FDA 510(k) Premarket Notification

K840359 POST-AURICULAR HEARING AID

CLEARED

Feb 21, 1984
26 days to decision

K840359 · Product code: **ESD** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k840359/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Jan 26, 1984
Decision date	Feb 21, 1984
Days to decision	26 days
Third-party review	No

APPLICANT

Company	**Phonic Ear, Inc.**
Location	Mchenry, IL, US
510(k) history	31 submissions · 31 cleared · 1982-2004