Natural Bizbenzoquinoline Derivatives Protect Zebrafish Lateral Line Sensory Hair Cells from Aminoglycoside Toxicity

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INTRODUCTION

Over 360 million people worldwide are affected by moderate to severe hearing impairment, including 37.5 million Americans (Blackwell et al., 2014; World Health Organization, 2015). Hearing loss can have devastating consequences including social isolation, decreased employment opportunities, and a loss of household income (Kochkin, 2007; Mick et al., 2014; Emmett and Francis, 2015). Hearing loss is caused by genetic mutations, aging, loud noise, and exposure to certain ototoxic medications such as the aminoglycoside antibiotics neomycin.
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Aminoglycosides are commonly prescribed to treat antibiotic resistant gram-negative bacterial infections such as tuberculosis (Durante-Mangoni et al., 2009), neonatal infections, and infections associated with cystic fibrosis (Rizzi and Hirose, 2007). For example, in cystic fibrosis patients, infection caused by *Pseudomonas aeruginosa* responds positively to aminoglycoside treatment (Vázquez-Espinosa et al., 2015). However, as a side effect of treatment, approximately 20–30% of patients suffer from ototoxic damage (Rizzi and Hirose, 2007; Xie et al., 2011; Schacht et al., 2012). Methods are needed to ameliorate this damage and promote safe use of these antibiotics.

Aminoglycoside-induced hearing loss results from damage to sensory hair cells of the inner ear (Schacht et al., 2012). Aminoglycosides kill hair cells via activation of multiple signaling cascades, including programmed cell death pathways (Forge and Schacht, 2000; Matsui et al., 2002; Jiang et al., 2006; Coffin et al., 2013b). Aminoglycoside exposure is correlated with increased reactive oxygen species, a loss of mitochondrial membrane potential, and subsequent hair cell death, sometimes accompanied by signs of classical apoptosis such as nuclear condensation and caspase activation (Forge and Li, 2000; Matsui et al., 2002, 2004; Hirose et al., 2004; Owens et al., 2007). However, several lines of evidence suggest that different aminoglycosides may activate different cell death pathways and that even a single aminoglycoside may act on multiple signaling pathways within a single sensory epithelium (Jiang et al., 2006; Owens et al., 2009; Coffin et al., 2013a,b). For example, Jiang et al. (2006) found variable cell morphology in cochleae from aminoglycoside-treated mice, indicative of multiple modes of cell death. Furthermore, they did not find evidence for caspase activation, but rather for activation of other proteases such as calpains and cathepsins. Similarly, aminoglycoside toxicity in the zebrafish lateral line is likely caspase-independent (Coffin et al., 2013b). Different aminoglycosides also activate only partially-overlapping cell death pathways in the lateral line, with neomycin activating mitochondrially-associated signaling via Bax, and gentamicin activating Bax-independent mechanisms that act through p53 (Owens et al., 2009; Coffin et al., 2013a). Compounds that modulate these intracellular signaling pathways offer therapeutic options for preventing aminoglycoside ototoxicity. However, given the complexity of the cell signaling events involved, it is often difficult to take an *a priori* approach to selecting a single molecular target for manipulation. We have therefore adopted an objective screen with the goal of identifying one or more natural compounds that prevent aminoglycoside ototoxicity.

Natural compounds such as plant extracts offer a novel source of otoprotective drugs. Natural compounds have been used in Eastern medicine for thousands of years and are still used today by people around the world (Ji et al., 2009). Recent evidence demonstrates their efficacy in some clinical scenarios. For example, the *Ginkgo biloba* extract EGB 760 attenuated neuronal loss in a mouse model of ischemic stroke and enhanced neurogenesis post-stroke (Nada et al., 2014). Furthermore, many natural compounds are available at low cost, allowing the possibility of relatively rapid transition to the clinical setting. We examined a library of natural compounds using the zebrafish (*Danio rerio*) lateral line as a model system to discover compounds that protect hair cells from aminoglycoside ototoxicity.

The zebrafish lateral line is an excellent model for high throughput screening of compounds that modulate hair cell survival, an intractable approach in mammalian systems (Harris et al., 2003; Owens et al., 2008; Coffin et al., 2013a,b). Lateral line hair cells are used to detect vibrations in the water for several behaviors, including predator avoidance and schooling (Partridge and Pitcher, 1980; Hoekstra and Janssen, 1983; Montgomery and Hamilton, 1997). These externally located hair cells are stereotypically organized into clusters known as neuromasts that are arrayed along the head and trunk of the fish (Metcalfe et al., 1985; Raible and Kruse, 2000). This external location provides an additional advantage, allowing for rapid access for pharmacologic manipulation similar to an *in vitro* preparation in an *in vivo* system. Zebrafish lateral line hair cells are structurally and functionally similar to mammalian hair cells. All vertebrate hair cells share core features, including an apical polarized hair bundle with mechanotransduction machinery (e.g., TMC proteins) and extracellular tip links composed of cadherin 23 and protocadherin 15 (Söllner et al., 2004; Kazmierczak et al., 2007; Pan et al., 2013; Maeda et al., 2014). Mutations in several of these proteins, for example the hair bundle motor protein Myosin VIIA, cause deafness in both mammals and zebrafish (Self et al., 1998; Ernest et al., 2000). Like inner ear hair cells, lateral line hair cells are bathed in a regulated ionic environment, with a gelatinous cupula overlying the apical hair bundles, similar to the cupula in the canal cristae of mammalian inner ears (Russell and Sellick, 1976; Valli et al., 1977; Van Netten, 1997). One key difference is that zebrafish hair cells, and indeed hair cells in all non-mammalian vertebrates, regenerate following ototoxic damage. This regeneration depends critically on supporting cells, rather than the hair cells themselves, such that while supporting cells suffer at the expense of supporting groups, the hair cells are highly similar (Brignull et al., 2009). Importantly, fish hair cells respond similarly to toxic insult, making it likely that otoprotectants found in zebrafish studies will translate to mammals (Harris et al., 2003; Nicolson, 2005; Ton and Parng, 2005; Ou et al., 2007, 2009; Owens et al., 2008; Esterberg et al., 2013). For example, prior screening efforts demonstrated that the small molecule PROTO1 and the acetylcholinesterase inhibitor tacrine protect both zebrafish and mammalian hair cells from neomycin damage, highlighting the tractability of using zebrafish to discover otoprotective drugs (Owens et al., 2008; Ou et al., 2009). Using the zebrafish lateral line, we screened a natural compound library and discovered four novel compounds that protect hair cells from aminoglycoside exposure.

**MATERIALS AND METHODS**

**Animals**

Zebrafish were maintained in zebrafish facilities at both Washington State University Vancouver and Amherst College.
All cell biology experiments were conducted on 5–6 day old larval zebrafish (*AB strain, brm3c:mGFP (Tg(pou4f3:gap43-GFP)), or myo6b:EGFP (Tg(myo6b:EGFP))) maintained at 28.5°C in defined E2 embryo medium (EM) containing 1 mM MgSO₄, 120 µM KH₂PO₄, 74 µM Na₂HPO₄, 1 mM CaCl₂, 500 µM KCl, 15 mM NaCl, and 500 µM NaHCO₃ in distilled water at a pH of 7.2 (Westerfield, 2000). We selected this age range because hair cells in 5 day-old fish show mature responses to ototoxic insult, and the small fish size allows for high throughput screening of compounds in small volumes (Murakami et al., 2003; Santos et al., 2006; Owens et al., 2008; Vlasits et al., 2012). Microphonic experiments used wild-type Ekkwil zebrafish (Ekkwil Waterlife Resources), which also exhibit neomycin-induced hair cell death similar to that observed in *AB fish (Kruger, unpublished data). All procedures were approved by the appropriate Institutional Animal Care and Use Committee at Washington State University and Amherst College.

**Aminoglycoside Treatment**

Neomycin or gentamicin were used to induce damage to hair cells. Neomycin (10 mg/ml) and gentamicin (50 mg/ml) were acquired from Sigma-Aldrich (St. Louis, MO, USA) and diluted in EM. Fish were incubated with either neomycin for 30 min (termed “acute exposure”), or gentamicin for 6 h (termed “continuous exposure”) at a range of concentrations from 50–400 µM. Exposed fish were then rinsed 4× with fresh EM. Neomycin treated fish were allowed to recover for 1 h prior to hair cell assessment, while gentamicin treated fish were assessed immediately. These acute and continuous exposure time courses were chosen because there are distinct, yet overlapping, cell death pathways activated by aminoglycosides (Owens et al., 2009; Coffin et al., 2013a,b). Therefore, we asked if natural compounds universally protected hair cells from aminoglycoside-induced death, or if protection was specific for neomycin or gentamicin.

**Natural Compound Screen**

We conducted two independent screens of the Enzo Natural Product Library (502 compounds, Enzo Life Sciences) to identify compounds that protect hair cells from neomycin or gentamicin damage. These two aminoglycosides were used because they activate different hair cell death mechanisms, allowing us to proactively screen for drug-specific protectants (Owens et al., 2009; Vlasits et al., 2012; Coffin et al., 2013a,b). Prior to treatment with each natural compound, fish were incubated in a 3 µM solution of the nuclear dye Yo-Pro-1 (Life Technologies) for 30 min to label hair cells (Santos et al., 2006; Owens et al., 2008). Yo-Pro-1 is a non-toxic dye that is highly amenable to screens of this nature because the dye is retained for at least 24 h (Coffin et al., 2009; Thomas et al., 2015). Therefore, hair cells could be labeled prior to treatment, allowing for rapid post-treatment assessment without the need to move the animal or add additional dyes. Three to four fish were placed in each well of a glass bottom 24-well plate; 2–3 wells each were used for positive (neomycin or gentamicin-only) and negative (DMSO-only) controls. 0.2% DMSO served as a negative control because it is the vehicle used to dissolve the natural compounds. Eighteen to 20 wells received aminoglycoside and one of the 502 natural compounds. Fish were pretreated for 1 h with a natural compound (4 µg/ml), co-treated for 1 h with the natural compound and 200 µM neomycin, or co-treated for 6 h with the natural compound and 100 µM gentamicin. Our treatment paradigm results in comparable hair cell loss with either aminoglycoside (Owens et al., 2009; Coffin et al., 2013a,b). Weight by volume (mg/ml) was used here to describe the concentration of the compounds in the Natural Products library instead of molarity because the library is standardized at 2 mg/ml for each compound. However, compounds in the library each have different molecular weights and therefore different molarities.

After treatment, fish were anesthetized with 0.001% buffered 3-amino-benzoic acid ethyl ester methanesulfonate (MS-222, Argent Labs, Redmond, WA, USA) prior to the assessment of anterior lateral line neuromasts (aLL) using a fluorescent dissection microscope (Leica M165FC, Leica Microsystems, Buffalo Grove, IL, USA). Given the rapid nature of the screen, the fluorescent intensity of every aLL neuromast was assessed holistically at once, as opposed to scoring specific neuromasts. A score of 0–3 was assigned to each fish: 0 (no visible hair cells); 1 (dim hair cell fluorescence); 2 (slightly bright hair cells); or 3 (fully bright hair cells; modified from Chiu et al., 2008). All compounds with a score of 2 or greater were rescreened in duplicate, and compounds that were confirmed to be protective through rescreening were analyzed via a concentration-response analysis.

**Hair Cell Assessment**

We used the mitochondrial dye 2-(4-(dimethylamino)styryl)-N-ethylpyridinium iodide (DASPEI) for the bulk of our hair cell assessment experiments because it specifically and robustly labels lateral line hair cells, making it the labeling method of choice for lateral line visualization in many fish species (Harris et al., 2003; Coffin et al., 2009; Owens et al., 2009; Van Trump et al., 2010; Brown et al., 2011). Eight to 15 fish per treatment were incubated in EM containing 0.005% DASPEI (Life Technologies, Grand Island, NY, USA) for 15 min, rinsed 3× with fresh EM and anesthetized in 0.001% buffered MS-222. The brightness of 10 anterior neuromasts (SO1, SO2, IO1-I04, M2, MI1, MI2, O2; Raible and Kruse, 2000) was assessed at 50× magnification using a Leica M165F fluorescent dissecting microscope. A scoring protocol was used to rank each neuromast as: 0 (no labeling); 1 (moderate labeling); or 2 (bright labeling). Neuromast scores were summed for a total score of 0–20 per fish (Harris et al., 2003; Owens et al., 2009).

Since our fluorescence assessment is semiquantitative, we validated the protection from aminoglycoside-induced hair cell death observed with DASPEI assessment through direct hair cell counts using immunocytochemistry. Fish were euthanized with an overdose of MS-222 and fixed with 4% paraformaldehyde overnight at 4°C. Fish were then blocked in 1% goat serum in phosphate-buffered saline (PBS; Life Technologies) with 0.1% Triton-X (PBST; Sigma-Aldrich) at room temperature for 2 h. To
label hair cells, fish were incubated in mouse anti-parvalbumin (EMD Millipore, Billerica, MA, USA), diluted 1:500 in PBST with 1% goat serum (Coffin et al., 2013a). Fish were then rinsed in PBST and labeled with either Alexa 488 or 568 goat anti-mouse secondary antibodies (Life Technologies; diluted 1:500 in PBST). Following 2–3 additional rinses in PBST, and then PBS, fish were stored in 1:1 PBS/glycerol at 4°C prior to assessment. Fish were visualized using a compound fluorescent microscope (Leica DMI4000 B or DMRB). Hair cell number was quantified in five neuromasts (IO1, IO2, IO3, M2, OP1) per fish, summed to calculate one value per animal, and averaged for each group. We further confirmed our hair cell counts using Brn3c:mGFP fish, which express membrane-bound GFP in all hair cells, and myo6b:EGFP fish, which express cytoplasmic GFP in all hair cells (Xiao et al., 2005; Namdaran et al., 2012; Suli et al., 2014). Hair cells were counted in live, anesthetized transgenic larvae or freshly fixed larvae as described above.

Concentration-Response Analysis
Four compounds scored a three during rescreening and were selected for further analysis. To determine the optimally protective concentration (OPC) for each natural compound, zebrafish were pretreated for 1 h with a range of compound concentrations (0.5–50 µM). The fish were then co-treated with the natural compound and 200 µM neomycin (acute) or 100 µM gentamicin (continuous) using the aminoglycoside treatment protocol described above. To determine the effectiveness of each natural compound at protecting hair cells, fish were pretreated with the OPC of each natural compound, then co-treated with the natural compound and 50–400 µM neomycin or gentamicin (Harris et al., 2003; Coffin et al., 2009). To assess broad applicability of potential protectants to diverse ototoxins, we also asked if these natural compounds conferred protection from cisplatin damage. Fish were pretreated with a range of protectant concentrations (0.25–10 µM) for 1 h, followed by co-treatment with protectant and 500 µM cisplatin for 6 h. Protection was assessed using DASPEI scoring and further validated by direct counts of anti-parvalbumin-labeled or GFP+ hair cells (for wildtype and Brn3c transgenic larvae, respectively). Experiments were conducted in triplicate.

Aminoglycoside Uptake
Gentamicin conjugated to the dye Texas Red (GTTR) was used to quantify uptake of aminoglycosides into hair cells (Stegyer et al., 2003; Wang and Steyger, 2009), and FM 1-43FX was used as a proxy for transduction channel function (Gale et al., 2001). Both GTTR and FM 1-43 are reported to primarily enter hair cells via the mechanoelectrical transduction (MET) channel, thereby offering two independent means of assessing channel block; FM 1-43FX is specifically taken up by the MET channel after a very brief exposure time (Alharazneh et al., 2011; Vu et al., 2013). Zebrafish were treated with 50 µM GTTR (made according to Stegyer et al., 2003) for 3 or 18 min or 1.5 µM FM 1-43FX (Molecular Probes, Eugene, OR, USA) for 30 s. The fish were then rinsed 3× in EM, euthanized, and fixed with 4% paraformaldehyde. The IO3 neuromast was imaged using a 20× dry objective with 5× digital zoom on a Leica SP8 laser scanning confocal microscope with the 552 nm laser and 550–630 nm detection for GTTR (peak detection 580 nm), or the 488 nm laser and 590–680 nm detection for FM 1-43FX (peak detection 620 nm); the laser power and gain were kept constant within each experiment. Image stacks were collapsed into maximum projection images for analysis. Neuromast regions of interest (ROIs) were used to quantify average fluorescent intensity of each region (neuromast fluorescence—background fluorescence). Fluorescent intensity was measured in arbitrary units (a.u.) using ImageJ. IO3 neuromasts were selected because they were used for previous hair cell assessment and were consistently visible in every fish.

Washout Experiment
We conducted a washout experiment to determine if natural compound treatment before or after, but not during aminoglycoside exposure, was sufficient to confer protection. We used gentamicin for this experiment because short-term exposure to gentamicin results in increasing hair cell damage over several hours, even after antibiotic removal (Owens et al., 2009). Fish were treated with 200 µM gentamicin for 30 min, followed by a 5.5 h recovery period in EM or a natural compound, and then assessed with DASPEI. Natural compound (or an equal volume of DMSO) was present before, during, and/or after gentamicin treatment, in order to assess the relative timing of the protective effect.

Cell Proliferation
It is possible that protective compounds promote hair cell regeneration, rather than preserving existing hair cells. To examine this possibility, we conducted a cell proliferation assay using bromodeoxyuridine (BrdU). Larvae were treated with a natural compound for 1 h, then co-treated with compound and 200 µM neomycin for 30 min, rinsed in fresh EM, and allowed to recover for 1 h. Ten millimolar BrdU (Sigma) was present for the entire 2.5 h time course. Larvae were then euthanized with an overdose of MS-222, fixed overnight in 4% paraformaldehyde at 4°C, and processed for anti-BrdU immunocytochemistry using the protocol in He et al. (2015). 4,6-diamidino-2-phenylindole (DAPI, Life Technologies) was used as a counter-label to visualize neuromast location. Using a Leica DMRB microscope we quantified the number of BrdU+ cells per neuromast (5 neuromasts per animal: IO1, IO2, IO3, M2, OP1, the same neuromasts that were used for hair cell counts).

Microphonics
MET channel function was assayed by performing extracellular recordings of microphonic potentials from the primary neuromasts (L1, L2, L3, L4) of the posterior lateral line (Trapani and Nicolson, 2010). Larvae were anesthetized with 0.016% MS-222 and pinned down on a silicone-lined (Sylgard-184, Dow Corning, Midland, MI, USA) recording chamber using homemade tungsten pins. The neurotoxin α-Bungarotoxin (125 µM; Abcam, Cambridge, MA, USA) was injected into the heart to prevent interference from muscle movement. To wash
out the MS-222, larvae were rinsed with normal extracellular solution (130 mM NaCl, 2 mM KCl, 2 mM CaCl₂, 1 mM MgCl₂ and 10 mM 4-(2-hydroxyethyl)-1-piperazineethane-sulfonic acid). Neuromasts were deflected ~20° bidirectionally at 20 Hz for 200 ms with a sinusoidal current of water produced by a high-speed pressure clamp (50 mM Hg maximum output, HSPC-1, ALA Scientific, Farmingdale, NY, USA). The water current was directed at a single neuromast with a waterjet micropipette, which was fabricated using a Flaming/Brown style pipette puller (Model P-1000, Sutter Instruments, Novato, CA, USA) to pull borosilicate capillary tubing (B150-110-10HP, Sutter Instruments) to a long micropipette that was then cleanly broken to produce a 30 µm diameter tip. The waterjet was aligned with the anteroposterior axis of the fish and positioned approximately 100 µm from the hair cell bundle with the bottom of the waterjet even with the top of the neuromast cupula. Waterjet position was confirmed using an upright epifluorescent microscope with a 40× water immersion objective (BX51-WI, Olympus, Olympus Center Valley, PA, USA). The recording electrode was also fabricated from glass capillary tubing (BF150-110-10HP, Sutter Instrument) and was positioned just adjacent to the base of the cupula at the level of the hair cell stereocilia. Microphonic potentials were sampled at 20 kHz with a 1 kHz low-pass filter using an Axon Instruments 200B amplifier (Molecular Devices, Sunnyvale, CA, USA) in current-clamp mode (I = 0) with 500× gain and were then amplified (100×) and filtered 100 Hz using a Model 440 amplifier (Brownlee Precision, Palo Alto, CA, USA). Data were acquired with an ITC-16 DAQ device and Patchmaster software (HEKA Elektronik). Microphonic traces in the figures represent an average of 200 consecutive sweeps collected for each recording. The average magnitude of the evoked microphonic potential (in microvolts) was calculated from the integral of the microphonic waveform divided by 0.21 s (microvolt • s/s). The 0.21 s interval was from stimulus onset (t = 150 ms) to 10 ms after the stimulus offset (t = 360 ms).

**Antibiotic Efficacy**

Kirby-Bauer antibiotic efficacy tests were used to determine if the otoprotective natural compounds affect the ability of neomycin or gentamicin to inhibit bacterial growth (Clinical and Laboratory Standards, 2009). Six millimeter filter article discs (Fisher Scientific, Waltham, MA, USA) were soaked overnight in either DMSO (control), 2 µg/ml neomycin (Owens et al., 2008) or 1 µg/ml gentamicin (Cohen et al., 1991; Ferguson, 2008), the OPC of the natural compound, or the natural compound and aminoglycoside. We used the minimum inhibitory concentration (MIC) of each antibiotic; a standard approach for assessing antibiotic efficacy (Clinical and Laboratory Standards, 2009). The MIC assumes that if the compound of interest (in this case, the natural compound) does not interfere with antibiotic efficacy at the lowest effective antibiotic concentration, then that same compound will likely not affect efficacy of higher antibiotic doses. However, we also tested a 15 mg/ml neomycin concentration to rule out compound/antibiotic interactions at higher antibiotic concentrations. The discs were removed from solution and allowed to dry for 1 h prior to plating to reduce saturation. The next day, E. coli strain ATCC25922 was plated on a 150 mm diameter agar plate (Mueller Hinton Agar; Fisher Scientific), and the pretreated discs were spaced evenly on the plate. The plate was then inverted and incubated overnight at 37°C. Images were taken of each plate using a Leica M165F microscope, and the area of E. coli growth inhibition around each disc was measured using ImageJ.

**Data Analysis**

Comparisons were made via one-way or two-way analysis of variance (ANOVA) or t-test with Bonferroni multiple comparison correction, as appropriate, using GraphPad Prism 6.0. Data are presented as mean ± SEM.

**RESULTS**

Using the zebrafish lateral line as a model of hair cell death, we screened a natural compound library for novel compounds that prevented neomycin- or gentamicin-induced hair cell death. Aminoglycoside treatment caused nuclear fragmentation and reduced neuromast fluorescence, while protective compounds largely prevented fragmentation and preserved labeling intensity (Figures 1A,B). From two independent screens, we found nine candidate natural compounds that protected hair cells from neomycin, 14 that protected from gentamicin, and two, hernandezine and decoyinine, that conferred protection from both aminoglycosides (Figure 1C). Overall, the percentage of otoprotective compounds that we initially detected from screening the Enzo Life Science Natural Product Library is comparable to similar otoprotection screens of other compound libraries (Ou et al., 2009; Vlasits et al., 2012).

Four compounds were selected for further evaluation (berbamine, E6 berbamine, hernandezine and isotetrandrine), because they consistently demonstrated 75% or greater otoprotection upon rescreening. We first determined whether 90 min of exposure to berbamine (Ber), E6 berbamine (E6), hernandezine (Her), or isotetrandrine (Iso) alone caused hair cell death (Figure 2A). We found that 25 and 50 µM of either Ber or E6 caused significant, albeit slight, damage to hair cells, with greater damage at 50 µM. We also found that 1 µM Iso slightly damaged hair cells (Figure 2A). However, all four compounds greatly protected hair cells from 200 µM neomycin. Ber, Her, and Iso were highly protective at 25 and 50 µM. In contrast, E6 was optimally protective at 0.5 µM, nearly 100-fold lower than the concentration that resulted in hair cell damage (Figure 2B). Based on these results, we selected the OPC of 0.5 µM for E6, and 25 µM for Ber, Her, and Iso, to further characterize the nature of protection. In order to determine the effectiveness of otoprotection, we asked if the OPC of each natural compound conferred protection from a range of aminoglycoside concentrations. All four natural compounds offered protection from neomycin and gentamicin-induced hair cell death, albeit with some ototoxicity at higher concentrations.
Our primary assessment of hair cell viability used DASPEI labeling, where fluorescent intensity is dependent on mitochondrial potential (Bereiter-Hahn, 1976). However, it is possible that changes in mitochondrial potential do not accurately reflect hair cell death. Therefore, to directly confirm the protection observed using DASPEI scoring, we also counted the number of viable hair cells remaining after treatment in immunocytochemically-labeled wild type larvae, transgenic brn3c:mGFP larvae, and transgenic myo6b:EGFP larvae. All validation measures clearly demonstrated that all four natural compounds protected hair cells from 200 μM acute neomycin exposure, with neomycin alone killing over 50% of the hair cells and each compound conferring approximately 90% protection (Figure 3). The results of these independent assessments confirm the hair cell protection observed and further validate the strong correlation between DASPEI scoring and hair cell counts (Harris et al., 2003; Coffin et al., 2013a).

The lateral line of larval zebrafish rapidly regenerates lost hair cells, with complete regeneration observed 72 h after neomycin damage (Harris et al., 2003; Ma et al., 2008). We therefore wanted to rule out that we were observing a combination of hair cell protection and increased hair cell regeneration. Following neomycin damage and 1 h of recovery, neuromasts had an average of 1.0 ± 0.1 BrdU+ cells per neuromast, and this number did not increase upon co-treatment with Ber, E6, or Her (0.9 ± 0.1, 0.4 ± 0.1, 1.2 ± 0.1 BrdU+ cells per neuromast, respectively, Figure 4). These data demonstrate that our bisbenzoxazinone derivatives protect hair cells from aminoglycoside damage, rather than promoting rapid hair cell regeneration.

**Natural Compounds Protect Hair Cells by Reducing Uptake of Aminoglycoside Antibiotics**

Aminoglycosides rapidly enter hair cells via MET channels that are located at the apical portion of the hair bundle (Alharazneh et al., 2011; Vu et al., 2013). Ou et al. (2012) showed that quinoline ring derivatives reduce uptake of aminoglycosides into hair cells, and all four of our protective compounds contain modified quinoline ring structures (bisbenzylisoquinoline rings; Figure 5). As such, we hypothesized that these natural compounds confer protection by reducing aminoglycoside access to hair cells through the MET channel. To test this hypothesis, we used gentamicin conjugated to Texas Red (GTTR) to quantify uptake of gentamicin into hair cells (Steyger et al., 2003). Application of 25 μM Ber, 0.5 μM E6, or 25 μM Iso significantly reduced uptake of 50 μM GTTR after 18 min of incubation (Figure 6). A similar reduction in uptake was seen after 3 min of incubation in GTTR (data not shown). We elected to remove Her from further investigation because it is 10-fold more expensive than the other natural compounds and is lethal to zebrafish after prolonged exposure (data not shown). For the GTTR experiment, Ca^{2+} was used as a positive control since increasing the calcium concentration decreases the probability of opening the MET channel, reducing the amount of GTTR that enters hair cells (Ricci and Fettiplace, 1998; Coffin et al., 2009).

To independently validate our hypothesis that the natural compounds provide protection by blocking uptake through the MET channels, we also used the fluorescent vital dye, FM 1-43FX, which rapidly increases in fluorescence upon entering a hair cell through the MET channels (Gale et al., 2001; Meyers et al., 2003). Log and half-log concentrations (0.01–100 μM) of E6, Ber, and Iso reduced FM 1-43FX uptake in a concentration-dependent manner (Figure 7). Additionally, a washout experiment with gentamicin showed that protection was abolished if zebrafish were removed from...
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FIGURE 2 | Four natural compounds protect zebrafish lateral line hair cells from aminoglycoside-induced death. (A) High concentrations of berbamine and E6 berbamine (25 or 50 µM) significantly damaged hair cells after 90 min of exposure (one-way ANOVA, berbamine: $F_{(5,84)} = 11.75, p < 0.0001$; E6 berbamine: $F_{(5,53)} = 23.25, p < 0.0001$). Iso also damaged hair cells, with significant loss at 1 µM (one-way ANOVA, $F_{(5,64)} = 2.485, p = 0.0405$). Her did not significantly damage hair cells (one-way ANOVA, $F_{(5,64)} = 1.839, p = 0.1178$) after 90 min. (B) Ber, E6, Her, and Iso robustly protect hair cells from acute 200 µM neomycin exposure, with optimal protective concentrations of 0.5 µM for E6 and 25 µM for Ber, Her, and Iso (one-way ANOVA, E6: $F_{(5,47)} = 47.02, p < 0.001$; Ber: $F_{(5,84)} = 64.81, p < 0.0001$; Her: $F_{(5,65)} = 127.1, p < 0.0001$; Iso: $F_{(5,59)} = 56.59, p < 0.0001$). (C) The optimally protective concentration (OPC) of Ber, E6, Her and Iso robustly protected hair cells from 50–400 µM acute neomycin exposure (two-way ANOVA, treatment effect, $F_{(4,317)} = 212.2, p < 0.0001$; interaction term $F_{(4,317)} = 74.47, p < 0.0001$). (D) All four compounds significantly protected hair cells from 50–400 µM continuous gentamicin exposure (two-way ANOVA, treatment effect, $F_{(4,281)} = 263.8, p < 0.0001$; interaction term $F_{(16,281)} = 20.24, p < 0.0001$). $N = 8–26$ and error bars are ±SEM.

gentamicin before adding Iso, suggesting a transient inhibitory mechanism consistent with an uptake block (Figure 8). In contrast, pre-treatment with Iso conferred significant, albeit reduced, protection from gentamicin toxicity, suggesting that the uptake block may be maintained after compound removal (Figure 8). Similar results were seen in washout experiments with Ber and E6 (data not shown). Lastly, we asked if E6 protected hair cells from cisplatin, another ototoxin that enters hair cells through the MET channel (Thomas et al., 2013). We elected to focus on E6 for the cisplatin experiments because all four natural compounds have similar structures and appear to confer protection via similar mechanisms. Furthermore, since DMSO complexes with cisplatin to increase ototoxic damage, we selected the natural compound with lowest OPC, which also equates to the lowest concentration (0.002%) of DMSO (Uribe et al., 2013). Following 6 h exposure to 500 µM cisplatin, cotreatment with 0.5 µM E6 did not confer protection from hair cell death, but 1 and 10 µM E6 were significantly protective (Figure 9). Collectively, these data support our hypothesis that these bisbenzylisoquinoline-containing alkaloids protect hair cells by reducing uptake of chemical ototoxins through the MET channel.

Protected Hair Cells are Still Functional after Exposure to E6 Berbamine

To determine whether E6 could elicit a protective effect against aminoglycoside ototoxicity without altering hair cell function, we assessed MET channel function using microphonic recordings. The MET channel is a non-selective cation channel...
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FIGURE 3 | Direct hair cell counts confirm otoprotection. Ber, E6, and Iso robustly protect hair cells from 200 µM neomycin after acute exposure. (A–F) Representative low magnification (A–C) and high magnification (D–F) images of myo6b:EGFP transgenic fish treated with the indicated conditions. White circles in (A–C) denote the neuromasts shown in (D–F). Scale bar in (A) = 40 µm and applies to panels (A–C). Scale bar in (D) = 10 µm and applies to panels (D–F). (G–I) Representative images of brn3c: mGFP hair cells treated with the indicated conditions. (G) The dotted line encircles a hair cell. Scale bar in (G) = 10 µm and applies to panels (G–I). (J) Direct counts of hair cells in five neuromasts per fish demonstrate significant hair cell protection in AB fish labeled with anti-parvalbumin (black bars, one-way ANOVA, \(F_{(4,60)} = 48.88, p < 0.0001\)), Brn3c fish (light gray bars, one-way ANOVA, \(F_{(4,53)} = 75.68, p < 0.0001\)), and myo6: EGFP fish (dark gray bars, one-way ANOVA, \(F_{(4,53)} = 14.86, p < 0.0001\)). \(N = 8–24\) and error bars are ±SEM. and \(****p < 0.0001\).

FIGURE 4 | Ber, E6, and Her do not enhance regeneration rates after neomycin damage. Fish were treated with either natural compound only or natural compound and 200 µM neomycin in the presence of BrdU. Fish were counterstained with DAPI to identify neuromasts based on characteristic morphology, as shown in the inset where DAPI-labeled nuclei are blue and red nuclei are BrdU+. The line encircles the neuromast and the arrow denotes a BrdU+ supporting cell at the neuromast periphery. Scale bar = 10 µm. There is no difference in the number of BrdU+ cells per neuromast in fish that received neomycin + compound (gray bars) vs. those that received compound only (black bars; 2-way ANOVA, treatment effect \(F_{(1,67)} = 1.882, p = 0.175\)). \(N = 8–10\) and bars are ±SEM.

that opens upon mechanical deflection of the hair bundle (Howard et al., 1988). When neuromast hair bundles are deflected, inward potassium and calcium currents through the MET channel result in depolarization of the receptor potential and a hyperpolarization of extracellularly-recorded microphonic potentials (Corey and Hudspeth, 1983; Nicolson et al., 1998).

Consistent with hair cell damage, neomycin treatment caused a significant reduction in microphonic potentials evoked by 200 ms of 20-Hz sinusoidal stimulation (Figure 10A). In contrast, hair cells co-treated with E6 and neomycin had robust, albeit significantly reduced, microphonic potentials, indicating that many of the E6-protected hair cells remained functional (Figure 10B). To assay whether E6 is able to acutely block hair cell transduction at the concentrations used for our protection experiments, we applied 0.5 µM E6 to the bath while recording microphonics. Direct application of E6 did not reduce microphonic potentials. As a positive control, we then confirmed acute block of microphonics with 1 mM dihydrostreptomycin (DHS), a known MET blocker (Marcotti et al., 2005), which reduced microphonic potentials significantly (Figures 10C,D). Increasing E6 to a concentration (25 µM) that significantly blocked FM 1-43X uptake also had no effect on microphonic potentials \((p = 0.2)\), nor did application of 25 µM Iso \((p = 0.9)\). Overall, these electrophysiological results demonstrate that these bizbenzoquinoline derivatives protected lateral line hair cells from aminoglycoside toxicity without affecting their function, making them candidates for therapeutic development.
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Protective Compounds do not Interfere with Aminoglycoside Antibiotic Efficacy

To determine if E6, Ber, or Iso altered antibiotic efficacy, we conducted a Kirby-Bauer agar diffusion test (Clinical and Laboratory Standards, 2009). We co-treated *E. coli* with the optimal protective concentration of each natural compound and the MIC of neomycin (2 µg/ml) or gentamicin (1 µg/ml) and measured the area of inhibited growth. None of the natural compounds affected neomycin’s or gentamicin’s ability to inhibit growth of *E. coli* strain ATCC25922 (Figure 11). Furthermore, none of the natural compounds alone inhibited bacterial growth. We observed similar results using 15 mg/ml neomycin. The higher neomycin concentration yielded a growth inhibition area of 34.0 ± 2.3 cm, while neomycin plus E6, Her, or Iso lead to inhibition areas of 35.2 ± 2.2, 34.9 ± 1.9, and 36.3 ± 1.7 cm, respectively.

**DISCUSSION**

Our initial independent otoprotection screens using neomycin and gentamicin detected 23 natural compounds that protected hair cells from ototoxic damage. To weed out false positives, the 23 natural compounds were rescreened in duplicate. We discovered four bisbenzylisoquinoline compounds from the two screens, berbamine, E6 berbamine, hernandezine, and isotetrandrine, that consistently protected hair cells from acute neomycin and continuous gentamicin exposure paradigms; the other 19 natural compounds were not protective upon repeated testing. All four protective compounds substantially reduced GTTR and FM 1-43FX uptake, two compounds that enter hair cells via the MET channel. Electrophysiological recordings of microphonic potentials confirmed that natural compounds alone do not disrupt hair cell function and also serve to
preserve hair-cell transduction following ototoxic insult. To our knowledge this is the first time that a library of natural compounds has been screened for otoprotection.

Our results suggest that these bisbenzylisoquinoline-containing natural compounds protect hair cells by blocking aminoglycoside uptake through the MET channel, which is the primary site of aminoglycoside entry in hair cells (Marcotti et al., 2005; Alharazneh et al., 2011; Vu et al., 2013). However, other routes of aminoglycoside entry have been reported, raising the possibility that these natural compounds act by blocking one of the alternative uptake mechanisms (Huth et al., 2011). One alternative is that these natural compounds interfere with receptor-mediated endocytosis, since kanamycin was shown to colocalize with cationic ferritin (a marker for receptor-mediated endocytosis) in regions immediately below the apical plasma membrane of chicken sensory hair cells (Hashino and Shero, 1995). It is also possible that these natural compounds are acting as antioxidants to protect hair cells, since isoquinoline alkaloids have some antioxidant abilities (Zarei et al., 2015). However, this antioxidant hypothesis seems unlikely as no protection was observed in fish treated with natural compounds after gentamicin exposure, a time when cell death is occurring. The uptake block may be temporary, as removing the natural compound prior to treatment with gentamicin attenuated protection. However, some protection was retained in our washout study, suggesting that partial block may be maintained. All four otoprotective compounds have modified quinoline-like ring structures, and previous studies demonstrate that other quinoline ring derivatives, such as the FDA approved bioactive compounds tacrine and amsacrine, also block uptake of aminoglycosides into hair cells (Ou et al., 2009, 2012). The unique structure of these natural compounds may provide new information about how quinoline rings attenuate aminoglycoside uptake. For example, E6 has an additional benzene ring with an attached nitrite group that may increase its affinity for the MET channel (or other uptake receptor) compared to Ber, consistent with E6 acting at a substantially lower concentration. It is possible that the quinoline rings structures are not necessary to confer protection,
FIGURE 10 | Co-treatment with E6 berbamine protects hair cells from neomycin-induced loss of microphonics. (A) Representative microphonic traces from control, E6, E6 + Neo, and neomycin-treated hair cells during 20 Hz stimulation for 200 ms. Treatment with 200 µM neomycin alone results in a loss of microphonic potentials. (B) Microphonic potentials for control and E6 only group are not significantly different from one another (one-way ANOVA, multiple comparisons, p > 0.05). However, there was a significant difference between the control and the E6 with neo group, where hair cells treated with E6 + neo had significantly reduced microphonics compared to control or E6-only-treated cells (one-way ANOVA, multiple comparisons, p < 0.05). Furthermore, E6 prevents the loss of microphonic potentials normally seen with neomycin (one-way ANOVA, $F_{(3,39)} = 6.609$, p = 0.001). (C) Acute application of 1 mM dihydrostreptomycin (DHS) significantly reduces microphonic potentials by nearly 87% (paired t-test, p < 0.0001). (D) Acute application of E6 does not affect microphonic potentials compared to control (paired t-test, p > 0.05). N = 6–16 and bars are ±SEM, *p < 0.05, ***p < 0.001, and ****p < 0.0001.
and future experiments will examine a suite of chemical analogs to determine the minimum chemical structure that confers otoprotection. Whether or not the modified quinoline ring structures are necessary, our data demonstrate that these compounds protect hair cells by preventing aminoglycoside entry. By temporarily blocking ototoxin uptake, the mode of action of these natural compounds may be an appealing way to prevent aminoglycoside-induced hair cell death, as modulating specific intracellular signaling pathways is often complex and may result in unintended consequences for non-target cells. The nature of the relationship between these bis-benzoquinoline derivatives and aminoglycoside uptake mechanisms offers an avenue for future exploration.

As stated above, E6 was chosen for additional study because it is relatively inexpensive and had the lowest optimal protective concentration of the four compounds and of any other quinoline derivative found to date, suggesting a greater potency (Ou et al., 2012). Although E6 reduced uptake of GTTR and FM 1-43FX, microphonic recordings from lateral line hair cells showed that E6-treatment alone did not disrupt normal hair cell transduction, with hair cells previously exposed to E6 and neomycin displaying robust microphonic recordings. Additionally, acute application of E6 did not significantly reduce microphonics, suggesting that if E6 protection is conferred by direct interaction with the MET channel, it is likely not disrupting cation flow through the channel. Consistent with an interaction with the MET channel, E6 application was able to block larger molecules, such as neomycin (614 Da), gentamicin (477 Da), GTTR (1165 Da) and FM 1-43FX (611 Da), which are presumably unable to enter the partially obscured channel. An alternative possibility is that the natural compounds have a faster on/off rate than the aminoglycosides, allowing them to outcompete the larger molecules for access to the channel during cotreatment. However, this competitive inhibition should have resulted in reduced microphonics during acute application of E6 at concentrations that blocked FM 1-43X uptake. Another possibility is that these compounds reduce aminoglycoside uptake by a more indirect interaction with the MET channel similar to amiloride (Jørgensen and Ohmori, 1988). Alternatively, these compounds could cause modulation of intracellular calcium which may alter the MET channel open probability, since quinoline derivatives have been shown to block calcium influx associated with depleted calcium pools in bovine endothelial cells (Low et al., 1996; Ricci and Fettiplace, 1998; Eatock, 2000). However, this alternative explanation is unlikely since we did not see a reduction in the microphonic potentials, which would be expected with changes in calcium dynamics (Nicolson et al., 1998).

Cisplatin also enters hair cells through the MET channel (Thomas et al., 2013), but 0.5 μM E6 did not protect hair cells from cisplatin toxicity, while this concentration confers essentially complete protection from aminoglycoside damage. However, increasing the concentration of E6 to 1 or 10 μM significantly protected hair cells from cisplatin toxicity.
This result is not surprising given that cisplatin molecules (300 Da) are smaller than the larger aminoglycoside compounds. Future research is needed to determine how these otoprotective natural compounds interact with the MET channel or other antibiotic entry route to reduce aminoglycoside uptake.

Other investigators have used the zebrafish lateral line to discover otoprotective compounds. Vlasits et al. (2012) screened a library of 640 FDA-approved compounds and identified 10 compounds that protected from acute neomycin exposure, with only seven of those 10 compounds protecting from continuous gentamicin exposure. It is not surprising that the Vlasits screen found more otoprotective compounds, as all of the compounds in the FDA library were known to activate specific cellular pathways. Similarly, a screen of a library containing 1040 biologically active compounds identified seven compounds that prevented neomycin ototoxicity (Ou et al., 2009). Screenin a library of 10,960 small molecules revealed two benzothiophene carboxamide compounds (PROTO-1 and PROTO-2) that protected hair cells from acute neomycin exposure (~0.02% hit rate), but not continuous gentamicin exposure (Owens et al., 2008). Neomycin and gentamicin activate overlapping, yet distinct, cell death pathways in zebrafish hair cells (Owens et al., 2009; Coffin et al., 2013a,b). Therefore, it was not unexpected that some screens found compounds that protect from one antibiotic or the other, suggesting that these compounds modulate specific intracellular pathways. However, all four compounds from the present screen conferred protection from both neomycin and gentamicin, suggesting that these natural compounds may be broadly applicable to prevent aminoglycoside ototoxicity.

Ber, Her, and Iso are bisbenzylisoquinoline alkaloids from the Thalictrum family (herbaceous perennial flowering plant; Schif, 1991; Pelletier, 1996); Ber, and Iso are also found in Berberis, a shrub (Weber and Fournet, 1989; Di et al., 2003). E6 berbamine is a synthetic derivative of berbamine. None of these natural compounds have been reported to affect hearing, and since they are not well studied, there is little information about bioavailability or serum concentrations in mammals. However, all four natural compounds have been studied to treat various disorders, including cancer and pain; bisbenzylisoquinoline alkaloids have anti-tumor properties (Kuroda et al., 1976) and Ber reduces inflammation and pain responses in mice with serotonin-induced hind paw edema (Küpelí et al., 2002). This research shows the promising applications that natural compounds have in treating a wide variety of medical conditions, and our data add hair cell protection to this growing list of therapeutic possibilities.

Our otoprotective alkaloids did not affect antibiotic efficacy of neomycin or gentamicin, making them ideal for future research in mammalian systems, and ultimately as potential therapeutics to prevent hearing loss in humans. One caveat of using the zebrafish lateral line as an ototoxicity model is the rapid rate of hair cell regeneration, where newly generated hair cells may be confused with protected hair cells. However, aminoglycoside-induced hair cell death occurred within hours of antibiotic application, whereas substantial regeneration is only evident 48 h after hair cell death (Harris et al., 2003; Ma et al., 2008). We did not observe an increase in cell proliferation after co-administration of natural compound and neomycin, consistent with our interpretation that our natural compounds confer true hair cell protection. In addition to determining if Ber, E6, and Iso protect mammalian hair cells, future work will include using the structural similarities of the four natural compounds to create more efficient otoprotective synthetic derivatives. For example, the synthetic berbamine, E6, protects hair cells at a lower concentration than Ber, Iso, or Her. Natural compounds offer a rich source of functionally and chemically diverse otoprotective compounds with the potential to reduce the ototoxic side effect of a highly effective class of antibiotics.

**AUTHOR CONTRIBUTIONS**

Participated in research design: MK, AJO, JGT, ABC. Conducted experiments: MK, RB, AJO, TFS, JGT, ABC. Performed data analysis: MK, JT, AC. Wrote or contributed to the writing of the manuscript: MK, RB, AJO, TFS, JGT, ABC.

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