FDA 510(k) Premarket Notification

K901860 INTRA III AND INTRA IV HEARING AIDS

CLEARED

Jul 20, 1990
87 days to decision

K901860 · Product code: **ESD** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k901860/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Apr 24, 1990
Decision date	Jul 20, 1990
Days to decision	87 days
Third-party review	No

APPLICANT

Company	**Starkey Laboratories, Inc.**
Location	Mchenry, IL, US
Contact	JERROLD LANG
510(k) history	60 submissions · 60 cleared · 1977-2020