

K953911 PHILIPS EASYVISIONJun 27, 1996
311 days to decisionK953911 · Product code: **LLZ** · Radiology
Source: <https://www.510kdatabase.net/k953911/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Aug 21, 1995
Decision date	Jun 27, 1996
Days to decision	311 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Philips Medical Systems, Inc.
Location	Mchenry, IL, US
Contact	PETER ALTMAN
510(k) history	111 submissions · 110 cleared · 1983-2010

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k953911/> 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 May 17, 2026