

K960229 RESOUND REAL EAR LOUDNESS MAPPING (RELM) SYSTEM

Mar 7, 1996
50 days to decision

K960229 · Product code: **ETW** · Ear, Nose, Throat
Source: <https://www.510kdatabase.net/k960229/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Calibrator, Hearing Aid / Earphone And Analysis Systems (ETW)
Date received	Jan 17, 1996
Decision date	Mar 7, 1996
Days to decision	50 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Resound Corp.
Location	Redwood City, CA, US
Contact	KRISTA M BUCKLES
510(k) history	41 submissions · 41 cleared · 1989-1999

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k960229/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 4, 2026