

Teprotumumab bests placebo for active thyroid eye disease

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Patient's eye before/after Tepezza treatment
Photos: Horizon Therapeutics

change in proptosis, diplopia response, and mean change in the Graves' ophthalmopathy-specific quality-of-life overall score. In six patients in the teprotumumab group, reductions in extraocular muscle, orbital fat volume, or both were seen. Adverse events were mostly mild or moderate in severity; in the teprotumumab group, two serious events occurred.

"The effect of teprotumumab was rapid for each outcome, which was evident at the first postbaseline assessment at week six, and the outcomes continued to improve over the 24-week treatment period," the authors write.

Several authors disclosed financial ties to biopharmaceutical companies, including Horizon Therapeutics, which manufactures teprotumumab and funded the study.

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(HealthDay)—Teprotumumab is associated with better outcomes than placebo among patients with active thyroid eye disease, according to a study published in the Jan. 23 issue of the *New England Journal of Medicine*.

Raymond S. Douglas, M.D., Ph.D., from the Cedars-Sinai Medical Center in Los Angeles, and colleagues conducted a randomized phase 3 trial involving [patients](#) with active thyroid eye disease. Patients were randomly assigned to receive either intravenous infusions of the insulin-like growth factor I receptor inhibitor teprotumumab or placebo every three weeks for 21 weeks (41 and 42 patients, respectively).

The researchers found that the percentage of patients with a proptosis [response](#) at week 24 was higher for teprotumumab than placebo (83 versus 10 percent), with a number needed to treat of 1.36. All secondary outcomes were significantly better with teprotumumab than [placebo](#), including overall response, Clinical Activity Score of 0 or 1, mean

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