FDA 510(k) Premarket Notification

K960295 LONG LIFE RX-12

CLEARED

Mar 22, 1996
60 days to decision

K960295 · Product code: **ESD** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k960295/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Jan 22, 1996
Decision date	Mar 22, 1996
Days to decision	60 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Rexton, Inc.**
Location	Mchenry, IL, US
Contact	DANIEL I ANDERSON
510(k) history	21 submissions · 21 cleared · 1983-1997