FDA 510(k) Premarket Notification

CLEARED

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

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

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