

**K960680 RESOUND PERSONAL HEARING SYSTEMS EDR  
ENCORE SERIES MODELS EDR-E, EDR-ES, EDR-EV, & EDR-P1**

Apr 10, 1996  
50 days to decision

K960680 · Product code: **ESD** · Ear, Nose, Throat  
Source: <https://www.510kdatabase.net/k960680/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Feb 20, 1996
Decision date	Apr 10, 1996
Days to decision	50 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k960680/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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