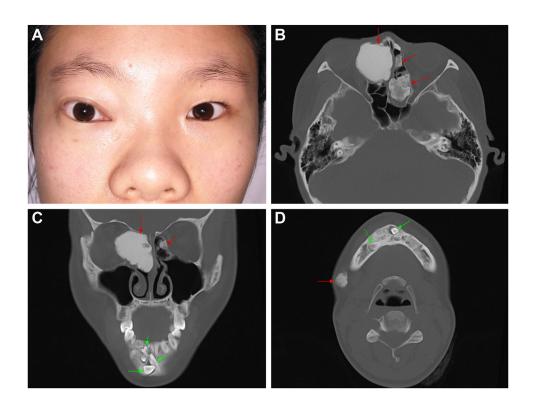
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Pictures & Perspectives



Atypical Gardner's Syndrome with Proptosis as the Primary Symptom

A 20-year-old woman presented with a 3-year history of progressive proptosis of the right eye (A). Her father had a history of multiple osteomas. Congenital hypertrophy of the retinal pigment epithelial was not found on retinal examination, except for retinal folds. Multiple well-circumscribed masses of high radiodensity (B-D), and multiple unerupted supernumerary teeth (C-D) were observed on computed tomography. Gardner's syndrome was diagnosed after these findings were assessed. The osteomas of the right ethmoidal sinus were removed due to the threat of the intraorbital structure. She was advised to follow up regularly to detect the colonic polyps and extraintestinal tumors (Magnified version of Figure A-D is available online at www.aaojournal.org).

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