

K822163 ENT RETRACTORSAug 11, 1982
19 days to decisionK822163 · Product code: **ESD** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k822163/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Jul 23, 1982
Decision date	Aug 11, 1982
Days to decision	19 days
Third-party review	No

APPLICANT

Company	Kelleher Corp.
Location	Mchenry, IL, US
510(k) history	94 submissions · 94 cleared · 1982-1983

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k822163/> 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 29, 2026