FDA 510(k) Premarket Notification

K920142 PHILIPS M60 SERIES

CLEARED

Jan 23, 1992
10 days to decision

K920142 · Product code: **ESD** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k920142/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Jan 13, 1992
Decision date	Jan 23, 1992
Days to decision	10 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Philips Int'L B.V.**
Location	The Netherlands, NL
Contact	ROBERT MARTIN
510(k) history	11 submissions · 11 cleared · 1985-1992