

K901860 INTRA III AND INTRA IV HEARING AIDSJul 20, 1990
87 days to decisionK901860 · Product code: **ESD** · Ear, Nose, ThroatSource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k901860/>, Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 15, 2026