FDA 510(k) Premarket Notification

K933640 WIDEX MODEL LOGO 12 (L12)

CLEARED

Aug 23, 1993
27 days to decision

K933640 · Product code: **ESD** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k933640/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Jul 27, 1993
Decision date	Aug 23, 1993
Days to decision	27 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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