, Sweden), as reported [41]. To identify \textit{mll} paralogues in Atlantic cod, PCR amplification was performed with degenerate primer sets that were designed against the most conserved regions of each \textit{mll} fish orthologues (Table 1). PCR reactions were performed using the Expand High Fidelity PCR System (Roche, Mannheim, Germany) with the following thermocycling conditions: initial denaturation at 94°C for 3 min, 35 cycles of amplification for 30 s at 94°C, 20 s at 56°C and 30 s at 72°C, and a final elongation step of 72°C for 5 min. PCR products were separated by electrophoresis on a 1\% (w/v) agarose gel, and the cDNA fragments of the predicted molecular weight were extracted from the gel using the QIAquick Gel Extraction Kit (Qiagen). Purified amplicons were cloned and sequenced as detailed elsewhere [41].

Bioinformatic analyses

To ascertain the identity of the cod cDNA sequences obtained, BLASTX searches were performed against the NCBI database (ncbi.nlm.nih.gov). Moreover, \textit{in silico} cloning using the Atlantic cod genome draft (codgenome.no) was performed to obtain longer \textit{mll} and \textit{setd1} sequences for phylogenetic analysis. Putative deduced amino acid sequences were aligned with the corresponding orthologues in various species (Table S1) using MUSCLE (drive5.com). To eliminate gaps and divergent regions, heuristic alignment was trimmed with Gblocks 0.91b (molevol.eumc.c-sic.es). The resulting multiple sequence alignments were used for Bayesian (MrBayes v3.1.2, mrbayes.csit.fsu.edu) and likelihood (PhyML 3.0, www.atgc-montpellier.fr/phylm) phylogenetic analyses. Bayesian phylogenetic trees were obtained using a mixed model of amino acid substitution (1,000,000 generations, sampling every 10\(^{4}\) generation and burning the first 10,000 trees) and the likelihood analysis was performed using the LG substitution model with 4 substitution rate categories and an estimated γ shape parameter. Graphical representations of phylogenetic trees were obtained with PhyloWidget (phylowidget.org). Synonym analyses of all \textit{mll} and \textit{setd1} genes were performed on the Genomicus v64.01 genome browser (www.dyogen.ens.fr/genomicus-64.01).

Semi-quantitative PCR (RT-PCR)

cDNAs were synthesized from total RNA extracted from brain, blood, muscle, gill, head kidney, kidney, heart, liver, spleen,
stomach, intestine, immature testes (gonado-somatic index, GSI = 1.9%) and ovaries (GSI = 1.5%) of two-year old Atlantic cod. Semi-quantitative RT-PCR was conducted for each cod mll parologue using the respective qPCR primer sets indicated on Table 1. Actb and eef1a were used as internal controls. Thermocycling parameters were 94°C for 3 min, followed by 35 cycles of 30 s at 94°C, 30 s at 60°C and 30 s at 72°C, with by a final elongation step of 72°C for 3 min. PCR products were analysed by electrophoresis on a 1% (w/v) agarose gel, visualised and photographed on a Kodak gel documentation system v.4.0.5 (Oslo, Norway).

Quantitative real-time PCR (qPCR)

Total fast muscle RNA and cDNA was obtained as above from six fish from each of the two different photoperiod groups at the start of the light treatment and 0.5, 1, 7, 30, 60, 120 and 180 days thereafter. Target and reference genes were amplified using the primer sets indicated on Table 1. These primers were designed with the GenScript Real-time PCR software (www.genscript.com) across exon borders determined by Spidey (www.ncbi.nlm.nih.gov/spidey) to avoid amplification of contaminating genomic DNA [42]. Quantification of transcript levels was performed by qPCR using the LightCycler® 480 SYBR Green I Master chemistry (Roche) on a LightCycler® 480 (Roche), as previously described [41]. Fifty-fold diluted muscle cDNA samples were run in duplicate, and minus reverse transcriptase and no template controls were included in the reactions. The PCR reaction was performed at 95°C for 15 min, followed by 45 cycles of 15 s at 94°C, 20 s at 60°C and 20 s at 72°C. Five-point standard curves of a 2-fold dilution series were prepared from pooled RNA in order to calculate amplification efficiencies [42]. Cycle threshold (Ct) values were determined by the LightCycler® 480 software with a fluorescence level arbitrarily set to one. The suitability of β-actin (actb), acidic ribosomal protein (arsp), eukaryotic elongation factor 1α (eef1a) and ubiquitin (ubi) as reference genes for this experimental setup was investigated [43] and raw target gene data were corrected with geNorm normalisation factors (med experimental setup was investigated [43] and raw target gene data were corrected with geNorm normalisation factors (med experimental setup was investigated [43] and raw target gene data were corrected with geNorm normalisation factors (med experimental setup was investigated [43] and raw target gene data were corrected with geNorm normalisation factors (med experimental setup was investigated [43] and raw target gene data were corrected with geNorm normalisation factors 

Supporting Information

Figure S1 Partial synteny map of the genomic region surrounding mll1. Orthologous genes in Gadus morhua, Oryzias latipes, Gasterosteus aculeatus, Takifugu rubripes, Tetraodon nigroviridis and Danio rerio are colour coded and represented by block arrows that show their orientation in the genome. Mll1 paralogues are indicated by the arrow. (TIF)

Figure S2 Partial synteny map of the genomic region surrounding mll3a. Orthologous genes in Gadus morhua, Oryzias latipes, Gasterosteus aculeatus, Takifugu rubripes, Tetraodon nigroviridis and Danio rerio are colour coded and represented by block arrows that show their orientation in the genome. Mll3a paralogues are indicated by the arrow. (TIF)

Figure S3 Partial synteny map of the genomic region surrounding mll3b. Orthologous genes in Gadus morhua, Oryzias latipes, Gasterosteus aculeatus, Takifugu rubripes, Tetraodon nigroviridis and Danio rerio are colour coded and represented by block arrows that show their orientation in the genome. Mll3b paralogues are indicated by the arrow. (TIF)

Figure S4 Partial synteny map of the genomic region surrounding mll4a. Orthologous genes in Gadus morhua, Oryzias latipes, Gasterosteus aculeatus, Takifugu rubripes, Tetraodon nigroviridis and Danio rerio are colour coded and represented by block arrows that show their orientation in the genome. Mll4a paralogues are indicated by the arrow. (TIF)

Figure S5 Partial synteny map of the genomic region surrounding mll4b. Orthologous genes in Gadus morhua, Oryzias latipes, Gasterosteus aculeatus, Takifugu rubripes, Tetraodon nigroviridis and Danio rerio are colour coded and represented by block arrows that show their orientation in the genome. Mll4b paralogues are indicated by the arrow. (TIF)

Figure S6 Partial synteny map of the genomic region surrounding mll5. Orthologous genes in Gadus morhua, Oryzias latipes, Gasterosteus aculeatus, Takifugu rubripes, Tetraodon nigroviridis and Danio rerio are colour coded and represented by block arrows that show their orientation in the genome. Mll5 paralogues are indicated by the arrow. (TIF)

Figure S7 Partial synteny map of the genomic region surrounding setd1a. Orthologous genes in Gadus morhua, Oryzias latipes, Gasterosteus aculeatus, Takifugu rubripes, Tetraodon nigroviridis and Danio rerio are colour coded and represented by block arrows that show their orientation in the genome. Setd1a paralogues are indicated by the arrow. (TIF)

Figure S8 Partial synteny map of the genomic region surrounding setd1ba. Orthologous genes in Gadus morhua, Oryzias latipes, Gasterosteus aculeatus, Takifugu rubripes, Tetraodon nigroviridis and Danio rerio are colour coded and represented by block arrows that show their orientation in the genome. Setd1ba paralogues are indicated by the arrow. (TIF)

Figure S9 Partial synteny map of the genomic region surrounding setd1bb. Orthologous genes in Gadus morhua, Oryzias latipes, Gasterosteus aculeatus, Takifugu rubripes, Tetraodon nigroviridis and Danio rerio are colour coded and represented by block arrows that show their orientation in the genome. Setd1bb paralogues are indicated by the arrow. (TIF)

Table S1 GenBank accession numbers for five mll and two setd1 paralogues and corresponding proteins.

Table S2 Orthologues of mll and SET domain genes from yeast to human.

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Author Contributions

Conceived and designed the experiments: JF. Performed the experiments: KN AG. Analyzed the data: KN JF. Contributed reagents/materials/analysis tools: JF. Wrote the paper: KN JF.

References