FDA 510(k) Premarket Notification

K930246 MAMMORX

CLEARED

Jul 26, 1993
188 days to decision

K930246 · 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