FDA 510(k) Premarket Notification

K832129 HEARING AID G3H & G3T

CLEARED

Jul 26, 1983
25 days to decision

K832129 · Product code: **ESD** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k832129/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Jul 1, 1983
Decision date	Jul 26, 1983
Days to decision	25 days
Third-party review	No

APPLICANT

Company	**Widex Hearing Aid Co., Inc.**
Location	Mchenry, IL, US
510(k) history	52 submissions · 52 cleared · 1976-2008