Research paper

Gentamicin is ototoxic to all hair cells in the fish lateral line system

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ABSTRACT

Hair cells of the lateral line system in fish may differ in their susceptibility to damage by aminoglycoside antibiotics. Gentamicin has been reported to damage hair cells within canal neuromasts, but not those within superficial neuromasts. This finding, based on SEM imaging, indicates a distinction in the physiology of hair cells between the two classes of neuromast. Studies concerned with the individual roles of canal and superficial neuromasts in behavior have taken advantage of this effect in an attempt to selectively disable canal neuromasts without affecting superficial neuromast function. Here we present an experimental test of the hypothesis that canal neuromasts are more vulnerable to gentamicin than superficial neuromasts. We measured the effect of gentamicin exposure on hair cells using vital stains (DASPEI and FM1-43) in the neuromasts of Mexican blind cave fish (*Astyanax fasciatus*) and zebrafish (*Danio rerio*). Contrary to the findings of prior studies that used SEM, gentamicin significantly reduced dye uptake by hair cells of both canal and superficial neuromasts in both species. Therefore, lateral line hair cells of both neuromast types are vulnerable to gentamicin ototoxicity. These findings argue for a re-evaluation of the results of studies that have used gentamicin to differentiate the roles of the two classes of neuromast in fish behavior.

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1. Introduction

Aminoglycoside antibiotics are ototoxic to hair cells within the inner ear and lateral line system of vertebrates (Wersäll, 1960; Kroese and van den Bercken, 1982; Forge and Schacht, 2000; Ernst et al., 1994). Because these antibiotics provide a cost-effective treatment for bacterial infections, biomedical researchers have investigated whether hair cells may be protected from this ototoxicity (Forge and Schacht, 2000). Comparative biologists have utilized aminoglycosides to experimentally ablate the lateral line system to examine its role in behavior (e.g. Montgomery et al., 1997; Coombs et al., 2001). It is therefore of interest to both biomedical and comparative research that that some hair cells in the fish lateral line system have been reported to be resistant to damage by the aminoglycoside gentamicin (Song et al., 1995).

Hair cells are found in two classes of lateral line receptor organs, canal neuromasts (CNs) and superficial neuromasts (SNs) (Fig. 1). In both classes, a gelatinous cupula covers the apical surface of the hair cells. The cupula transmits the motion of the surrounding fluid to the underlying hair cells to allow fish to sense water flow (Dijkgraaf, 1963). CNs are recessed within channels within the dermal layer and range in diameter from 100 to 500 µm with tens to hundreds of hair cells. SNs reside on the surface of the skin and are smaller (~50 µm in diameter) with around a dozen hair cells (Münz, 1989). The two classes are innervated separately and have distinct developmental origins (Webb and Shirey, 2003; Ghysen and Dambly-Chaudière, 2004; López-Schier and Hudspeth, 2005; Nuñez et al., 2009). In addition, the response properties of the two receptors differ, which creates two submodalities within the lateral line system. SNs generally are sensitive to low-frequency (<30 Hz) deflections that are representative of water velocity and CNs respond best to high-frequency deflections that are dependent on pressure gradients (Coombs and van Netten, 2006). Although these differences can largely be understood as a result of the mechanical filtering properties of the overlying cupula and canal, it remains possible that differences in the physiology of hair cells contribute to their distinct properties.

Song et al. (1995) found that the hair cells within CNs are damaged by exposure to gentamicin while those within SNs were not. Although the hair cells of both classes are similar in ultrastructure, scanning electron microscopy (SEM) revealed that exposure to gentamicin results in damage to the ciliary bundles of only the hair cells within canal neuromasts. This difference in susceptibility has
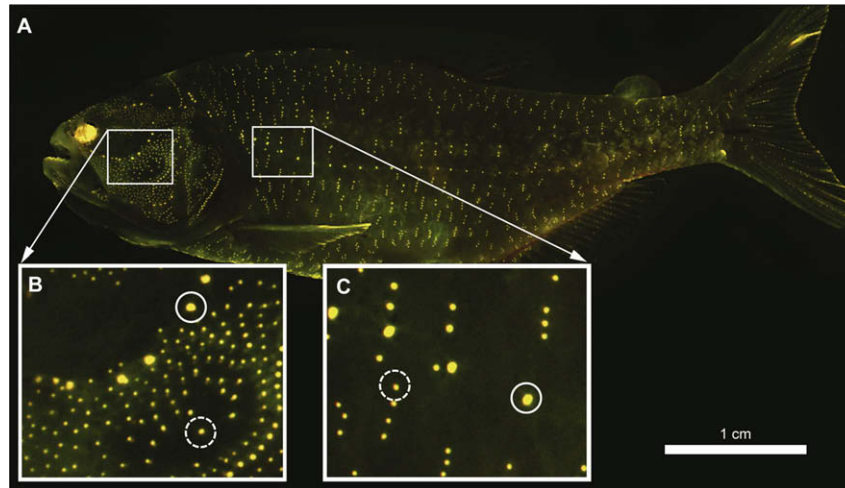
Fig. 1. The lateral line system of a Mexican blind cave fish stained with DASPEI under epifluorescent illumination. (A) The neuromasts over the surface of the body appear as discrete yellow circles. They are shown in detail for a region in the head in the infraorbital region (B) and trunk (C). (B–C) Canal neuromasts (solid circle) appear larger than superficial neuromasts (dashed circle) because they possess a greater number of hair cells.

been used to distinguish hair cell types in fish (Yan et al., 1991) and mammals (Lindeman, 1969). Thus, Song's result (later replicated by Montgomery et al., 1997; Baker and Montgomery, 1999a; and Coombs et al., 2001 in different species) provided the basis for the hypothesis that the hair cells within SNs are a physiologically distinct type from those within CNs. The goal of the present study was to use vital-staining techniques (Raible and Kruse, 2000; Meyers et al., 2003) to examine the effect of gentamicin on hair cell viability in CNs and SNs.

A number of styryl pyridinium dyes, like DASPEI (2-(4-(dimethylamino)styryl)-N-ethylpyridinium iodide) and FM1-43 ((n-(3-triethylammoniumpropyl)-4-(4-(dibutylamino)styryl) pyridinium dibromide), allow for the visualization of hair cells in vivo. FM1-43 enters hair cells by rapid entry through the mechanotransduction channels at the stereociliary tips of hair cells (reviewed in Meyers et al., 2003). When the mechanotransduction channels are blocked or otherwise compromised, this dye fails to permeate the cell; as such, dye uptake indicates functioning hair cells. Similarities in conformation and labeling behavior make it likely that DASPEI is also rapidly taken up through the mechanotransduction channels. FM1-43 has an additional advantage in that it will fluoresce much longer than DASPEI (e.g. over many days, Meyers et al., 2003). Thus, FM1-43 can be applied shortly before drug treatment and then visualized afterwards to determine if the hair cells survive the treatment.

In this study, the effects of gentamicin on the lateral line system were explored in zebrafish (Danio rerio) and the Mexican blind cave fish (Astyanax fasciatus). The zebrafish has emerged as a major model for studying the effects of aminoglycoside antibiotics on hair cells (Harris et al., 2003; Murakami et al., 2003; Santos et al., 2006; Owens et al., 2007; Chiu et al., 2008). The adults have little pigmentation in the head, which allows canal neuromasts to be visualized. The Mexican blind cave fish (also known as Astyanax mexicanus) completely lacks pigmentation, has thousands of large neuromasts, and is a common subject for studying the role of the lateral line system in behavior (e.g. von Campenhausen et al., 1981; Weissert and von Campenhausen, 1981; Tyke 1990; Baker and Montgomery, 1999a; Sharma et al., 2009). Additionally, cave fish have been used in studies of hair cell damage and recovery (Repass and Watson, 2001; Berg and Watson, 2002). Therefore, both of these species are ideal animal models for investigation of the effects of gentamicin on hair cells within the lateral line.

2. Materials and methods

2.1. Animals

Experiments were conducted independently at Bowling Green State University with Mexican blind cave fish and at the University of California at Irvine with zebrafish. Fish at both locations were maintained with standard techniques for these species. Mexican blind cave fish (standard length: 4–6.7 cm) were obtained from commercial aquarium suppliers and maintained in freshwater at 20–25 °C in darkness. Zebrafish (AB line, standard length: 2.7–3.5 cm) were raised in a colony in the lab of MJM, where they were maintained at 27–29 °C on a 14/10 h light/dark cycle. The Institutional Animal Care and Use Committees of both universities approved all protocols for experiments and fish care.

2.2. Gentamicin treatment

The effect of gentamicin on the number of neuromasts was assessed by a comparison between treated and control groups. Fish within both groups were handled to the same degree and were placed in a well-aerated aquarium for 24 h. During this period, the control group was exposed to standard tank water, whereas the water for the treatment group included a 0.001% concentration of gentamicin sulfate (Fisher Scientific, Pittsburgh, PA). This treatment was a lower concentration and a shorter exposure than in previous studies (Song et al., 1995 used 0.002% for 4 days) because we learned from a pilot study that a 0.002% gentamicin concentration prevented dye labeling in all neuromasts in both species within 24 h of exposure. Our 0.001% treatment assured that some neuromasts would still label, thus permitting a quantitative comparison between CNs and SNs and maximizing the potential for observing differential effects.

2.3. Vital dye labeling

Two vital stains were used to determine the presence of viable hair cells within neuromasts. Both stains were applied by exposing animals to a bath solution of dye under darkened conditions. Treatment and control fish of both species were stained with 0.008% DASPEI (Invitrogen Molecular Probes, Eugene, OR) solution for 10 min (Mexican blind cave fish) or 1 h (zebrafish) immediately following the 24 h gentamicin or sham exposure. In order to

determine if measured differences in DASPEI staining after genta-micin exposure resulted from damage to the integrity of the hair cell or a mere blockage of transduction channels, we stained addi-tional groups of control and treated zebrafish to 3 μM FM1-43 (Invitrogen Molecular Probes, Eugene, OR) 1 h prior to the genta-micin or sham treatment. This stain allowed us to measure the number of pretreatment cells that survived gentamicin exposure.

2.4. Imaging

All imaging was performed on adult fish anesthetized with 200 mg l^{-1} tricaine methanesulfonate (i.e. MS-222 or Finquel; Ar-gent Labs, Redmond, WA). Neuromasts were observed under epi-fluorescent illumination. Observations on Mexican blind cave fish were conducted on a stereomicroscope (Olympus SZX12, New Hyde Park, NY) with a GFP filter set (excitation 450–490 nm and barrier 515 nm) to visualize DASPEI. Photographs of these fish were captured with a CCD camera (Olympus Q Color-3 with Image Pro Software, V. 6). A similar microscope (Zeiss Discovery V.20 with GFP filter set and an AxioCam HRc camera, Carl Zeiss, Thorn-wood, NY) was used to visualize DASEPI in zebrafish. FM1-43 was made visible on this microscope using the same filter set. All obser-vations were made within 1.5 h of the completion of the 24 h treatment.

2.5. Counting neuromasts

Three methods for counting neuromasts were used due to dif-ferences in microscope optics, the degree of pigmentation in the two species, and the specificity of the two stains. Individual hair cells were most visible by DASPEI staining in zebrafish, which al-lowed us to differentiate brightly stained, presumably active hair cells, from weakly stained inactive hair cells and surrounding sup-port cells (as in Harris et al., 2003). A zebrafish neuromast stained with DASPEI was scored as "viable" if it possessed more than two brightly stained hair cells, "damaged" if it included one or two brightly stained hair cells, or "non-viable" if no brightly stained cells could be identified. Individual hair cells were not visible in DASPEI-stained cavefish and we consequently counted every neu-romast with visible labeling without differentiating the intensity of labeling. Therefore, neuromast counts for the blind cave fish in-cluded both weakly and strongly labeled neuromasts with varying numbers of viable hair cells. We similarly counted all identifiable neuromasts within a specified region in FM1-43 stained zebrafish because it was difficult to obtain reliable hair cell counts at 24 h following labeling. This was due to diffusion of the dye resulting in the labeling of nearby cells around the neuromasts. However, neuromasts were clearly identifiable under high magnification, as observed previously (Santos et al., 2006).

Neuromasts from all over the body were included in our counts. For DASPEI-stained Mexican blind cave fish, CNs were sampled from the supraorbital, infraorbital, post otic, preopercular, mandib-ular canals on the head and from the main trunk canal on the body (see Fig. 2). Superficial neuromasts were sampled from the ventral surface of the lower jaw (mandibular tip location) and along the rostral-most portion of the trunk canal from the fifth trunk canal scale through the tenth. SNs from scales dorsal and ventral to each of these trunk scales were also sampled. Because zebrafish possess far fewer neuromasts compared to blind cavefish, we were able to score all visible CNs over the entire body and all SNs on the trunk for both DASPEI and FM1-43 stained fish. On the head, samples of SNs from DASPEI-stained fish were limited to a randomly selected group of 50 because they were more numerous and distributed with greater variability in this region. In FM1-43 stained fish, sam-ples of SNs were restricted to the ventral section of the preopercu-lum, where there was a densely populated and easily identified line of SNs.

2.6. Statistical design

Our experimental design permitted us to test the effect of gen-tamicin on SNs and CNs. We used a two-factor ANOVA (imple-mented in Matlab v. 2007a, with the Statistics Toolbox) to test for significant differences in the quantity of labeled SNs and CNs (factor 1) in treatment and control groups (factor 2) (Sokal and Rohlf, 1995). Neuromast counts for specific body regions are re-ported, but body region was not included as a factor in the analysis because regional differences were not a concern for the present study. Thus, samples from different body regions were summed for each neuromast type and treatment group prior to statistical analysis. For DASPEI-stained zebrafish, only those neuromasts with brightly labeled hair cells (scored as 'viable' or 'damaged') were in-cluded in our ANOVA, whereas for DASPEI-stained blind cavefish and FM1-43 stained zebrafish, all neuromasts with any visible staining were included. In principle, this means that our criteria for gentamicin-induced effects were more conservative for the lat-ter two cases.

3. Results

Neuromasts labeled with DASPEI were highly visible through-out the body in untreated Mexican blind cave fish (Figs. 1 and 2A–E). Gentamicin exposure substantially reduced the degree of visible labeling of both CNs and SNs on all regions of the body (Fig. 2F–J). Close inspection of CNs revealed weak labeling at the margins of neuromasts, where non-sensory, support cells normally reside. Little fluorescence was observed in the center of neuro-masts, where hair cells are normally located (Fig. 2E vs. J). A similar response to gentamicin was apparent in zebrafish. The higher res-olution of our zebrafish images revealed that some brightly labeled hair cells were visible in treated zebrafish (Fig. 3). In zebrafish stained with FM1-43, labeled neuromasts showed no observable difference in brightness between control and treated fish, however there were far fewer remaining labeled neuromasts in treated indi-viduals (Fig. 4).

Both gentamicin-treated zebrafish and Mexican blind cave fish exhibited significantly fewer labeled neuromasts than control fish in both neuromast categories (Fig. 5). In DASPEI-stained cavefish, treated fish ($n = 5$) had significantly fewer visible neuromasts than control fish ($n = 6$) ($p < 0.001$, Table 1, Fig. 5A), even though weakly stained neuromasts were included in counts. In DASPEI-stained zebrafish, greater imaging resolution made it possible to separate individual hair cells allowing us to consider only neuromasts with brightly labeled hair cells. Consequently, a more significant differ-ence between treatment ($n = 8$) and control ($n = 8$) groups for both classes of neuromast ($p \ll 0.001$, Table 2, Fig. 5B) was observed. On average, none of 11 CNs and 1 of 159 SNs were scored as viable in zebrafish after the gentamicin treatment. The control group, by contrast, had neuromasts scored as viable in 15 out of 15 cases for CNs and in 194 out of 240 cases for SNs (Table 2). In addition, zebrafish stained with FM1-43 prior to treatment exhibited a sig-nificantly lower neuromast count ($p = 0.003$, Table 3, Fig. 5C) in the gentamicin treatment group ($n = 5$) relative to the control ($n = 5$).

4. Discussion

Our results demonstrate that gentamicin disrupts hair cell func-tion throughout the lateral line system and thereby refute the hypothesis that hair cells within SNs are resistant to gentamicin.
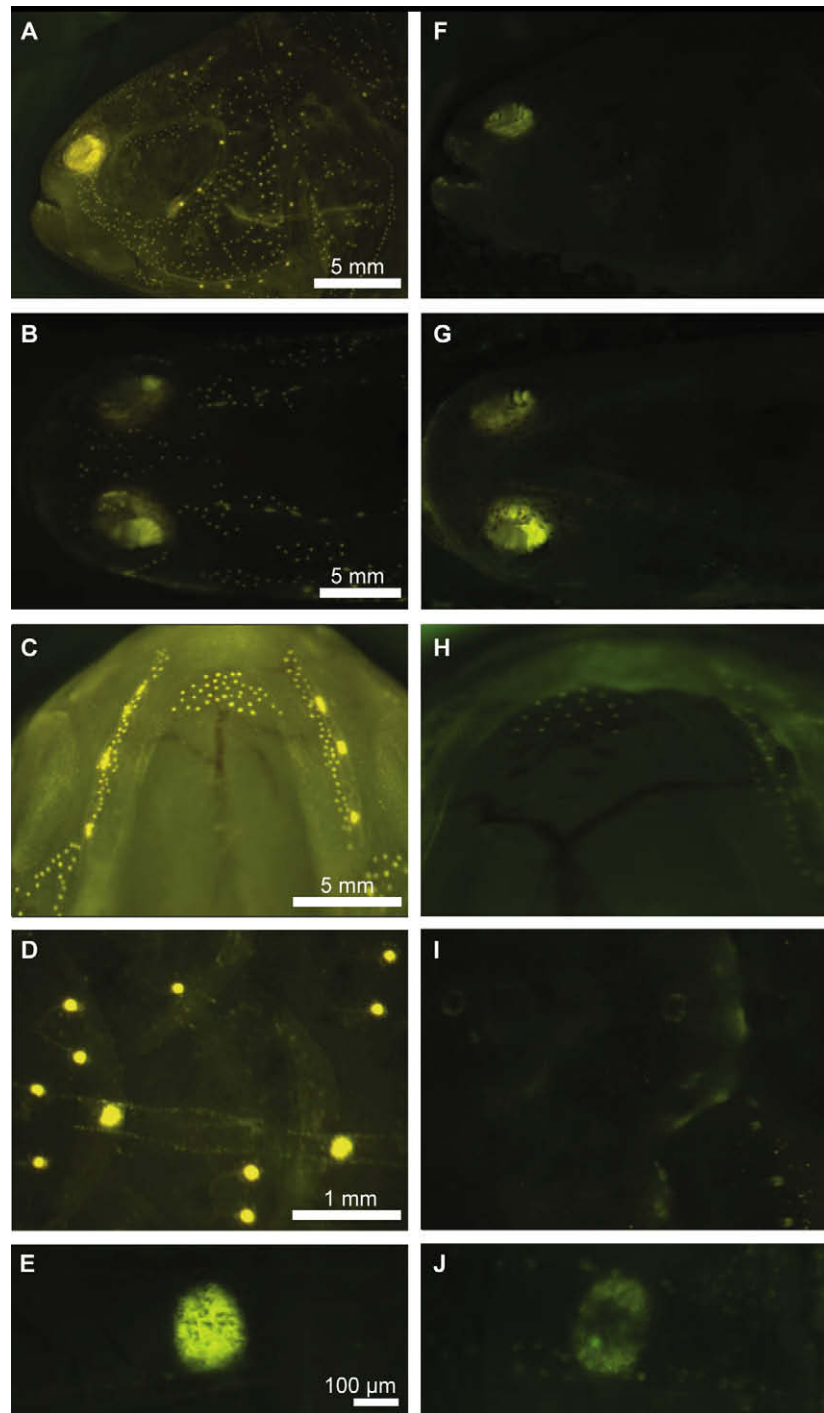
Fig. 2. DASPEI staining of the lateral line system in a Mexican blind cave fish from the control (A–E) and treatment (F–J) groups. These photographs were taken from lateral (A, F), and dorsal (B, G) perspectives of the head, a ventral view of the mandible (C, H), a lateral view of the trunk (D, I), and a single trunk canal neuromast (E, J) showing the pattern of weak surround labeling in a treated neuromast (J).

These results were obtained despite using doses and exposure durations much lower than those previously reported to cause destruction of ciliary bundles in CN, but not SN hair cells (Song et al., 1995). Moreover, this finding is supported by the work of two independent laboratories, using three quantification procedures, with two vital stains, and in two species. These findings have implications for our understanding of aminoglycoside ototoxicity and the behavioral roles of the two submodalities of the lateral line system.

4.1. Ototoxic effects of gentamicin in lateral line hair cells

In contrast to our findings, studies that have visualized neuromasts with SEM reported morphological disruption of hair cells in CNs, but not SNs in three different species: oscar *Astronotus ocellatus* (Song et al., 1995; Montgomery et al., 1997), Mexican blind cave fish, *A. fasciatus* (Baker and Montgomery, 1999a) and mottled sculpin, *Cottus bairdi*, (Coombs et al., 2001). This pattern even persisted for several days following gentamicin
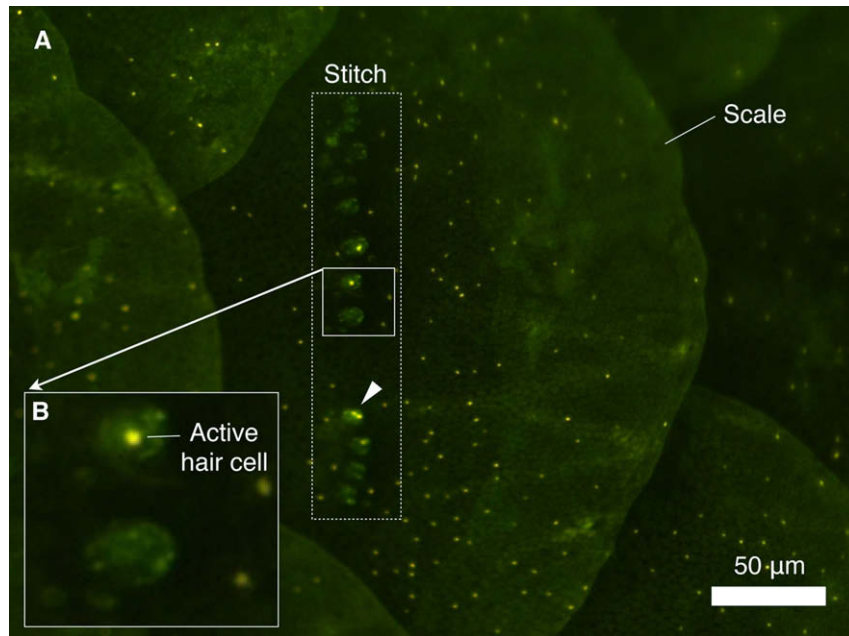
Fig. 3. The location of weakly labeled superficial neuromasts is visible in an individual zebrafish that was treated with gentamicin and subsequently stained with DASPEI. The stain fluoresces strongly in one or two active hair cells and weakly in surrounding cells within the neuromast. (A) 14 neuromasts are arranged in a vertical series, called a stitch (dashed box). Only one neuromast is revealed to have more than a single active hair cell (arrowhead). (B) A detailed view of two weakly labeled neuromasts illustrates staining of a single (above) and no (below) active hair cells.

exposure, until CN hair cells eventually regenerated, approximately 10–20 days after the initial gentamicin treatment (Coombs et al., 2001). Therefore, the prior evidence showing a differential effect of gentamicin is compelling. Understanding the discrepancy between vital dye and SEM results requires a consideration of what each method indicates. Gentamicin is capable of disabling a neuromast either by blocking the mechanotransduction channels (Kroese and van den Bercken, 1982; Forge et al., 1989; Moczydlowski et al., 1985; Ernst et al., 1994), or triggering a process that leads to the degeneration of apical cilia, and hair cell apoptosis (Williams et al., 1987; Forge and Schacht 2000). Either mechanism disrupts hair cell function

and therefore prohibits DASPEI staining, but only cilia degeneration or apoptosis would generate the observed SEM findings. Therefore, the discrepancy between the present DASPEI staining results and previous findings could be explained if gentamicin blocks the channels of both CN and SN hair cells, but causes cilia degeneration in CNs only. Under such conditions, neither group of hair cells would be labeled by DASPEI and only CN hair cells would appear damaged in SEM. However, our results from FM1-43 staining oppose this interpretation. Neuromasts pre-labeled with FM1-43 were significantly less numerous in the treatment group compared to the control in both CNs and SNs (Figs. 4, 5C, and Table 3). This suggested that in addition to possibly

Fig. 4. FM1-43 staining of the lateral line system in the head of a zebrafish from the control (A) and treatment (B) groups. These images were produced about 24 h after staining to allow the time necessary for the gentamicin treatment. Note the high contrast of neuromasts in the control fish (A). Staining can be observed in ~18 superficial neuromasts along the operculum (dashed ellipse). In contrast, treated fish (B) retain fewer neuromasts with labeled hair cells.

Fig. 5. Comparison of numbers of neuromasts visible with vital staining. For each fish, we included counts of superfic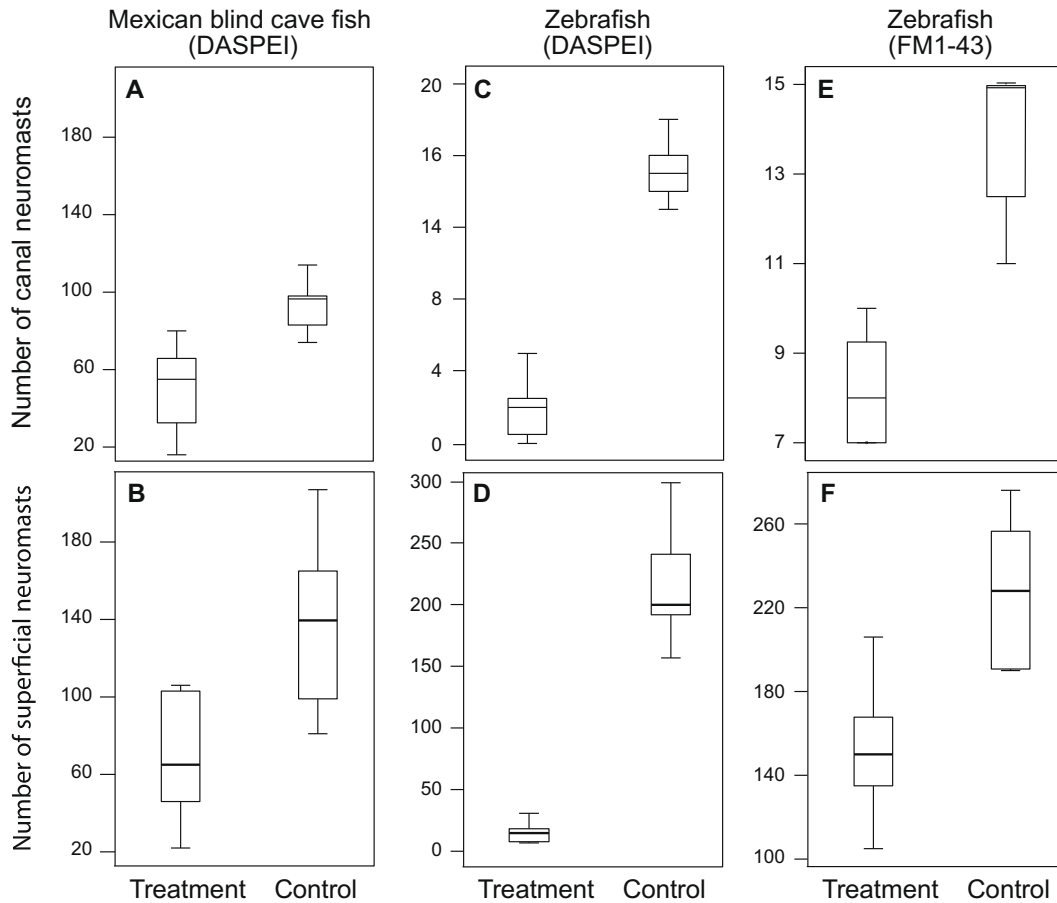ial and canal neuromasts as the sum from all regions of the body. The variation among individuals is represented by the boxplots that denote the median (central line), 25th and 75th percentiles (outer margins of box) and the range (whiskers). Counts of canal (A) and superficial (B) neuromasts were visualized with DASPEI in Mexican blind cavefish from both treatment ($n = 5$) and control ($n = 6$) groups (Table 1). DASPEI staining in zebrafish revealed a similar difference between treatment ($n = 8$) and control ($n = 8$) groups in both canal (C) and superficial (D) neuromasts. These counts included the sum of viable and damaged neuromasts (Table 2). A similar difference was measured between treatment ($n = 5$) and control ($n = 5$) fish stained with FM1-43 in both canal (E) and superficial (F) neuromasts (Table 3). In all cases, a two-factor ANOVA found significant differences between the treatment and control groups.

Table 1
Number of neuromasts stained with DASPEI in Mexican blind cavefish.

Treatment					Control				
	Cranial	Trunk	Mandible	Trunk		Cranial	Trunk	Mandible	Trunk
Individual	CN	CN	SN	SN	Individual	CN	CN	SN	SN
1	48	32	17	89	1	50	24	30	69
2	29	32	14	88	2	46	52	24	107
3	6	10	0	22	3	50	48	42	165
4	12	26	11	43	4	45	50	29	119
5	34	21	18	47	5	52	62	41	124
					6	49	34	31	50
Mean	26	24	12	58		49	45	33	106
1SD	17	9	7	30		3	14	7	41

The treatment group was exposed to a gentamicin solution (see Section 2 for details).

blocking the mechanotransduction channels, gentamicin exposure leads to loss of hair cells in both types of neuromasts.

Although gentamicin is ototoxic to all hair cells in the lateral line system (Fig. 5), developmental differences between CNs and SNs could create different patterns of damage from gentamicin exposure. CNs appear in embryonic development and, unlike SNs, do not increase in numbers as the fish matures (Webb and Shirey, 2003; López-Schier and Hudspeth, 2005; Nuñez et al., 2009). It is therefore likely that an adult fish will possess some SNs that contain a greater proportion of functionally immature and regenerat-

ing hair cells than in its CNs. Immature and recently regenerated hair cells are resistant to aminoglycoside antibiotics (Dai et al., 2006; Hashino and Salvi, 1996; Murakami et al., 2003) because they lack a complete compliment of functioning mechanotranduction channels. These channels provide the major means for entry by the antibiotics to damage the cell (Kroese et al., 1989; Marcotti et al., 2005). If gentamicin exposure ceased prior to the maturation of these cells, then SN could appear resistant to gentamicin by virtue of possessing a greater number of immature and regenerating hair cells.

Table 2
Number of neuromasts stained with DASPEI in zebrafish.

Treatment group							Control group						
	CNs			SNs				CNs			SNs		
Individual	Viable	Damaged	Non-Viable	Viable	Damaged	Non-Viable	Individual	Viable	Damaged	Non-Viable	Viable	Damaged	Non-Viable
1	0	1	10	0	17	137	1	13	1	0	179	21	17
2	0	5	8	0	13	98	2	13	0	0	167	32	18
3	1	2	9	5	26	148	3	13	1	0	163	22	46
4	0	2	10	0	18	146	4	17	1	0	259	12	7
5	0	0	11	0	9	184	5	16	0	0	267	32	36
6	0	0	16	0	7	168	6	16	0	0	189	22	16
7	0	2	11	0	19	209	7	15	0	0	142	15	22
8	0	2	12	0	7	182	8	15	0	0	184	16	26
Mean	0	2	11	1	15	159		15	0	0	194	22	24
1SD	0	2	2	2	7	34		2	1	0	45	7	12

The treatment group was exposed to a gentamicin solution (see Section 2 for details).

Table 3
Number of neuromasts stained with FM1-43 in zebrafish.

Treatment			Control			
	Cranial	Cranial	Trunk		Cranial	Cranial	Trunk
Individual	CN	SN	SN	Individual	CN	SN	SN
1	8	13	92	1	15	18	173
2	10	13	137	2	13	10	218
3	7	15	130	3	15	26	250
4	9	14	192	4	15	24	226
5	7	14	141	5	11	18	172
Mean	8	14	138		14	19	208
1SD	1	1	36		2	6	34

The treatment group was exposed to a gentamicin solution (see Section 2 for details).

It remains possible that SN and CN hair cells possess subtle distinctions in their mechanism of ototoxicity. Recently, it has been shown that apoptosis may occur in lateral line hair cells through separate rapid (within 1 h of exposure) and/or slow pathways (after 6 h of exposure) (Coffin et al., 2009; Owens et al., 2009). If SNs were more susceptible to rapid cell death than CNs, then most of the hair cells within SNs would die within the first hour of exposure. SN hair cells would thus have an opportunity to regenerate prior to visualization (4 days later, as in Song et al., 1995). These recently regenerated hair cells would be resistant to gentamicin (Hashino and Salvi, 1996; Dai et al., 2006). CNs respond to both time-courses of cell death, as seen in the presumptive canal neuromasts of zebrafish larvae (Owens et al., 2009). Therefore, CNs may lose many of their hair cells at a later time, leaving less time for the regeneration of new hair cells.

Although not entirely reconcilable with previous SEM results, we contend that vital stains offer a superior method for assessing ototoxicity because they provide a comprehensive view of functioning hair cells *in vivo*. From these data, we cannot determine whether or not gentamicin blocks the mechanotransduction channels of hair cells in both neuromast classes. However, our findings do suggest that gentamicin has similar, short-term effects on the overall viability of lateral line hair cells in both canal and superficial neuromasts. If differences do exist in their response to gentamicin, then it is likely a more subtle, and time-dependent, distinction than previously believed.

4.2. Gentamicin and the role of lateral line submodalities in fish behavior

Gentamicin has been used as a tool for selectively blocking CNs to test the role of this submodality in fish behavior. Our finding that gentamicin is ototoxic to both CNs and SNs is cause for the results of these studies to be re-evaluated. Because gentamicin disables both submodalities, we predict that gentamicin-treated fish should behave similarly to fish treated with other non-selective blockers of lateral line function, such as streptomycin sulfate (Kaus, 1987) or cobalt chloride (Karlsen and Sand, 1987). This prediction is consistent with the results of some previous studies, whereas others require re-interpretation. Here we survey the findings of a few studies that have investigated the relative contributions of CN and SN submodalities by gentamicin treatment.

4.2.1. Object entrainment

Montgomery et al. (2003) found that trout (*Oncorhynchus mykiss*) entrain their swimming behind a cylinder in flow to a similar degree when treated with either gentamicin or streptomycin. This behavior occurred less often in treated fish than in the control group. The authors concluded that both classes of neuromast contributed to entrained swimming under the assumptions that (1) both submodalities were blocked with streptomycin but (2) only the CN submodality was blocked with gentamicin. Our results suggest that the second assumption is incorrect and that, while the lateral line system contributes to this behavior (as confirmed by Liao, 2006), the relative role of each submodality remains inconclusive.

4.2.2. Rheotaxis

The ability of fish to orient to flow, rheotaxis, has been ascribed exclusively to sensory input from the SN submodality. Ablation of the entire lateral line system by treatment with either cobalt chloride or streptomycin sulfate caused three species of fish (including the Mexican blind cave fish) to be less capable of orienting their body towards flow (Montgomery et al., 1997; Baker and Montgomery, 1999a,b). Despite our prediction that gentamicin should have a similar influence, the number of individuals exhibiting rheotaxis was significantly greater for the gentamicin group than the cobalt chloride or streptomycin treated groups. Furthermore, there were no significant differences between control and gentamicin-treated fish at any flow velocity. Thus, it is difficult to reconcile our current findings and predictions with the gentamicin results of these studies. Nevertheless, the conclusion that rheotaxis is mediated by SNs at low current velocities may still be correct. This is because the physical ablation of SNs revealed that fewer individuals exhibited rheotaxis than in control groups.

4.2.3. Prey capture

Previous studies have suggested distinct roles for the two lateral line submodalities during prey capture. Coombs et al. (2001) found that prey-orienting responses to a vibrating sphere were significantly reduced in gentamicin-treated mottled sculpin (*C. bairdi*)

compared to a control group. The treated fish were also less capable of this behavior than a group that had its SNs physically ablated. This SN-ablated group behaved just as effectively as the control. Therefore, the authors' conclusion that prey-orienting behavior depends on CNs, but not SNs, is not invalidated by our discovery that gentamicin blocks both SNs and CNs. Montgomery et al. (2002), found that both gentamicin treatment and physical ablation of SNs caused Mexican blind cave fish to take a significantly longer time to capture prey. The authors concluded that neuromasts of both types were important to predation, but it now appears that while predation success in this species is indeed influenced by SNs, the contribution of CNs remains untested and uncertain. Therefore, the relevant roles of CNs and SNs in prey capture remains enigmatic, but differences between fish species appear likely.

In summary, our finding that gentamicin is ototoxic to all lateral line hair cells means (1) that hair cells in the SNs of the lateral line can no longer be regarded as functionally resistant to gentamicin toxicity, (2) that this drug should therefore no longer be used as a pharmacological tool for selective blocking of CN, but not SN hair cells, and (3) that the conclusions of some previous studies need to be re-evaluated. As evidenced by the frequent citation of these papers in review articles (e.g. Coombs and Montgomery, 1999; Bleckmann, 2008) and book chapters (e.g. Janssen, 2004; Coombs and van Netten, 2006; and Higgs et al., 2006), these studies have been influential in shaping our understanding of lateral line mediated behavior. Our results suggest that much of our understanding for the individual roles of canal and superficial neuromasts in behavior remains unresolved.

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