

K800012 RIONET HEARING AID HA-61Jan 29, 1980
25 days to decisionK800012 · Product code: **ESD** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k800012/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Jan 4, 1980
Decision date	Jan 29, 1980
Days to decision	25 days
Third-party review	No

APPLICANT

Company	Rion Acoustic Instruments, Inc.
Location	Mchenry, IL, US
510(k) history	51 submissions · 51 cleared · 1979-1988

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k800012/> 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 July 2, 2026