

**K951379 RESOUND PERSONAL HEARING SYSTEM MODELS
ED3A & ED3A/S W/REMONTE CONTROL OPTION**Apr 27, 1995
31 days to decisionK951379 · Product code: **ESD** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k951379/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Mar 27, 1995
Decision date	Apr 27, 1995
Days to decision	31 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.netDevice record: <https://www.510kdatabase.net/k951379/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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