FDA 510(k) Premarket Notification

K931372 RESOUND PORTABLE PRESCRIPTIVE PROGRAMMING SYSTEM

CLEARED

Nov 3, 1993
231 days to decision

K931372 · 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