Cellular Expression of *Midkine-a* and *Midkine-b* during Retinal Development and Photoreceptor Regeneration in Zebrafish

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**ABSTRACT**

In the retina of adult teleosts, stem cells are sustained in two specialized niches: the ciliary marginal zone (CMZ) and the microenvironment surrounding adult Müller glia. Recently, Müller glia were identified as the regenerative stem cells in the teleost retina. Secreted signaling molecules that regulate neuronal regeneration in the retina are largely unknown. In a microarray screen to discover such factors, we identified *midkine-b* (*mdkb*). Midkine is a highly conserved heparin-binding growth factor with numerous biological functions. The zebrafish genome encodes two distinct midkine genes: *mdka* and *mdkb*. Here we describe the cellular expression of *mdka* and *mdkb* during retinal development and the initial, proliferative phase of photoreceptor regeneration. The results show that in the embryonic and larval retina *mdka* and *mdkb* are expressed in stem cells, retinal progenitors, and neurons in distinct patterns that suggest different functions for the two molecules. Following the selective death of photoreceptors in the adult, *mdka* and *mdkb* are coexpressed in horizontal cells and proliferating Müller glia and their neurogenic progeny. These data reveal that Mdkb and Mdkb are signaling factors present in the retinal stem cell niches in both embryonic and mature retinas, and that their cellular expression is actively modulated during retinal development and regeneration. J. Comp. Neurol. 514: 1–10, 2009. © 2009 Wiley-Liss, Inc.

Indexing terms: *mdka; mdkb; growth factors; development; regeneration; neurogenesis; zebrafish*

Midkine is a secreted heparin-binding growth factor that is highly conserved throughout the animal kingdom (Kadomatsu et al., 1988, 1990; Winkler et al., 2003; Kadomatsu and Muramatsu, 2004). Numerous functions have been ascribed to this molecule: neurogenic, mitogenic, neurotrophic, chemotactic, fibrinolytic, and antiapoptotic (for review, see Muramatsu, 2002; Kadomatsu and Muramatsu, 2004). In mammals, Midkine is expressed in many tissues during embryonic development, most prominently in the developing neural tube and at epithelial-mesenchymal boundaries (Mitsiadis et al., 1995). In mammalian tissues injury elevates the expression of Midkine (Miyashiro et al., 1998; Obama et al., 1998; Kikuchi-Horie et al., 2004; Sakakima et al., 2006; Jochheim-Richter et al., 2006), and in zebrafish the expression of the “a” paralog of Midkine is increased in regenerating heart and fin (Lien et al., 2006; Schebesta et al., 2006). These results suggest a role for Midkine in regeneration and repair of vertebrate tissues (Ohta et al., 1999). Supporting this idea, mice deficient in Midkine show decreased regenerative capacity following partial hepatectomy (Ochiai et al., 2004), and in albino rats intravitreal injection of recombinant Midkine promotes survival of photoreceptors following light-induced injury (Unoki et al., 1994; Masuda et al., 1995).

The zebrafish genome contains two *midkine* genes, *mdka* and *mdkb*, encoding proteins that share 68% amino acid identity (Winkler et al., 2003). In early embryos the two midkines have distinct patterns of expression and biological actions (Winkler and Moon, 2001; Winkler et al., 2003; Schäfer et al., 2005; Liedtke and Winkler, 2008).

The zebrafish retina develops from cells of the anterior neural plate that form the optic cups, which by 24 hours postfertilization (hpf) are well developed and consist of proliferating neuroepithelial cells (Schmitt and Dowling, 1994; Additional Supporting Information may be found in the online version of this article.

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Neuronal differentiation begins at 28–32 hpf within a precocious patch, ventronasal to the optic stalk (Burrill and Easter, 1994; Schmitt and Dowling, 1994; Schmitt and Dowling, 1999; Hu and Easter, 1999). Retinal cell differentiation and lamina formation proceed at a fast pace in sequential waves that originate in the ventronasal patch and sweep dorsally and then temporally through the different layers. By 72 hpf, the retina is fully laminated and functional (Easter et al., 1996; Schmitt and Dowling, 1999; Hu and Easter, 1999).

After the initial differentiation of the retinal neuroepithelium, new neurons are added to the retina throughout the life of the animal. This neurogenesis persists in two regions, specialized niches that harbor stem cells and their immediate progeny: in the ciliary marginal zone (CMZ), at the border between differentiated retina and the iris, and in the differentiated retina, where resident stem cells give rise to a lineage of cells that exclusively generates rod photoreceptors (Hitchcock et al., 2004; Raymond et al., 2006). Within the CMZ, developmental time is spatially recapitulated, stem cells are located peripherally in the CMZ, adjacent to the iris, and progenitors with increasingly restricted competence are located more centrally. This spatial pattern is evidenced by a regionalized expression of genes that sequentially specify cellular identities (Raymond et al., 2006; see also Harris and Perron, 1998).

The teleost retina can regenerate photoreceptors and neurons in a process that generally recapitulates cellular and molecular events during late retinogenesis (Otteson and Hitchcock, 2003; Hitchcock et al., 2004; Raymond et al., 2006). Recent studies identified Müller glia as stem cells responsible for both persistent rod genesis (see above) and neuronal regeneration (Yurco and Cameron, 2005; Raymond et al., 2006; Fausett and Goldman, 2006; Bernardos et al., 2007; Kassen et al., 2007; Fimbel et al., 2007; Thummel et al., 2008).

Following the death of extant neurons, Müller glia de-differentiate, re-enter the cell cycle, and give rise to multipotent progenitors, which continue to proliferate, migrate, and differentiate to replace the missing neurons. Genes that guide developmental neurogenesis are re-expressed in regenerative Müller glia and their progeny (Raymond et al., 2006, and references therein). Little is known, however, about secreted signaling molecules regulating these regenerative events.

We identified mdkb in a screen for genes whose expression is regulated by the selective death and regeneration of photoreceptors. As a first step to understand the role of Midkines in the retina, we analyzed the cellular expression of mdkb and its paralog, mdka, during both retinal development and adult photoreceptor regeneration. During development, mdka is expressed in stem cells and progenitors, transiently expressed in developing Müller glia, and constitutively expressed in horizontal cells. In contrast, mdkb is expressed by newly postmitotic cells and constitutively expressed by retinal ganglion and amacrine cells. During regeneration, in addition to their constitutive patterns of expression, both midkines are expressed in horizontal cells and proliferating Müller glia and their neurogenic progeny. This study describes the expression of these two secreted factors in the developing, adult, and regenerating retina and establishes the foundation for future studies to investigate the function of these molecules.

**MATERIALS AND METHODS**

**Care of zebrafish and embryos**

Zebrafish (*Danio rerio*; 6–8 months old, 2–3 cm long) were purchased from suppliers and maintained in aquaria at 28.5°C with a 10/14-hour dark/light cycle. Three strains were used: albino, wildtype, and Tg(gfap:GFP)jin002001 (to identify Müller glia; a gift from Dr. Pamela Raymond). Embryos were generated by natural mating at light onset and reared at 28.5°C in embryonic rearing solution (Westerfield, 2000). Procedures for handling animals were approved by the University of Michigan’s Committee for the Use and Care of Animals.

**Light treatments**

Two light-lesion paradigms were used in this study. Each selectively kills photoreceptors, which induces proliferation and photoreceptor regeneration that follows a common time-course. First, for the gene chip analysis, *albino* zebrafish (University of Oregon, Eugene, OR) were treated as described previously (Vihtelic and Hyde, 2000; Vihtelic et al., 2006). Second, to induce photoreceptor death in pigmented fish, wild-type animals were dark adapted for 8 days and exposed to constant fluorescent light (28000–31000 lux). Control animals were sacrificed immediately following dark adaptation, and experimental animals were sacrificed at 24 hours, 48 hours, 72 hours, 5 days, 7 days, or 10 days after light onset. Additional animals were returned to normal lighting and sacrificed at 14, 21, or 28 days after light onset. This experiment was performed in triplicate.

**RNA isolation and microarray analysis**

Oligonucleotide microarrays were used to identify genes that were differentially expressed after light-induced photoreceptor death. For these experiments, retinas were sampled at a single timepoint, 72 hours after light onset, which is after the death of photoreceptors and at a time when the retina is replete with injury-induced photoreceptor progenitors (Vihtelic and Hyde, 2000). Control animals were maintained in normal light conditions. After animals were sacrificed, retinas were dissected free from the surrounding ocular structures and retinal RNA was isolated and processed in eight separate pools (four control, four experimental; 12 retinas each). From each pool of RNA, probes were synthesized and hybridized to a single chip. Briefly, total RNA was amplified to yield double-stranded antisense RNA (aRNA), which was biotinylated using the Affymetrix GeneChip Expression IVT Labeling Kit (Affymetrix, Santa Clara, CA). Ten micrograms of labeled aRNA were fragmented and hybridized to Zebrafish Genome Arrays (Affymetrix). Chips were scanned using the GeneChip Scanner 3000 (Affymetrix). The fluorescence intensity readouts were sorted into CHP files with the Affymetrix Microarray Suite v. 5.0. The array data discussed in this publication have been deposited in the National Center for Biotechnology Information Gene Expression Omnibus (Edgar et al., 2002) and are accessible through GEO Series accession number GSE13999 (http://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE13999).

A false-discovery rate-adjusted confidence interval (FDR-Cl) was used to identify statistically significant changes in fluorescence intensity as described (Hero et al., 2004; Benjamini and Yekutieli, 2005). This approach employs robust multiarray averaging (RMA) to normalize the data, assigns
fold-intensity differences, and utilizes a statistical method that provides an FDR-CI for each differentially labeled probe set (Hero et al., 2004; see also Yoshida et al., 2002). This analysis generated a rank-ordered list of probe sets that showed 2-fold or greater changes in fluorescence intensity on the chip images. Analysis of gene ontologies was performed using the Database for Annotation, Visualization, and Integrated Discovery (DAVID) (Dennis et al., 2003; http://david.abcc.ncifcrf.gov).

RNA isolation and quantitative RT-PCR

For quantitative reverse-transcriptase polymerase chain reaction (QRT-PCR), both albino and wildtype zebrafish were sacrificed and eyecups were removed and retinas dissected and carefully separated from the retinal pigment epithelium. RNA was isolated from control and light-lesioned albino zebrafish in experiments independent of those used for the microarray screen. Four retinas per group from 3–4 different zebrafish were pooled. Retinas were homogenized and RNA was isolated according to the manufacturer’s protocol (Ambion, Austin, TX). RNA was quantified with a spectrophotometer and RNA quality was assessed on ethidium bromide-stained agarose gels; 500 ng of total RNA was used to synthesize cDNA following the manufacturer’s protocol (Superscript II, Invitrogen, Carlsbad, CA). The resulting first-strand reaction was diluted 1:4 and used as a template for the subsequent QRT-PCR reaction (iQ SYBR Green Supermix, Bio-Rad, Hercules, CA) in the iCycler Real-Time PCR detection System (Bio-Rad). The following amplification and melt curve analysis protocol was used: 95°C 3 minutes, 40 x (95°C: 20 sec, 57°C: 20 sec, 72°C: 30 sec), 95°C: 1 minute, 90 x 55°C: 10 sec. The following primers were used: rhodopsin, forward: 5’ agccatcataaatcaccaca and reverse: 5’ agccttcatgctcgc-gatt; opsin-1, forward: 5’ aacaccagggagcagt and reverse: 5’ tgggtgtgcaacaagctg; proliferating cellular nuclear antigen (pcna), forward: 5’ catccagacactagctgaaga and reverse: 5’ ctgctgtggagcttgtt; connexin 52.6, forward: 5’ tgacagatcgtctctgcc and reverse: 5’ tgtgtcttgaaagccttg (Zoidl et. al. 2004); mdka, forward: 5’ tgaagttttgtctctgtgtggag and reverse: 5’ acgcagatcaatgactgcttg; mdkb, forward: 5’ gcctgttgagctgctgttg; reverse: 5’ catcacaatgctgctgttg; reverse: 18s RNA forward: 5’ tcgcctacccatcaaggaagagtac and reverse: 5’ tggctgcaattgacgctgtggc. The threshold cycle (Ct) was determined by the iCycler using the maximum curvature approach and then maintained constant for subsequent runs. Relative levels of gene expression were determined by the ∆∆Ct method using either connexin 52.6 or 18s RNA as an endogenous reference. Specificity of the amplification products was verified by agarose gel electrophoresis.

Tissue preparation, in situ hybridization, and immunohistochemistry

At selected developmental times, embryos and larvae were fixed by immersion in 4% paraformaldehyde in 0.1 M phosphate buffer, pH 7.2, cryoprotected in 20% sucrose, frozen, and cryosectioned. At set times following light treatment, adult fish were anesthetized in 0.05% 3-amino benzoic acid-ethyl ester (Sigma, St. Louis, MO), eyes were enucleated, lenses removed, and eyecups were processed identically to the embryos and larvae. Sense and antisense riboprobes for mdka or mdkb were synthesized from plasmids containing the full-length mdka and mdkb cDNAs (gift from Dr. Christoph Winkler [Winkler et al., 2003]; accession numbers NM_131070 [mdka] and NM_131716 [mdkb]). Plasmids were linearized and digoxigenin (DIG)-labeled riboprobes were generated by in vitro transcription using the DIG RNA Labeling kit (Roche Diagnostics, Indianapolis, IN). In situ hybridization was performed as described previously (Hitchcock and Kakuk-Atkins, 2004) using DIG-labeled probes and Fast Red (Roche Diagnostics) as the enzymatic substrate. The enzymatic reaction was monitored using a fluorescence microscope and stopped after 2–5 hours, when the signal was distinct and the background low. Fresh Fast Red staining solution was prepared and added to the slides each hour. The anti-DIG antibodies are Fab fragments isolated from sheep immunized against the whole digoxigenin molecule and conjugated with alkaline phosphatase (Hoffman-La Roche, Nutley, NJ; Cat. No. 11093274910). To combine in situ hybridization with immunohistochemistry, the Fast Red color reaction was first allowed to develop, then slides were processed immediately for immunohistochemistry using standard procedures.

Other commercially available antibodies were also used in this study. The antibody against Proliferating Cell Nuclear Antigen (PCNA; Sigma-Aldrich, St. Louis, MO) was used at a 1:1,000 dilution. This antibody was isolated from mouse immunized with recombinant rat PCNA protein and validated by its capacity to immunoprecipitate PCNA protein from whole cell extracts (Waseem and Lane, 1990). In our study, tissues stained with this antibody produce a pattern that is identical to that previously reported (e.g., Raymond et al., 2006). Antibodies against Green Fluorescent Protein (GFP; ab6656; Abcam, Cambridge, MA) were used at a 1:1,000 dilution. These antibodies were raised in rabbit against purified recombinant GFP made in Escherichia coli. In Western blots, this antibody recognizes a single 67-kDa band in lysate from whole cells expressing GFP-fusion protein. Again, tissues stained with this antibody produce a pattern that is identical to that previously reported (Bernardos et al., 2007). The Prox-1 antibodies were raised in rabbit against a synthetic peptide from the C-terminus of mouse Prox-1 (EIKFSPNCQELLLH), which in Western blots recognizes a single 83-kDa band, and used at 1:2,000 dilution (AB5475, Chemicon International, Temecula, CA). Zebrafish Prox-1 possesses 84% identity at the amino acid level with the mouse homologue (Glasgow and Tomarev, 1998), and this antibody has been shown to label Prox-1-expressing cells in zebrafish (Ober et. al., 2006). In addition, all sections were also stained with the fluorescent nuclear stain, 4,6-diamidino-2-phenylindole dihydrochloride (DAPI, Invitrogen-Molecular Probes, Eugene, OR). Prior to PCNA immunostaining, sections were processed for antigen retrieval as previously described (Raymond et al., 2006). Secondary antibodies, were goat, antirabbit or goat, antimouse were used at a dilution of 1:500 (Invitrogen-Molecular Probes).

Photographic images

Images were taken with a Nikon DMX 1200 digital camera mounted on a Nikon Eclipse E800 epifluorescent compound microscope. Adobe Photoshop CS2 (Adobe Systems, San Jose, CA) was used to construct the figures. The layer tool was used to generate image overlays, and the channel mixer tool was used to change the red signal to magenta. In some images, the clone stamp tool was used to remove unwanted scale bars. Images in Figure 2 and Supplemental Figure 1
were taken with an AxioCam mRM digital camera and a Zeiss Axio Imager epifluorescent compound microscope (Carl Zeiss Microimaging, Thornwood, NY). Images were false-colored using the Zeiss AxioVision 4.0 software and exported into Adobe Photoshop CS2 and treated as described above.

**RESULTS**

**Expression analyses**

The RMA/FDR-CI identified 671 experimental probe sets that showed a 2-fold or greater difference in fluorescence intensity (hereafter referred to as "gene expression") when compared to control probe sets. To characterize the global changes in gene expression, the differentially expressed genes were submitted to DAVID (Dennis et al., 2003) for functional annotation clustering. Table 1 lists gene ontology biological process terms that were statistically significantly overrepresented within this dataset. This analysis showed that many of the differentially expressed genes are associated with DNA replication and, very broadly, enzyme activity and cellular metabolism. Another ontology group in this collection, "visual perception," corresponds to genes encoding opsins. Functional categories were also identified from the gene ontology analysis, and after removing the entries lacking annotation, the number of genes that showed an increase or decrease in expression within each category was determined (Fig. 1A). Genes known to control the mitotic cell cycle, to regulate photoreceptor physiology, or to encode growth factors were separately evaluated. As anticipated, the expression of each of the 17 cell cycle control genes is increased, which reflects the accretion of mitotic photoreceptor progenitors, whereas the expression of each of the eight photoreceptor-specific genes is decreased, reflecting the selective death of rods and cones. Genes encoding seven differentially expressed growth factors (mida b [see below], granulin a, granulin 1, granulin 2, dickkopf 1, galectin 1-like 2, matrixmetalloproteinase 9, follistatin) were also identified. The direction of change in the expression for four of these genes was independently validated by in situ hybridization (data not shown; see Results). Finally, QRT-PCR was performed for genes encoding Rhodopsin, Red Opsin, PCNA, MdkA, and MdkB (Fig. 1B; Suppl. Fig. 1). In the light lesioned retinas, the levels of mRNA encoding rhodopsin and red opsin were reduced, whereas the levels of mRNA encoding pcna and mdkb were increased. The microarray analysis did not detect a change

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<th>Table 1. Gene Ontology (GO) Process Terms That Were Statistically Significantly Overrepresented among the Differentially Expressed Genes</th>
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<td>GO Biological Process Term</td>
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<td>Intracellular part</td>
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<td>Intracellular</td>
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<td>Enzyme regulator activity</td>
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<td>DNA-dependent DNA replication</td>
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<td>DNA replication initiation</td>
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<td>Protein polymerization</td>
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<td>Cysteine-type endopeptidase activity</td>
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<td>DNA replication</td>
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<td>Intracellular nonmembrane-bound organelle</td>
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<td>Nonmembrane-bound organelle</td>
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<tr>
<td>Chromosomal part</td>
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<td>Calcium-dependent phospholipid binding</td>
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<td>DNA metabolic process</td>
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<td>Enzyme inhibitor activity</td>
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<td>Chromatin</td>
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<td>Intracellular organelle</td>
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<td>DNA-dependent ATPase activity</td>
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<td>Nucleosome</td>
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<td>Cysteine-type peptidase activity</td>
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<td>Nucleotide-triphosphatase activity</td>
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<td>Microtubule-based movement</td>
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<td>Cytoskeleton-dependent intracellular transport</td>
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<td>Lipid binding</td>
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<td>Translation elongation factor activity</td>
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<td>GTPase activity</td>
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<td>Hydrolase activity</td>
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<td>Hydrolase activity, acting on acid anhydrides</td>
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<td>Sensory perception of light stimulus</td>
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<td>Visual perception</td>
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<td>ATPase activity</td>
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<td>Biogenic amine metabolic process</td>
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<td>Nucleosome assembly</td>
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The cellular expression of \textit{mdka} and \textit{mdkb} during retinal development. \textit{a–e}: In situ hybridizations that illustrate the retinal expression of \textit{mdka} at 30–120 hpf, respectively. The white arrowheads identify \textit{mdka}-expressing cells in the circumferential marginal zone and retina. The white arrows in panels \textit{c–e} identify the columnar cells within the inner nuclear layer that express \textit{mdka}. The yellow arrowhead in panel \textit{d} identifies the optic nerve. The yellow arrowheads in panel \textit{e} identify presumptive horizontal cells. \textit{f–j}: In situ hybridizations that illustrate the expression of \textit{mdkb} at 30–120 hpf, respectively. The arrow in panel \textit{g} identifies the ventralonal patch. The arrows in panels \textit{g–j} demarcate regions of transient \textit{mdkb} expression. The arrowheads in panels \textit{h} and \textit{i} identify the circumferential marginal zone, which does not express \textit{mdkb}. Note the prominent expression of \textit{mdkb} within the gcl and inl in panels \textit{h–j}. \textit{onl}, outer nuclear layer; \textit{inl}, inner nuclear layer; \textit{ipl}, inner plexiform layer; \textit{gcl}, ganglion cell layer. Scale bar = 50 \textmu m.

Figure 2.

in the expression of \textit{mdka}, whereas the QRT-PCR revealed a modest decrease in the level of mRNA encoding this protein.

\textit{mdka} and \textit{mdkb} are expressed in distinct cells during retinal development

To determine the cellular pattern of expression of \textit{mdka} and \textit{mdkb} in the developing retina, in situ hybridization was performed on tissue sections from embryos and larvae between 24 and 120 hpf (Fig. 2). Sense probes did not show any specific hybridization (Suppl. Fig. 2). Neither \textit{mdka} nor \textit{mdkb} transcripts are detected in the eye at 24 hpf (data not shown), consistent with previous observations (Winkler and Moon, 2001; Winkler et al., 2004). However, retinal expression of \textit{mdka} is detected at 30 hpf. At this time, a low level of expression is present throughout the retinal neuroepithelium and more intense expression is observed at the retinal margin, presaging the site of the CMZ (Fig. 2a). At 48 hpf, \textit{mdka} is expressed at the retinal margin and more broadly in the inner nuclear layer (INL) (Fig. 2b), but \textit{mdkb} is not expressed in the ventral and central retina where differentiated cells and laminae are present. Between 48 hpf and 72 hpf, \textit{mdka} expression becomes progressively restricted to the retinal margin. At 72 hpf, \textit{mdka} expression appears centrally in columnar cells spanning the INL with morphology suggestive of Müller glia (Fig. 2c). \textit{mdka} expression in these radially oriented cells persists through 120 hpf (Fig. 2c–e) when \textit{mdka} transcription begins in presumptive horizontal cells (Fig. 2e, see below). Interestingly, in addition to this dynamic pattern of expression in the neural retina, \textit{mdka} is strongly expressed in cells within the nascent optic nerve (Fig. 2d) and the lens epithelium.

The spatial pattern of expression for \textit{mdkb} is distinctly different from \textit{mdka} and suggests that this gene is transcribed in newly postmitotic cells that are integrated into the inner retinal layers. \textit{mdkb} is first expressed at 48 hpf in a ventronasal patch and more broadly in the laminated central retina (Fig. 2g). At 72 hpf, \textit{mdkb} is expressed in differentiated ganglion and amacrine cells, which straddle the inner plexiform layer (IPL) (Fig. 2h) and in a broad annulus of cells central to the CMZ (Fig. 2h). Between 96 hpf and 120 hpf, expression persists within the inner nuclear and ganglion cell layers, whereas the annulus of \textit{mdkb} expression becomes progressively restricted to cells just central to the CMZ (Fig. 2h–j). In contrast to \textit{mdka}, \textit{mdkb} is not expressed by retinal progenitors in the CMZ (Fig. 2h,j). \textit{mdkb} appears not to be expressed in cells destined for the outer nuclear layer (ONL). These data suggest that \textit{mdkb} is expressed by inner retinal cells as they differentiate and constitutively expressed by mature ganglion and amacrine cells. This pattern of expression is maintained in the adult (see below).

To analyze the expression of \textit{mdka} and \textit{mdkb} relative to proliferating cells, we combined in situ hybridization with immunohistochemistry for PCNA, a cofactor of DNA polymerases that is expressed during the late G1, S, and early G2 phases of the cell cycle (Kurki et al., 1986; Moldovan et al., 2007). At 72 hpf, when both mature and proliferating cells are present in the retina, \textit{mdka} is expressed in PCNA-positive cells within the CMZ (Fig. 3a–c). In contrast, \textit{mdkb} is coexpressed with PCNA in only a few cells at the interface between the CMZ and the differentiated retina (Fig. 3d–f).

Together, these data show that, during early retinal development, \textit{mdka} is expressed in mitotic retinal progenitors, whereas \textit{mdkb} is expressed in newly postmitotic cells. In addition, \textit{mdka} is transiently expressed in presumptive Müller glia (see next section). Finally, both genes are constitutively expressed in subsets of neurons within the inner retinal layers.

\textit{mdka} is transiently expressed in Müller glia and constitutively expressed in horizontal cells

During larval development, the \textit{mdka}-expressing cells in the INL have centrally located large nuclei with \textit{mdka} mRNA in radially extending processes (Figs. 2c–e, 3a, 4a). This morphology suggests \textit{mdka} is expressed by Müller glia. To test
During photoreceptor regeneration, both midkine genes are expressed by proliferating Müller glia and photoreceptor progenitors

To investigate the expression of the midkine genes during photoreceptor regeneration, in situ hybridization was used to compare the cellular patterns of mdka and mdkb expression in control and experimental retinas. The patterns of midkine expression in adult, control retinas are similar to that observed at 120 hpf: mdka is expressed in horizontal cells (Figs. 4m, 5a, and Suppl. Fig. 2), whereas mdkb is expressed by cells in the vitreal aspect of the INL and the GCL (Fig. 5g, Suppl. Fig. 2). In the adult, unlesioned retina, neither probe labels dividing cells (Suppl. Fig. 3). Housing wildtype, pigmented fish in constant bright light results in the thinning of the ONL, due to the death of photoreceptors, and the corresponding decrease in mRNA encoding opsin genes. The anatomical loss of photoreceptors is fully restored after fish are returned to a normal lighting environment, although mRNA levels do not return to normal levels (Suppl. Fig. 1). Seventy-two hours after light onset corresponds to the peak of injury-induced proliferation and the peak of PCNA expression (Vihitelic and Hyde, 2000; Suppl. Fig. 1), therefore, we sampled this timepoint to examine the relationship between the expression of mdka and mdkb and the injury-induced cell proliferation (Fig. 5). In the light-lesioned retinas, the expression of mdka persists in horizontal cells and is newly expressed by proliferating, PCNA-positive photoreceptor progenitors in both the inner and outer nuclear layers (Fig. 5b–d). In contrast to mdka, the domain of mdkb expression expands from the inner tier of (amacrine) cells within the INL to include all cells of the INL, including horizontal cells (Fig. 5h–j). In addition, mdkb is expressed in PCNA-positive photoreceptor progenitors. Light lesions using the adult Tg(gfap:GFP)M2001 zebrafish confirm that proliferating cells expressing mdka and mdkb are Müller glia and their mitotic progeny (data not shown). These altered patterns of gene expression begin to return to that of the controls by 10 days after light onset (Fig. 5e,f,k,l). Together, data from the in situ hybridizations show that photoreceptor death induces changes in the cellular pattern of mdka expression and a marked increase in the number of cells that express mdkb. Further, following photoreceptor death, and in contrast to that observed in developing and adult retinas, the expression of both midkines becomes coincident in horizontal cells and photoreceptor progenitors.

DISCUSSION

Light-induced death and regeneration of photoreceptors in zebrafish is a well-described phenomenon (Vihitelic and Hyde, 2000). Photoreceptors die between 24 and 48 hours after light onset. Overlapping with the death of photoreceptors, injury-induced proliferation commences at about 24 hours and continues for the next several days, enlarging a pool of photoreceptor progenitors. From this pool of mitotic cells, the first regenerated photoreceptors begin to differentiate at about 2000). Photoreceptors die between 24 and 48 hours after light onset. Overlapping with the death of photoreceptors, injury-induced proliferation commences at about 24 hours and continues for the next several days, enlarging a pool of photoreceptor progenitors. From this pool of mitotic cells, the first regenerated photoreceptors begin to differentiate at about 4 days postinjury (Bernardos et al., 2007). We used oligonucleotide microarrays to identify genes that are differentially expressed at 72 hours after light onset, near the peak of the proliferative phase. The light-lesion model was selected because, 1) it selectively kills photoreceptors without introducing ancillary injury to the eye; 2) we infer that degeneration of this speculation, we combined in situ hybridization with immunostaining for green fluorescent protein (GFP) on retinal sections from Tg(gfap:GFP)M2001 transgenic zebrafish (Bernardos and Raymond, 2006). These fish express GFP under the control of the glial fibrillary acidic protein (gfap) regulatory elements, which in the retina selectively marks Müller glia. The mdka-positive cells in the INL (Fig. 4a,c,d,f) overlay precisely with GFP-positive cells (Fig. 4b,c,e,f), confirming that mdka is transcribed in Müller glia.

To test our inference that mdka is expressed by horizontal cells, we combined in situ hybridization and antibodies against Prox-1. Prox-1, a homeodomain protein required for horizontal cell development (Dyer et al., 2003) and is expressed in chicks and mammals by horizontal, amacrine, and bipolar cells (Belecky-Adams et al., 1997; Dyer et al., 2003; Edqvist and Hallbook, 2004). At 72 hpf, Prox-1 labels the narrow band of horizontal cell bodies at the outer boundary of the INL, although mdka expression in these cells is not detected at this time (Figs. 2c, 4g–i). At 120 hpf, however, mdka expression colocalizes with Prox-1 immunostaining in the horizontal cells (Fig. 4j–l). Note that mdka expression is reduced or absent in the dorsal retina, which is less mature than ventral retina, showing that mdka is not expressed by newly differentiated horizontal cells. In the adult retina, mdka is constitutively expressed in Prox-1-positive horizontal cells (Fig. 4m–o).
Figure 4.  
*mdka* is expressed in Müller glia. a,d: In situ hybridizations of *mdka* expression in Tg(gfap:GFP)Z40X fish at 120 hpf. b,e: Müller glia immunostained with antibodies against green fluorescent protein. c: The digital overlay of panels a and b. f: The digital overlay of panels d and e. In each panel the three arrows in identify the same three Müller glia. In panel d the asterisks identify *mdka* expression in presumptive horizontal cells. g: An in situ hybridization showing the expression of *mdka* at 72 hpf. h: The same section as in panel g, but immunostained with antibodies against Prox1. i: The digital overlay of panels g and h. Note that at 72 hpf, horizontal cells synthesize Prox1, but do not yet express *mdka*. j: An in situ hybridization showing the expression of *mdka* at 120 hpf. k: The same section as in panel j, but immunostained with antibodies against Prox1. l: The digital overlay of panels j and k. Note the colocalization of *mdka* mRNA and Prox1 protein. m: An in situ hybridization showing the expression of *mdka* in the adult retina. n: The same section as in panel j, but immunostained with antibodies against Prox1. o: The digital overlay of panels d and e. Arrows in j–o identify horizontal cells that express *mdka* and are immunostained for Prox1, onl, outer nuclear layer; inl, inner nuclear layer; gcl, ganglion cell layer; DAPI, nuclear stain 4,6-diamidino-2-phenylindole dihydrochloride; ipl, inner plexiform layer; gcl, ganglion cell layer; ipl: inner plexiform layer. Scale bar = 50 μm.

Figure 5.  
In the light-lesioned retina, *mdka* and *mdkb* are expressed by horizontal cells and injury-induced photoreceptor progenitors. a: An in situ hybridization showing the expression of *mdka* in a control retina. The white arrow identifies *mdka*-expressing horizontal cells. b: An in situ hybridization showing *mdka* expression in a retina following 72 hours of light exposure. c: The same section as in panel b, immunostained with antibodies against PCNA. d: The digital overlay of panels b and c. Arrowheads and arrows in panels b–d identify double-labeled cells in the ONL and INL, respectively. e,f: *mdka* expression at 10 days after light onset and 18 days (28 days after light onset) after returning the fish to normal lighting, respectively. Note that by 10 days the cellular expression of *mdka* begins to resemble the controls, and PCNA-positive cells no longer express this gene. g: An in situ hybridization showing the expression of *mdkb* in a control retina. h: An in situ hybridization showing *mdkb* expression in a retina following 72 hours of light exposure. The arrowhead and arrow identify *mdkb*-expressing cells in the inner and outer nuclear layers, respectively. i: The same section as in panel f, immunostained with antibodies against PCNA. j: The digital overlay of panels h and i, in panels b–d and h–j, arrowheads and arrows identify double-labeled cells in the ONL and INL, respectively. ONL, outer nuclear layer; INL, inner nuclear layer; GCL, ganglion cell layer; PCNA, Proliferating Cellular Nuclear Antigen; DAPI: nuclear stain 4,6-diamidino-2-phenylindole, dihydrochloride. Scale bar = 50 μm.
only two cell types (rods and cones) will result in less complex changes in the retinal transcriptome; and 3) the selective death of photoreceptors, while sparing the other cell types, models aspects of photoreceptor degeneration in mouse genetic models and human disease. The specific goal of the array experiment was to identify growth factors expressed by the regenerative Müller glia and their progeny that we could hypothesize function to regulate aspects of photoreceptor regeneration. Several growth factors were identified in the array screen, and we chose to characterize the developmental and injury-induced expression of *mdkb*, which was shown to be significantly upregulated both in the microarrays and QRT-PCR analyses, and its paralog, *mdka*, which we show is also expressed in the retina and is upregulated in regenerating heart and fin (Lien et al., 2006; Schebesta et al., 2006).

During retinal development, the expression of the two midkines is differentially regulated, with each gene exhibiting distinct temporal and spatial patterns of cellular expression. During embryonic retinal development, midkines are expressed in adjacent, nonoverlapping domains, which, as retinal neurons differentiate, become restricted to annuli at the retinal margin. Double labeling with antibodies against PCNA showed that *mdka* is expressed by retinal progenitors, whereas *mdkb* is expressed in nearby, newly postmitotic cells. The adjacent cellular expression of these paralogous genes in the early retinal neuroepithelium is similar to their spatial patterns of expression in the neural tube. In early embryos, *mdka* and *mdkb* are expressed at the forebrain/midbrain and midbrain/hindbrain boundaries, where they form contiguous, nonoverlapping domains and function to pattern the neural tube (Winkler et al., 2003).

During larval retinal development, each midkine is expressed in a spatial pattern that persists into adulthood. *Mdka* is expressed by mitotic cells in the CMZ and horizontal cells in the INL. In the optic tectum of adult zebrafish, *mdka* transcripts are present in the subventricular zone (Winkler et al., 2003), a site of persistent neurogenesis (Marcus et al., 1999). In mammals, Midkine is expressed in neural progenitors, in both the normal and injured brain, and functions to promote proliferation (Kikuchi-Horie et al., 2004; Zou et al., 2006). Our finding that *mdka* is expressed in the neurogenic CMZ of the zebrafish retina suggests that *mdka* is a component of neural stem cell compartments in vertebrates. The persistent expression of *mdka* in horizontal cells is intriguing. Horizontal cells are among the first retinal neurons to differentiate, and it has been proposed that they play a pioneering role in the development of photoreceptors (Messersmith and Redburn, 1990; Hagedorn et al., 1998). In contrast to *mdka*, *mdkb* is expressed in newly postmitotic cells destined for the inner retina. The initial expression of *mdkb* follows the wave of differentiation, and this pattern of expression is then recapitated among each generation of newly postmitotic cells as they exit the CMZ. *mdkb* is then also constitutively expressed by amacrine cells and cells in the ganglion cell layer.

Several extrinsic signaling molecules have been identified that regulate development of the teleost retina. IGFs and Wnts regulate proliferation of retinal progenitors, whereas sonic hedgehog, FGFs, retinoic acid, and Notch-mediated signaling regulate aspects of neuronal differentiation (Stenkamp, 2007, and citations therein; Vinokhukumar et al., 2008). The function of the Midkines in the teleost retina is not yet established; however, based on the tight spatial association of the two Midkines with proliferating and differentiating retinal neurons, respectively, it can be speculated that Mdka is a molecular component of the retinal stem cell niche (Raymond et al., 2006; see also Livesey et al., 2004) and may play a role in regulating proliferation, whereas Mdkb may play a role in aspects of neuronal differentiation.

During photoreceptor regeneration, both genes show altered patterns of cellular expression. The expression of *mdkb* expands from the inner tier of the INL to include all neurons of the INL. In addition, *mdka* is expressed in the injury-induced progenitors. The marked expansion in the number of cells within the INL that express *mdkb* can account for the upregulation of *mdkb* expression detected by the microarrays and QRT-PCR. In contrast, the expression of *mdka* showed no significant change in the microarrays, and QRT-PCR showed that *mdka* expression is slightly downregulated, even though the cellular expression of *mdka* in the injured retina is consistently altered. There is no obvious explanation for these seemingly contradictory results, although it should be noted that the circadian clock regulates *mdka* expression (Calinescu et al., in prep.), and this could influence either the microarray or QRT-PCR analyses. Nonetheless, the coincident expression of both midkines in horizontal cells and injury-induced photoreceptor progenitors, patterns of expression distinct from that observed in the developing retina, suggests that each gene product may have a common function during retinal injury and repair. In zebrafish, *mdka* expression is upregulated during regeneration of the fin and heart (Schebesta et al., 2006; Lien et al., 2006). The cellular expression of *mdka* in the regenerating fin has not yet been described. In the regenerating heart, *mdka* is expressed in cells that surround the wound site, but it is not expressed in the proliferative cardiomyocytes that replace the ventricular tissue (Lien et al., 2006). These results and the data reported here show that Midkine paralogs can be differentially regulated during the regeneration of specific tissues, but that both Midkines function as injury-induced growth factors. Further, both midkines may play an essential role in injured tissues with the capacity to regenerate to full anatomical integrity and functional recovery.

In summary, the expression of *mdka* and *mdkb* is actively and differentially regulated during retinal development and injury-induced photoreceptor regeneration. During development, the two midkine genes are expressed in distinct populations of cells—stem cells, retinal progenitors, and mature neurons, suggesting that these secreted molecules subserve different functions. Following retinal injury, the spatial pattern of midkine expression is altered and both genes are expressed in horizontal cells and Müller glia and their neurogenic offspring. This is the first description of the cellular expression of midkines in the vertebrate retina, and this study adds Mdka and Mdkb to the molecular signature of Müller glia that exhibit features of neural stem cells. Across the vertebrate phylum, Midkine is upregulated following injury, and in humans Midkine is integral to the growth of numerous carcinomas. An emerging concept in biology is that the same regulatory proteins control common cellular events during development, tissue repair, and carcinogenesis (Beachy et al., 2004; Gardiner, 2005). Our studies support this concept and add Mdka and Mdkb to the family of signaling molecules present in the developing and regenerating retina.
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