FDA 510(k) Premarket Notification

K904730 QUALITONE ULTRA CANAL GENESIS

CLEARED

Dec 7, 1990
51 days to decision

K904730 · Product code: **ESD** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k904730/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Oct 17, 1990
Decision date	Dec 7, 1990
Days to decision	51 days
Third-party review	No

APPLICANT

Company	**Qualitone**
Location	Mchenry, IL, US
Contact	A BROWN
510(k) history	39 submissions · 39 cleared · 1976-1996