FDA 510(k) Premarket Notification

K790064 TINNITUS DEVICES

CLEARED

Feb 16, 1979
32 days to decision

K790064 · Product code: **ETS** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k790064/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Generator, Electronic Noise (for Audiometric Testing) (ETS)
Date received	Jan 15, 1979
Decision date	Feb 16, 1979
Days to decision	32 days
Third-party review	No

APPLICANT

Company	**Vicon Instrument Co.**
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1976-1981