Teprotumumab-associated chronic hearing loss screening and proposed treatments

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SUMMARY
We report a case of a woman in her 50s with chronic teprotumumab-associated sensorineural hearing loss. The patient presented with chronic thyroid eye disease with proptosis and diplopia despite systemic thyroid control and orbital decompression. She was started on teprotumumab but developed tinnitus after the third dose, followed by frank hearing loss after the fifth dose. Her audiogram showed bilateral mild to moderate-severe hearing loss, which was significantly worse compared with her baseline audiogram obtained prior to treatment. Teprotumumab was immediately stopped, however repeat audiogram 6 weeks later showed no improvement. Given potentially irreversible sensorineural hearing loss, we recommend close monitoring with regular audiometric testing before, during and after teprotumumab therapy and propose potential treatment to reverse its effects in the ear.

BACKGROUND
Thyroid eye disease, also known as Graves’ ophthalmopathy, is an orbital autoimmune condition associated with inflammation, tissue fibrosis, infiltration and expansion. Signs of thyroid eye disease include proptosis, lagophthalmos and strabismus, which can lead to debilitation, disfigurement, and in the worst cases permanent visual loss.

The pathogenesis of thyroid eye disease is not fully understood, however there is growing evidence that increased activity of insulin-like growth factor-1 receptor (IGF-1R) is involved. Patients with active disease demonstrate elevated IGF-1R expression in orbital fibroblasts as well as activating autoantibodies targeting these receptors. 1 Teprotumumab (Horizon Therapeutics, Dublin, Ireland) is a human monoclonal antibody with activity against IGF-1R. It is the first drug approved by the US Food and Drug Administration for the treatment of thyroid eye disease. Clinical studies have found teprotumumab effective for improving proptosis, diplopia, clinical activity score and quality of life measures. 2–4 Insulin-like growth factor-1 (IGF-1) is a ubiquitous molecule found throughout the body playing various roles in multiple systems. Consequently, side effects involving virtually every body system have been reported with teprotumumab use, including muscle spasms, hearing loss, hyperglycemia and inflammatory bowel disease. 2–5,8–9

Notably, the Phase III Horizon clinical study reported subjective hearing loss in 10% of patients that was purportedly reversible without intervention; however this was not substantiated by audiometric testing. 2 Subsequent literature has found the incidence of otological side effects to be as high as 30%. 7 One clinical study reported borderline audiometry in two patients with tinnitus, however baseline pretreatment audiometry was not obtained for comparison. 9 Multiple published cases of teprotumumab-associated hearing loss objectively demonstrated persistent and potentially permanent sensorineural hearing loss. 10–12 IGF-1 plays a significant role in the development, maintenance and protection of hearing function within the inner ear. 13 The inhibition of the IGF-1 signalling pathway by teprotumumab would explain its association with sensorineural hearing loss.

Here, we present a case of sensorineural hearing loss after partial teprotumumab therapy. We recommend practice guidelines for patient screening and suggest potential therapeutic modalities for teprotumumab-associated hearing loss.

CASE PRESENTATION
A woman in her 50s with a history of chronic Graves’ disease presented with persistent diplopia and proptosis despite orbital decompression. Her exam was notable for extracocular muscle restriction in up gaze, right proptosis and right hypoglossus. Her thyroid stimulating immunoglobulin was stable but elevated at 431 (normal <140). Her thyroid stimulating hormone was 1.11 ulU/mL (normal 0.40–4.50 ulU/mL), free T4 was 1.00 ng/dL (normal 0.7–1.7 ng/dL) and free T3 was 2.89 pg/mL (normal 2.3–4.2 pg/mL). The patient was on methimazole 2.5 mg daily. MRI of the head and orbits with and without contrast showed marked right greater than left enlargement of the medial and inferior rectus muscles and proptosis consistent with thyroid ophthalmopathy. There was bony evidence of the prior orbital decompression of the medial wall and floor.

After considerable discussion, the patient was started on teprotumumab with the plan to receive an initial dose of 10 mg/kg, followed by 20 mg/kg every 3 weeks to complete a total of eight infusions.

INVESTIGATIONS
Prior to her first dose of teprotumumab, the patient underwent baseline otological and audiological evaluation, which showed a normal otological examination and normal tympanogram. Baseline audiometry demonstrated mild hearing loss at 4000–8000 Hz bilaterally (normal hearing threshold -250–3000 Hz) v consistent with normal age-related hearing loss (figure 1). The right ear was slightly worse than the left ear by 10 dB at 250–1000 Hz. Of note, the patient had a history of...
benign paroxysmal positional vertigo, she reported that her right ear had not felt the same since her first episode several years ago. After the third infusion, the patient reported intermittent ringing in her ears. On the fifth dose, she had noticeable hearing loss. Repeat audiogram revealed bilateral mild to moderate-severe down sloping sensorineural hearing loss with a 15–30 dB decline at 500–8000 Hz compared with her baseline (figure 2).

**OUTCOME AND FOLLOW-UP**

After discussion of these findings with the patient and her otolaryngologist, the decision was made to stop teprotumumab, forgoing the remaining doses to avoid potential dose-dependent ototoxicity. Repeat audiogram 6 weeks after cessation of teprotumumab showed no improvement (figure 3). The patient continues to be followed closely by her otolaryngologist with plans to consider hearing aids should there be no improvement after 3 months.

**DISCUSSION**

This case describes a patient who objectively developed chronic sensorineural hearing loss after partial teprotumumab therapy for the treatment of thyroid eye disease.

Teprotumumab is a biological agent that inhibits the IGF-1 pathway by targeting the IGF-1R. IGF-1 is a neuroprotective agent that regulates the development and maintenance of the inner ear. It participates in cellular metabolism, growth, proliferation, differentiation and prevention of cell death of hair cells. IGF-1 deficiency is associated with profound sensorineural hearing loss in mice models and humans.\(^\text{13-15}\)

Objective teprotumumab-associated chronic sensorineural hearing loss has been documented in the literature. In all cases, audiometry obtained following the initiation of treatment was performed only after the onset of subjective symptoms of hearing impairment, which varied from the third to the fifth dose. The timing of audiometric testing after the development of symptoms also varied widely from immediately to months later when symptoms persisted or worsened with subsequent infusions.\(^\text{11-12}\)

The magnitude of hearing loss in these cases was typically mild to moderate-severe, with no improvement on audiometry several months after discontinuation of therapy.\(^\text{10,11}\)

**Patient screening and monitoring**

Given the findings above, it is imperative to counsel patients on the risk of potentially irreversible sensorineural hearing loss prior to the initiation of teprotumumab. Risk factors should be considered such as history of otological insult, including age-related hearing loss, loud noise exposure and other conditions that degrade baseline otological function. Based on these factors, patients in collaboration with clinical providers may wish to defer or elect alternative treatments. A multidisciplinary team involving ophthalmology, otolaryngology, audiology and endocrinology is important for the care of patients with thyroid eye disease.
disease considering teprotumumab therapy. Baseline otological examination and audiometry should be established prior to treatment to identify susceptible patients and detect changes associated with drug therapy. This practice is standard with other medications with known ototoxicity potential such as cisplatin.17

The relationship between serum IGF-1 level and hyperthyroidism is unclear. Several studies suggest hyperthyroidism is associated with elevated serum IGF-1, while other studies found no difference in serum IGF-1 between hyperthyroid and euthyroid patients.18–21 In some animal models, low serum IGF-1 was associated with greater severity of hyperthyroid disease.22 Interestingly, one study found 18.4% of patients with newly diagnosed Graves’ disease had IGF-1 deficiency. Low IGF-1 was associated with higher thyroglobulin level and more severe free T3 hyperthyroidism at diagnosis.23 Serum IGF-1 testing may thus be useful to identify individuals predisposed to teprotumumab-associated hearing loss in the setting of a normal baseline audiogram. Interestingly, serum IGF-1 in our patient was elevated at 501 ng/mL (normal 37–208 ng/mL) 6 weeks after stopping teprotumumab, suggesting that elevated IGF-1 may not be protective either.

The published cases of teprotumumab-associated hearing loss performed audiograms after patients reported symptoms of hearing loss.10,11 Given that subjective hearing loss does not reliably reflect objective hearing loss, the onset of otological dysfunction in relation to drug dose and duration is unclear.12 In addition to baseline testing, we propose regular surveillance audiometric testing ideally after every infusion throughout the therapeutic course and after drug cessation. This degree of monitoring may not be practical but does provide the ability to identify early, subclinical hearing impairment. It may also help characterize patterns of dose-dependent toxicity. We recommend at a minimum audiology testing pre, mid and post treatment.

Patients taking teprotumumab should be counselled and monitored closely for symptoms of early hearing dysfunction (ear fullness, muffled sounds, tinnitus, autophony, hypoacusis). Ototoxic medications and loud noise exposure should also be avoided.24

Potential treatment
Sensorineural hearing loss is primarily caused by damage or dysfunction of cochlear hair cells, which have little ability to proliferate or regenerate in postnatal mammalians. Systemic corticosteroids with or without local intratympanic corticosteroid injection has traditionally been used to treat idiopathic sudden onset sensorineural hearing loss, with variable success.25–26 Unfortunately, steroids would not be expected to help teprotumumab-associated hearing loss since its mechanism is not inflammatory. Oral steroids were used in one reported case of teprotumumab-associated hearing loss but were ineffective.10 Hearing aids and cochlear implantation are options to improve functional hearing but do not address the underlying

Figure 2 Audiogram after the fifth dose of teprotumumab demonstrates bilateral mild to moderate-severe down sloping sensorineural hearing loss with a 15–30 dB decline at 500–8000 Hz compared with baseline.
Local IGF-1 therapy has shown promising results in the treatment of refractory sensorineural hearing loss but is not yet currently widely used. It is given intratympanically and diffuses through the round window membrane to enter the inner ear (figure 4). Risks of any intratympanic injection include pain, headache, vertigo, nausea, infection and tympanic membrane perforation.27–31 This targeted therapy should be considered for treatment of teprotumumab-associated sensorineural hearing loss. Acting on the same signalling pathway in a timely manner could theoretically negate the local effects of teprotumumab in the ear. This could potentially prevent or reverse teprotumumab-associated sensorineural hearing loss. Prophylactic treatment in susceptible patients could also be considered.

Stable drug delivery to the inner ear remains a challenge. Systemic IGF-1 is not well tolerated and is unlikely to pass through the blood-labyrinth barrier.12 External topical application such as ear drops would theoretically be the safest and least invasive route of delivery. However, this is significantly limited by the impermeable nature of the tympanic membrane which separates the outer and middle ear. Small peptide-linked particles have been shown to be actively transported across the intact tympanic membrane in in vivo rat models.33 Chemical permeation enhancers (bupivacaine, limonene, sodium dodecyl sulfate) may also aid increased drug permeability across the tympanic membrane, while hydrogel formulations maintain direct contact for maximal delivery.34 A chemical permeation enhanced hydrogel formulation of IGF-1 applied to tympanic membrane in the external ear could be a novel treatment strategy for teprotumumab-associated hearing loss. Drug delivery to the middle ear via the eustachian tube could also be considered.35 36 Currently, intratympanic injection remains the most widely used strategy for drug delivery to the inner ear.

Additionally, entry into the inner ear is limited by the permeability of the round window membrane. Several strategies have been employed to increase permeability including ultrasound microbubbles and drug carriers.37 Combined with these strategies to increase penetration into the inner ear, local application of topical IGF-1 could potentially be beneficial for patients with teprotumumab-associated hearing loss specifically.
Alternatives

Novel agents continue to be investigated as an alternative to teprotumumab for the treatment of thyroid eye disease. A low volume subcutaneous formulation of IGF-1R antagonist is currently in development, which could significantly improve convenience and cost of treatment. The longer half-life and greater stability of this agent would allow for fewer or less frequent dosing. \(^{19}\) \(^{35}\) Stable serum concentration would also minimize peaks and troughs, limiting drug exposure and dose-dependent toxicity to other organ systems. This is particularly important for areas with limited regenerative potential such as the inner ear.

In summary, teprotumumab has been associated with sensori-neural hearing loss, through its antagonistic action on the IGF-1 signalling pathway, which engages in otological health. Further investigation is needed to better characterize teprotumumab-associated hearing loss, including its risk factors, frequency, magnitude and reversibility in relation to individual susceptibility, drug dosage and duration of treatment. This would potentially help establish safer, clearer guidelines and eligibility criteria for teprotumumab therapy.

Learning points

- Teprotumumab is associated with potentially irreversible sensorineural hearing loss.
- Careful patient selection and counselling regarding these risks is imperative. Potential risk factors should be assessed including age, prior otological history and serum insulin-like growth factor-1 (IGF-1).
- We recommend regular otological examination and audiometric testing before treatment, after each infusion, and after treatment, as well as close monitoring of hearing loss symptoms.
- Topical IGF-1 therapy may be helpful in treating teprotumumab-associated hearing loss, however this is still undergoing evaluation.

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Case reports provide a valuable learning resource for the scientific community and can indicate areas of interest for future research. They should not be used in isolation to guide treatment choices or public health policy.

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REFERENCES


5. Amanika L, Sears CM, Claus K. Teprotumumab-Related adverse events in thyroid eye disease. Poster presented at: the American Society of ophthalmic plastic and reconstructive surgery 52nd annual fall scientific symposium; November 11–12, 2021; New Orleans, Louisiana, United States.

6. Chiu CA, Rechel BE, Freitag SK. Understanding the side effect profile of Teprotumumab. poster presented at: the American Society of ophthalmic plastic and reconstructive surgery 52nd annual fall scientific symposium; November 11–12, 2021; new Orleans, Louisiana, United States.


38 Viridian Therapeutics, Inc. VRDN-002, a second-generation insulin-like growth factor-1 receptor (IGF-1R) antagonist antibody for thyroid eye disease: preclinical pharmacokinetic profile and clinical promise. Presented at: the 90th annual meeting of the American thyroid association. Virtual event; September 30, 2021, 2021.