

K931372 RESOUND PORTABLE PRESCRIPTIVE PROGRAMMING SYSTEMNov 3, 1993
231 days to decisionK931372 · Product code: **ESD** · Ear, Nose, Throat
Source: <https://www.510kdatabase.net/k931372/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Mar 17, 1993
Decision date	Nov 3, 1993
Days to decision	231 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k931372/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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