FDA 510(k) Premarket Notification

K790878 SCREENING AUDIOMETER, HIGH FREQUENCY

CLEARED

Aug 3, 1979
88 days to decision

K790878 · Product code: **EWO** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k790878/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Audiometer (EWO)
Date received	May 7, 1979
Decision date	Aug 3, 1979
Days to decision	88 days
Third-party review	No

APPLICANT

Company	**Vicon Instrument Co.**
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1976-1981