Cochlear protein biomarkers as potential sites for targeted inner ear drug delivery

James G. Naples 1 · Lauren E. Miller 2 · Andrew Ramsey 3 · Daqing Li 3,4

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Abstract
The delivery of therapies to the cochlea is notoriously challenging. It is an organ protected by a number of barriers that need to be overcome in the drug delivery process. Additionally, there are multiple sites of possible damage within the cochlea. Despite the many potential sites of damage, acquired otologic insults preferentially damage a single location. While progress has been made in techniques for inner ear drug delivery, the current techniques remain non-specific and our ability to deliver therapies in a cell-specific manner are limited. Fortunately, there are proteins specific to various cell-types within the cochlea (e.g., hair cells, spiral ganglion cells, stria vascularis) that function as biomarkers of site-specific damage. These protein biomarkers have potential to serve as targets for cell-specific inner ear drug delivery. In this manuscript, we review the concept of biomarkers and targeted-inner ear drug delivery and the well-characterized protein biomarkers within each of the locations of interest within the cochlea. Our review will focus on targeted drug delivery in the setting of acquired otologic insults (e.g., ototoxicity, noise-induced hearing loss). The goal is not to discuss therapies to treat acquired otologic insults, rather, to establish potential concepts of how to deliver therapies in a targeted, cell-specific manner. Based on our review, it is clear that future of inner ear drug delivery is a discipline filled with potential that will require collaborative efforts among clinicians and scientists to optimize treatment of otologic insults.

Keywords Inner ear · Targeted-drug delivery · Biomarkers · Hearing loss

Introduction
The cochlea is a privileged organ within the temporal bone that is protected from the external environment by the otic capsule bone, the round window membrane (RWM), and blood-labyrinth barrier. There are various cell types within the cochlea, which are particularly sensitive to disruptions of the local environment. Small changes within the local environment of the cochlea can introduce acquired insults, which present as a sensorineural hearing loss (SNHL) in the clinical setting and can be accompanied by tinnitus and vertigo. These characteristics introduce significant challenges when treating acquired cochlear disorders. There are various well-established etiologies of acquired cochlear insult that include excessive noise exposure, ototoxic medications, and idiopathic causes such as Meniere’s disease and sudden SNHL. While each of these various etiologies produces a similar clinical effect of SNHL, there is evidence to suggest that many of the causes of acquired SNHL preferentially introduce insults to specific locations within the cochlea (inner hair cells (IHC) [1], outer hair cells (OHC) [2], stria vascularis (SV) [3], spiral ganglion neurons (SG) [4], and supporting cells [5]). The specificity of ototoxicity is not managed differently in the clinical setting based on location of insult, and our current clinical therapies are non-specific. Despite these challenges, clinical and basic science research has led to early foundational understandings of the various mechanisms by which therapies can penetrate the protective barriers and reach locations within the cochlea.
However, the ability of treatments to target only the specific sites of insult remains elusive and is an area of active research that necessitates identification of target-specific biomarkers.

A biomarker is defined as a biological molecule that can be used as an indicator of disease state. This concept can be applied more specifically in the study of acquired diseases of the cochlea to indicate and identify site-specific insults to the structures within the cochlea. In theory, each of the various cell types within the cochlea (IHC, OHC, SV, SG, and supporting cells) has specific molecules (biomarkers—typically proteins) that differentiate it from other cells. Further, many of these acquired cochlear insults cause preferential injury to specific sites within the cochlea (e.g., cisplatin injury to OHC [7], noise-toxicity to spiral ganglion cells in cochlear synaptopathy [8], loop diuretics to the stria vascularis [9]).

As such, the concept of cochlear biomarkers opens the potential for targeted, cell-specific therapies in the setting of acquired cochlear injuries. While this concept holds great potential to introduce targeted drug delivery to the inner ear, one of the necessary requirements is that the biomarker target has to be accessible for interaction. From a structural standpoint, this suggests that the biomarker has to be located within the cell’s outer membrane with an available extracellular domain for specific interactions with the treatment or delivery vehicle of interest (Fig. 1). Intracellular targets are theoretically available but would possess limited specificity due to potential nonspecific entry into the cell. The ideal components necessary for targeted inner ear drug delivery are listed in Table 1.

In the clinical setting, acquired cochlear insults induce permanent, irreversible SNHL. Unfortunately, by the time these disorders are recognized, subjects have already sustained some cochlear damage. Thus, prevention and early recognition are the only options for avoiding insult. If the disorder is identified in a timely manner, oral or intratympanic (IT) therapies become available, although therapy to reverse these acquired insults is currently limited to corticosteroids [14]. Even if treatment is initiated, the outcomes are highly variable. Two factors that would likely improve preservation of cochlear structures in the setting of acquired insults are (1) rapid initiation of therapies (or pre-treatment in the context of known ototoxic medication administration) and (2) targeted, specific delivery of therapies to inner ear structures. If both factors can be addressed, there is significant potential to improve therapeutic efficiency and hearing outcomes in the clinical setting. The timing of the initiation of therapy is dependent upon patient and clinician recognition of the problem and will not be discussed in this review. Targeted delivery of drugs to specific locations within the cochlea is achievable because of the characterization of potential cell-specific biomarkers, and thus will be the focus of this manuscript.

At this time, there is significant translational animal research exploring the concept of targeted local delivery for cochlear disorders [10, 15]; however, there are no targeted therapies available for treatment of the inner ear in the clinical setting. In this review, we aim to discuss potential inner ear biomarkers that may serve to facilitate more efficient mechanisms of inner ear drug delivery. We will discuss potential biomarkers by cell type and provide context as to which clinical disorder may be addressed by targeted therapeutics to each specific cell type. Furthermore, brief information regarding whether potential ligands are available for targeted binding to the biomarker of interest will be provided. Ultimately, biomarker-ligand interactions have the scientific and clinical potential that in the future may lead to therapeutic options for acquired cochlear insults. While we aim to be comprehensive, limited information is available on many of the potential biomarkers; thus, we will detail specifics where possible, and discuss potential utility in other instances.
Biomarkers of congenital cochlear disorders will not be included in this review as this is outside the scope of the discussion. Targeted therapeutics in the management of congenital hearing loss (e.g., Connexin-26, GJB2, related hearing loss) are not currently available because subjects are often deaf at birth, and it is challenging to administer therapy in a timely manner. Therapeutic agents for acquired cochlear disorders will not be specifically discussed, as the focus will be on the biomarkers that offer potential targets for therapeutic delivery. It is our hope that this review provides the most up-to-date information regarding the concept of cochlear biomarkers and their role as potential targets for specific delivery of drugs to the inner ear. Emphasis will be placed on the significant translational potential for this research.

**Methodology for literature search**

A literature search was performed using the PubMed database. Keywords incorporating each specific cell type of the cochlea were included in our initial search. The cell types we searched were “inner ear,” “outer hair cells,” “inner hair cells,” “spiral ganglion,” and “stria vascularis.” These cochlear cell types were searched with various combinations of the words: “protein,” “target,” and “biomarker.”

Each of the titles and abstracts for these search queries was reviewed by one of the authors. Papers that included proteins primarily responsible for congenital or genetic types of SNHL were excluded from this review. Additionally, any abstracts that were repeated during the search were only included once. The Uniprot (https://www.uniprot.org/) and human protein atlas (www.hprd.org/) databases were then used to determine whether the protein was exposed to the extracellular environment. Papers that had information regarding specific proteins within each cell type were evaluated further to determine whether the protein of interest had been fully characterized. It was noted that many of the proteins identified are found in more than one cell type within the inner ear. In this case, the proteins were characterized, nonetheless, because of their potential utility and availability as biomarkers for targeted drug delivery. In cases where the proteins and genes have more than one name, the name listed on the two protein websites will be used in this review.

Following evaluation within the protein databases, a search for ligand-binding domains and ligands available for each protein was performed (Fig. 2). If a protein was known to have available ligands, they were evaluated for their potential utility to target the drug delivery vehicles to the biomarkers of interest.

**Results**

**Inner and outer hair cells**

Inner and outer hair cells are among the most vulnerable cell types within the inner ear and are often the primarily damaged cell type in the setting of insult. Despite their vulnerability,

| Table 1 Components of an ideal biomarker for targeted drug delivery |
|---------------------------------------------------------------|
| **Ideal situation** | **Challenges** |
| **Accessibility** | 1. Biomarker target is transmembrane protein with an easily accessible extracellular domain |
| | 2. Targeted therapy is delivered prior to or shortly after induced damage |
| | 3. Targeted therapy has properties that allow it to enter the inner ear |
| **Biomarker/ligand interaction** | 1. Many known biomarkers and cellular targets are intracellular, limiting specificity and accessibility for therapeutic use |
| | 2. Delivering therapeutics to the target biomarker prior to significant cellular damage |
| | 3. Many therapies have properties that restrict their entrance to the privileged site of the inner ear [10] |
| **Biomarker specificity** | 1. Few biomarker targets have well-characterized ligand-binding domains |
| | 2. Few compounds are available that bind efficiently to the available domains within biomarker |
| | 3. Many of the potential biomarkers are essential for cochlear function. Ligand-protein interaction may alter cellular function |
| **Analyzability** | 1. Many of the potential targets available are non-specific and exist in various cell types within the cochlea [11] |
| | 1. Few techniques are available that offer analysis of drug delivery in the clinical setting, and are limited to animal models [12] |
| | 2. Clinical outcomes with audiometry are variable and may not reflect what is happening at the cellular level [13] |
they are well characterized both molecularly and functionally, thus offering the realistic potential for specific, targeted delivery to potential hair cell biomarkers. Molecularly, there are a variety of transmembrane protein channels that meet criteria for biomarkers. Functionally, OHCs are largely supportive in function, while IHCs are responsible for signal transduction that converts sound to an electrical signal via its interactions with the cochlear nerve. We will detail some of the well-characterized proteins within the inner and outer hair cells that have significant potential as biomarkers. Additional potential biomarkers are listed in Table 2.

**Hair cell-specific protein biomarkers**

**Prestin**

Prestin is a well-described outer hair cell protein that functions as a cochlear amplifier [32]. Unlike many of the other cochlear biomarkers, prestin is specific to the OHCs and does not exist within any other cell type within the cochlea [25]. This makes it particularly appealing as a biomarker for acquired disorders of the OHCs such as cisplatin ototoxicity which preferentially induces damage to the OHCs.

Currently, research is underway investigating targeted delivery of therapies to the OHCs via prestin. This work began after phage display experiments produced a series of peptides that bound to prestin. These peptides were successfully used in the targeted binding of polymersomes to rat cochlear explants [26]. One of the peptides has subsequently been used in vivo to deliver targeted liposomes containing a JNK inhibitor to outer hair cells in a murine model undergoing a severe acoustic insult. The use of the prestin biomarker enabled the payload to be successfully targeted to the outer hair cells and greatly enhanced the protection of the mice’s hearing from the insult [27].

Unfortunately, prestin expression in outer hair cells is downregulated in cells undergoing kanamycin-induced apoptosis. This may limit its usefulness in the treatment of aminoglycoside-induced apoptosis in animal models if therapy is not introduced early enough in the clinical setting [33].

**Large conductance voltage and calcium-activated potassium (BK) channels**

These protein channels are located in the outer membranes of inner and outer hair cells. These channels play an important role in processing the neural feedback from the brain [18]. Clinically, the BK channel has been investigated as a potential channel involved in a tinnitus pathway [19].

A 9-amino acid peptide has been identified which binds tightly to BK channels [19] and can be used both to modulate BK channel activity and potentially as a targeting peptide. Collectively, this work provides the foundation for the concept...
of BK channels as a potential biomarker available for targeted delivery of therapies aimed at addressing tinnitus.

**Spiral ganglion-specific protein biomarkers**

The spiral ganglion cells and neural tissue within the cochlea are an important site for targeted drug delivery because many acquired cochlear disorders have a primary or secondary effect at this location. Synaptopathy of the neuronal elements within the cochlea is a rapidly growing topic within acoustic research [34], and clinically presentation is variable depending on the severity of damage. The few, well-characterized specific protein biomarker targets within the spiral ganglion cells will be discussed, and the remaining will be listed in Table 3.

**Trisialoganglioside clostridial toxin receptor**

This trisialoganglioside clostridial toxin receptor (GT1b) is expressed by a large number of neuronal cells within the central and peripheral nervous system. Within the cochlea, it is a receptor located in the spiral ganglion neurons [43, 44]. It serves as a potential prototype biomarker for targeted delivery to spiral ganglion cells due to its specificity to this cell type within the cochlea. Additionally, phage display has identified the Tet1 peptide that binds specifically to the receptor [41]. The advantage of this synthetic peptide is that binding to the trisialoganglioside clostridial toxin receptor does not alter cell signaling within the spiral ganglion. The Tet 1 peptide has been conjugated to polymersomes and delivered to the inner ear using a cochleostomy. The targeted polymersomes bound to the auditory nerve and also to, or adjacent to, the spiral ganglion’s NF-200-positive nerve fibers [42]. Clinically, targeted delivery of therapies to this receptor as a biomarker may provide treatment options for neural-based disorders such as cochlear synaptopathy [4] in the setting of noise exposure. Consequently, the Tet1 peptide is currently the most promising technique to deliver therapeutic payloads to the spiral ganglion or auditory nerve.

**BDNF/NT-3 growth factors receptor (TrkB)**

Within the cochlea, BDNF/NT-3 growth factors receptor is another spiral ganglion-specific biomarker protein for which local delivery of targeted therapies offer potential utility. As with the trisialoganglioside clostridial toxin receptor, phage display has been used to develop binding peptides for these receptors. Recent work has demonstrated that nanoparticles targeted with the TrkB binding A371 peptide bind preferentially to the spiral ganglion’s neuronal cells [35], and that this targeting peptide has enabled Rolipram, a phosphodiesterase inhibitor, to be delivered to the mouse spiral ganglia as a potential agent that improves cell survival [35, 45]. Additional work has suggested the possibility of targeted delivery of the TrkB agonist, 7,8-dihydroxyflavone (DHF) as a means for increasing neurite growth in the setting of kainic-acid cochlear damage [46]. Similarly, targeted delivery of

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**Table 2** Potential hair cell-specific biomarker proteins

| Protein                                         | Gene         | Cell type | Function | Targeting | Ref  |
|-------------------------------------------------|--------------|-----------|----------|-----------|------|
| Chloride intracellular channel protein 5        | CLIC5        | OHC, IHC  | Channel  | None      | [16] |
| GABA receptors                                  | GABBR1, GABBR2 | OHC, IHC  | Receptors| None      | [17] |
| Large conductance voltage and calcium activated potassium (BK) channel | KCNMA1        | OHC, IHC  | Channel  | Peptide   | [18, 19] |
| LHFPL tetraspan subfamily member 5 protein      | LHFPL5       | OHC, IHC  | Structural protein | None | [20, 21] |
| Mechanoelectrical transducer (MET) channel      | Various      | OHC, IHC  | Channel  | None      | [22] |
| Otoferlin                                       | OTOF         | IHC       | Synapse protein | None | [23] |
| Phospholipid-transporting ATPase IC             | ATP8B1       | OHC, IHC  | Transport protein | None | [24] |
| Prestin                                         | SLC26A5      | OHC       | Motor protein | Peptide | [25-27] |
| Transmembrane channel-like protein 1            | TMC1         | OHC, IHC  | Channel  | None      | [28, 29] |
| Transmembrane channel-like protein 2            | TMC2         | OHC, IHC  | Channel  | None      | [28] |
| Vesicular glutamate transporter 3               | SLC17A8      | IHC, SG   | Transport protein | None | [30, 31] |

**Table 3** Potential spiral ganglion-specific protein biomarkers

| Protein                                         | Gene | Cell type | Function | Targeting | Ref  |
|-------------------------------------------------|------|-----------|----------|-----------|------|
| BDNF/NT-3 growth factors receptor               | NTRK2| SG        | Receptor | Peptide   | [35, 36] |
| NMDA receptors                                  | Various | IHC ribbon synapses | Synapse protein | Peptide | [37, 38] |
| Substance-P receptor                            | TACR1 | SG        | Receptor | Peptide   | [39, 40] |
| Trisialoganglioside clostridial toxin receptor  | Not Listed | SG   | Receptor | Peptide   | [41, 42] |
neural growth factors such as BDNF [47] may offer the potential for neural regeneration.

**N-methyl-D-aspartate receptors**

These excitatory glutamate receptors are found in a wide range of neural cells that are responsible for opening ion channels and neural signaling. However, in the inner ear, they are found within the synapses of the inner hair cells and thought to play a role in the development of tinnitus [37]. There is significant potential for these receptors as biomarkers for neural tissues at the level of the synapse, and clinical work has already led to the identification of the therapeutic peptide Rapastinel (formally Glyx-13) as a potential therapy for tinnitus [38]. Additional work in animal models has demonstrated ribbon synapse protection from IT application of N-methyl-D-aspartate (NMDA) antagonists [37].

**Stria vascularis-specific protein biomarkers**

**Na, K-ATPase**

This is a transmembrane protein expressed by a wide range of organs. However, in the cochlea, it is expressed principally in the stria vascularis. The function of Na, K-ATPase in the inner ear is to pump potassium ions into the endolymph to maintain the high potassium concentration within the fluid and the endocochlear potential. In other organs, the pNaKtide peptide has demonstrated binding to the Na, K-ATPase, making it highly likely that the peptide will bind to Na, K-ATPase in the stria vascularis [48]. Interestingly, some research has suggested that this pNaKtide does not alter downstream activity [49]. The role of the stria vascularis in maintaining endocochlear potential is vital to the function of the cochlea, and damage to the stria has been implicated in a variety of acquired disorders. Classically, otoxicity induced by loop diuretics affects the stria [50] and disrupts the endocochlear potential. Newer evidence supports the concept that cisplatin may impact the stria [51], although the mechanism of cisplatin’s effect here is not yet understood. Clinically, therapies targeted at stria vascularis-specific biomarkers may offer options for prevention of ototoxicity induced by loop diuretics and/or cisplatin.

In addition to the Na, K-ATPase, the Na, K, Cl Co-Transporter, and Barttin proteins are specific to the stria vascularis and may potentially serve as biomarkers, although targeting peptides are currently not available (Table 4).

**Non-specific protein biomarkers**

Ideally, a biomarker for targeted delivery of therapies should be specific to a particular cell type and location within the cochlea. Nonetheless, the number of protein biomarkers expressed in more than one cell type exceeds the number of cell-specific protein biomarkers. Although non-specific proteins may not be as useful as the cell-specific biomarkers, they can play important roles in the delivery of therapeutics to the inner ear. For example, these proteins can be useful to enhance the delivery of materials to the appropriate inner ear sites in instances where multiple cell types have suffered an insult. Alternatively, they may offer non-specific delivery of therapies to the inner ear cells for which there is currently not a targeting peptide and in scenarios where some delivery is more advantageous than no delivery at all (Table 5). Several non-specific biomarkers are briefly described below.

**Calcium channels**

Calcium channels are ubiquitous within the various cochlear cells. The ability to limit cellular calcium uptake in cochlear cells that have recently experienced an insult has the potential to reduce the number of cells that undergo apoptosis and consequently lessen the hearing deficit induced by the insult [83–85]. L- and T-type calcium channels are the two major forms of calcium channels within the cell [67]. Several natural peptides have been identified that target these calcium channels [68, 74]. In addition to these natural peptides, there is also a range of synthetic calcium channel binding compounds that have been developed by the pharmaceutical industry that may be used as targeting compounds.

**Neural growth factor receptors**

The tyrosine kinase receptor A (TrkA) (gene NTRK1) is a receptor involved in signaling responsible for neural growth and differentiation. Although it is similar in function to TrkB, TrkB is activated by BDNF and is specific to neural tissue, while TrkA is activated by neural growth factor (NGF), and the TrkA receptor is located throughout various cell types within the organ of Corti [65]. The non-specific location of

| Protein                | Gene | Cell type | Function             | Targeting | Ref       |
|------------------------|------|-----------|----------------------|-----------|-----------|
| Barttin                | BSND | SV        | Ion channel subunit  | None      | [52, 53]  |
| Na, K-ATPase           | Multiple Genes | SV | channel              | Peptide   | [48, 54]  |
| Na, K, Cl Co-transporter | NKCC1 | SV | Transport protein    | None      | [55, 56]  |
this receptor suggests that the NGF ligand exerts its neurotrophic effects at sites outside of the neural tissue. An additional advantage of this receptor is that peptide fragments of the ligand have been demonstrated to bind to the receptor and could be used to target the protein [66].

### Serine protease proteins

Type II transmembrane serine protease (TMPRSS1/hepsin) (Gene HPN) is expressed in the spiral ganglion and the stria vascularis [78]. Outside the cochlea, the protein acts as a prostate cancer biomarker, and small molecules that bind to the proteins have been identified. These compounds could be used to target the spiral ganglion and the stria vascularis [79].

### Discussion

There are a wide range of cochlear insults that present with permanent SNHL, tinnitus, and/or vertigo. Many of the insults that result in damage have location-specific sites of injury within the cochlea. Despite the fact that many of the locations and mechanisms of these insults have been established, clinical medicine has been unable to reliably translate this work to specifically target therapies to these sites of injury. In this review, we present an overview of potential cell-specific biomarker proteins within the cochlea as a way for scientists and clinicians to logically think about targeted inner ear delivery. Our goal with this review was to offer a more specific thought process and approach to addressing acquired forms of inner ear damage by suggesting available protein targets for therapeutic delivery to each of the various cell types within the cochlea.

The concept of targeted, location-specific delivery and biomarkers for each of the cell types within the cochlea is a challenging order for scientists and clinicians to address. The privileged nature of the cochlea and its sensitivity to minor changes in the local environment create a logistical challenge of delivering therapies to the correct location without inducing more damage. It is likely necessary to optimize local delivery of therapeutics to the cochlear cells using a variety of techniques that include intratympanic (IT) delivery [86], direct round window membrane (RWM) delivery [27], and intracochlear delivery [87]. Clinically, only IT delivery is routinely used; however, the efficiency of this technique varies based on a number of drug- and patient-related factors. There

| Table 5 | Potential non-specific protein biomarkers |
|---------|------------------------------------------|
| Protein | Gene          | Cell type                             | Function     | Targeting | Ref     |
| Aquaporin 4 | AQP4          | Deiters’ and Hansen’s cells           | Channel      | None      | [57]    |
| ATP-sensitive inward rectifier K channel 10 | KCNJ10        | SV, SG Deiters’                       | Channel      | None      | [58, 59]|
| Basigin  | BSG           | SV, auditory nerve                    | Maintenance protein | None      | [60]    |
| CD44 antigen | CD44          | Outer pillar and Claudius cells       | Hyaluronic acid receptor | None      | [61]    |
| Clarin 1  | CLRN1         | OHC, IHC, ‘SG’                        | Structural protein | None      | [62, 63]|
| Fibroblast growth factor receptor 3 | FGFR3         | OHC, pillar and Deiters’ Cells        | Growth factor | None      | [64]    |
| High affinity nerve growth factor receptor | NTRK1         | OHC, IHC, ‘SG’ supporting cell        | Receptors    | Peptide   | [65, 66]|
| L-type calcium channel | CACNA1C,CACNB1,OTOA | Ubiquitous                           | Calcium channel | Peptides, Small Molecules | [67–69]|
| Otoancorin | SLC26A4       | Spiral limbus, tectorial membrane,    | Anchoring protein | None      | [70]    |
| Pendrin   | SYP           | Sulcus, spiral prominence             | Ion transport | None      | [71, 72]|
| Synaptophysin | SYP           | OHC, IHC, ‘SG’                        | Synapse protein | None      | [73]    |
| T-type calcium channel | CACNA1H,CACNA1G | Ubiquitous                           | Calcium channel | Peptides, Small Molecules | [67, 74]|
| Transient receptor potential cation channel subfamily V member 4 | TRPV4        | SG,SV                                 | Calcium channel | Small molecules | [75–77]|
| Type II transmembrane serine protease | HPN           | Support cells                         | Protease     | Small molecule | [78–80]|
| Vang-like protein 2 | VANGL2        | OHC, IHC, supporting cells            | Anchoring protein | None      | [81]    |
| Zinc transporter ZIP8 | SLC39A8       | Mainly OHC, IHC, SV, supporting cells | Transport protein | None      | [82]    |
| Zinc transporter ZIP14 | SLC39A14      | Mainly OHC, IHC, SV, supporting cells | Transport protein | None      | [82]    |
has been an explosion of research evaluating novel vehicles for therapeutic delivery to the inner ear such as liposomes, hydrogels, and nanoparticles [27, 88–90], and ultimately, identifying the optimal delivery vehicle will improve the yield on our ability to get targeted therapies to the specific bio-markers of interest.

Once a delivery vehicle is optimized, ligands that specifically bind to the biomarker proteins of interest need to be developed. Some of the potential biomarkers within the various cell types have available ligands (see tables); however, many do not. It is reasonable to predict that the next revolution in inner ear drug delivery research may be focused on the production of ligands for proteins that are specific to the various cell types within the cochlea. The clinical implications of cell-specific ligands for targeted delivery of therapies to the inner ear cannot be understated. Currently, non-specific therapies such as corticosteroids are the only available clinical treatment in use to address otologic insults. The mechanism of corticosteroid activity within the inner ear is complex [86, 91, 92] and likely has non-specific interactions that are beyond what we can currently measure. If ligand specificity is developed and refined, this concept would offer the potential to treat various forms of otologic insult and hearing loss in a specific manner. This concept could just as easily be applied to the spiral ganglion cells in the treatment of various causes of cochlear synaptopathy and stria vascularis for loop diuretics.

It warrants mentioning that this review has focused on the use of peptides to target the biomarkers rather than pharmaceutical products (e.g., FDA-approved therapies, commercially available drugs). While pharmaceutical products can bind to the biomarkers as tightly and with the same specificity as peptides, they can be potentially much more challenging to conjugate to nanoparticles containing the appropriate therapeutic payload. The methodology to conjugate peptides to the nanoparticle’s coat is well-established and relatively straightforward to perform [94]. Although the methodology to conjugate pharmaceutical compounds to nanoparticles is possible, it is not as widely used, and largely limited to applications in oncology [95].

The simplicity of the concept of targeted delivery to cell-specific biomarkers is currently overshadowed by the complexity of execution. There are layers of complexity to each step of this process that necessitate discussion. From a clinical perspective, it is a risk to initiate treatment on subjects who have any residual cochlear function because the cochlear cells are particularly sensitive to changes within the local environment [96, 97]. Thus, many subjects with acquired forms of otologic insult and some remaining natural hearing are unlikely to be candidates in clinical trials for the initiation of targeted treatments [98]. Next, finding the appropriate combination of delivery vehicle, ligand, and therapy can be an overwhelming task. While there are a handful of vehicles for drug delivery (i.e., hydrogels, liposomes, etc.), the number of potential ligands is extensive when both natural and synthetic ligands are accounted for. Even if the correct combination of vehicle and
ligand is discovered, there is potential that conjugating the vehicle, ligand, and therapy may compromise the function or binding efficiency to the target. Similarly, science has not achieved reliable success in determining therapies that preserve cochlear function or reverse cochlear insult. Thus, it seems that the application of these concepts is not as close as many think. Nonetheless, there is potential to apply some of these concepts to hearing preservation during the introduction of a cochlear implant (CI) [99]. As indications for CI expand to include subjects with residual hearing, it offers a realistic opportunity to introduce these concepts to subjects that have little cochlear reserve, and thus, little to lose. In fact, some research is already evaluating the role of BDNF and cochlear implant for SGN survival [100, 101].

In addition to the challenges listed above, the physiological response to cellular insult introduces additional challenges. The first challenge is that response to insult and delivery of therapy is time sensitive. For the treating clinician, this represents a logistical challenge of getting subjects treated before irreversible damage develops. This typically occurs within the first few days after insult [102]. Similarly, cells within the cochlea respond to insult by altering their homeostasis, which manifests as a change in protein production and function. For example, prestin expression in outer hair cells is downregulated in cells undergoing kanamycin-induced apoptosis [33]. Thus, despite the theoretical utility of prestin as an OHCh-specific protein biomarker, its utility may be limited in the setting of aminoglycoside-induced apoptosis in both animal models and in the clinical setting if therapy is not introduced early enough [33]. Alternatively, proteins can be upregulated which may introduce new biomarker targets, as in the setting of cisplatin ototoxicity [103]. These concepts will likely warrant consideration as the concept of biomarker targeting is refined.

Another consideration with this concept is that we are currently only able to present this treatment approach for acquired forms of otologic insults in the setting of normal structure and function of the cochlea prior to insult. Many of the well-characterized forms of hearing loss are congenital or present at birth [104], which may eliminate the potential for non-gene-based therapies to reverse the process. Fortunately, there is currently a significant amount of animal research being undertaken that explores treatment and potential therapies of congenital and syndromic forms of hearing loss [105, 106]. Future considerations should aim at targeting specific sites of dysfunction in these genetic forms of hearing loss, particularly as gene therapies become available [107].

Ideally, the binding of the targeting peptide to its biomarker should not affect the biomarker’s cellular function. However, as the peptides are expected to target outer membrane proteins that are exposed to the extracellular environment, it is highly likely that many of the targeted biomarkers will be receptors or channel proteins. In such cases, the targeting peptides may act as an agonist or antagonist modulating cellular function [108]. For instance, care has to be taken employing the pNaKtide peptide to target the stria vascularis as the protein acts as an agonist modifying Na/K-ATPase-mediated amplification of ROS signaling [48, 109]. This concern of altered cellular function has previously been demonstrated where analogs of NGF bind TrkA and inhibit neurite outgrowth [66]. Refining and understanding ligand-target interactions will be necessary to avoid these unwanted interactions as progress is made.

This review has identified a large number of proteins that have the potential to act as biomarkers. Unfortunately, ligands and targeting peptides are not available for the majority of these proteins. This is an indication that the study of targeted delivery of therapeutic material to inner ear cells is in its infancy, and the future for targeted nanoparticles to deliver therapies to the inner ear holds great potential. As more work is accomplished, this technology should also enable genetic material to be delivered to cells in order to treat patients with genetically induced progressive hearing loss in addition to the acquired forms of hearing loss that we discuss.

Overall, cell-specific targeting for inner ear drug delivery would fulfill the following criteria: (1) a protein biomarker that is specific to a single cell type (IHC, OHC, SV, SG, supporting cells) should be available; (2) this specific biomarker should have an accessible extracellular domain; (3) a ligand that interacts with the cell-specific biomarker protein should be available; and (4) the ligand-protein interaction should lead to internalization of the therapy for downstream action. These ideal characteristics are summarized in Fig. 3. Despite the challenges that lie ahead in the evolution of targeted delivery of therapies for acquired inner ear disorders, progress is occurring. Our understanding of the function and structure of the cells within the inner ear has evolved tremendously within the last couple of decades. Biomolecular research has elucidated the function of many of the proteins that we present here as biomarkers and targets for therapeutic delivery. It is conceivable that ligand discovery is not far behind. The essential component to all of this work will be the collaborative efforts on the part of clinicians and scientists to make this concept a reality. Clinicians need to remain unsatisfied with the status quo of otologic therapies, and scientists need to continue to explore applications of their work to widespread burden of otologic disorders. This idea presents an opportunity to apply the discoveries of basic science to the treatment of otologic disease. Importantly, it also represents an opportunity to begin the next phase of inner ear therapeutics that addresses the shortcoming of previous generations of treatments and combines targeted, specific technology as medicine moves into an era of personalized medicine [110].

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The authors declare that they have no conflict of interest.

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