

K943670 MO6 SERIESAug 18, 1994
23 days to decisionK943670 · Product code: **ESD** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k943670/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Jul 26, 1994
Decision date	Aug 18, 1994
Days to decision	23 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Philips Hearing Instruments Co.
Location	Mahwah, NJ, US
Contact	ROBERT MARTIN
510(k) history	20 submissions · 20 cleared · 1991-1997

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k943670/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 5, 2026