FDA 510(k) Premarket Notification

K810702 EXMOOR AURAL GROMMET AG/T2L

CLEARED

Apr 14, 1981
29 days to decision

K810702 · Product code: **ETD** · Ear, Nose, Throat
Source: 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