

K810702 EXMOOR AURAL GROMMET AG/T2LApr 14, 1981
29 days to decisionK810702 · Product code: **ETD** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k810702/>**SUBMISSION DETAILS**

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

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

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Device record: <https://www.510kdatabase.net/k810702/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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