FDA 510(k) Premarket Notification

K884871 RESOUND HES, DIGITALLY PROGR., ITE HEARING AID

CLEARED

Feb 22, 1989
93 days to decision

K884871 · Product code: **ESD** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k884871/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Nov 21, 1988
Decision date	Feb 22, 1989
Days to decision	93 days
Third-party review	No

APPLICANT

Company	**Resound Corp.**
Location	Redwood City, CA, US
Contact	VINCENT PLUVINAGE
510(k) history	41 submissions · 41 cleared · 1989-1999

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k884871/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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