

**K810703 EXMOOR AURAL GROMMET AG/T2**Apr 14, 1981  
29 days to decisionK810703 · Product code: **ETD** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k810703/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tube, Tympanostomy (ETD)
Date received	Mar 16, 1981
Decision date	Apr 14, 1981
Days to decision	29 days
Third-party review	No

**APPLICANT**

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Company	<b>Exmoor Plastics , Ltd.</b>
Location	Walker, MI, US
510(k) history	43 submissions · 43 cleared · 1981-2010

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Device record: <https://www.510kdatabase.net/k810703/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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