

Use of Adalimumab in Treatment of Autoimmune Sensorineural Hearing Loss: A Word of Caution

To the Editor:

We read with interest the report of Morovic Vergles, *et al* outlining a case of autoimmune sensorineural hearing loss and rheumatoid arthritis (RA) treated successfully with adalimumab¹. Infliximab has also been reported to be effective in the treatment of autoimmune sensorineural hearing loss². Etanercept has had contrasting results in this setting, mostly negative^{3,4}. We recently encountered 2 cases of sensorineural hearing loss temporally related to adalimumab. These are the first such cases in the literature that we know of. There has been 1 previous account of acquired sensorineural hearing loss in the setting of etanercept usage that was judged unlikely to be related to etanercept⁵. We counsel caution in the usage of anti-tumor necrosis factor (anti-TNF) agents in this setting until further studies are undertaken.

The first case was a 44-year-old woman with a 5-year history of refractory seronegative arthritis who had little response to methotrexate or leflunomide. She had an anaphylactic reaction to infliximab after 1 year of usage. She began taking adalimumab in March 2003. In late 2003 she developed sensorineural hearing loss in her left ear and was reviewed by the otorhinolaryngology service. A magnetic resonance image (MRI) of her brain was normal; specifically, there was no evidence of acoustic neuroma. She was taking concurrent leflunomide treatment. Anti-TNF treatment was stopped; she has had no recovery of hearing and remains deaf in her left ear.

The second case was a 34-year-old woman diagnosed with inflammatory spondyloarthritis. She had a poor response to methotrexate and sulfasalazine. She had an excellent response after taking adalimumab, but developed left-side hearing loss identified after the fifth injection. This was confirmed as unilateral sensorineural hearing loss following otorhinolaryngology review and brain MRI. Adalimumab was stopped; there was no improvement in hearing over the first 2 months following cessation, but over the subsequent 2 months her hearing improved to normal.

Anti-TNF agents may be associated with the development of sensorineural hearing loss. This hearing loss may not be reversible in all cases.

However, autoimmune hearing loss has been reported to occur in association with autoimmune diseases such as RA^{1,6}. The sensorineural hearing loss in our cases may represent cases of autoimmune hearing loss in association with inflammatory arthritis but this is usually bilateral. Clinicians should consider this as a rare cause of unexplained hearing loss in a patient receiving an anti-TNF drug.

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