

**K934917 RESOUND PERSONAL HEARING SYSTEMS
MODIFICATION**Jan 19, 1995
463 days to decisionK934917 · Product code: **ESD** · Ear, Nose, Throat
Source: <https://www.510kdatabase.net/k934917/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Oct 13, 1993
Decision date	Jan 19, 1995
Days to decision	463 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k934917/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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