FDA 510(k) Premarket Notification

K822163 ENT RETRACTORS

CLEARED

Aug 11, 1982
19 days to decision

K822163 · Product code: **ESD** · Ear, Nose, Throat
Source: 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