FDA 510(k) Premarket Notification

K863787 PA EQUALIZER

CLEARED

Oct 14, 1986
18 days to decision

K863787 · Product code: **ESD** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k863787/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Sep 26, 1986
Decision date	Oct 14, 1986
Days to decision	18 days
Third-party review	No

APPLICANT

Company	**Electone, Inc.**
Location	Mchenry, IL, US
Contact	MICHAEL D UNDERHILL
510(k) history	73 submissions · 73 cleared · 1981-1997

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k863787/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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