

K930246 MAMMORXJul 26, 1993
188 days to decisionK930246 · Product code: **IYE** · Radiology
Source: <https://www.510kdatabase.net/k930246/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accelerator, Linear, Medical (IYE)
Date received	Jan 19, 1993
Decision date	Jul 26, 1993
Days to decision	188 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Diacor, Inc.
Location	Salt Lake City, UT, US
Contact	GLENN N WATERMAN
510(k) history	10 submissions · 10 cleared · 1986-2013

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k930246/> 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