

**K920142 PHILIPS M60 SERIES**Jan 23, 1992  
10 days to decisionK920142 · Product code: **ESD** · Ear, Nose, ThroatSource: <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	<b>Philips Int&amp;apos;L B.V.</b>
Location	The Netherlands, NL
Contact	ROBERT MARTIN
510(k) history	11 submissions · 11 cleared · 1985-1992

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k920142/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated July 6, 2026