FDA 510(k) Premarket Notification

K050888 INTEGRA IMMOBILIZATION SYSTEM

CLEARED

May 23, 2005
46 days to decision

K050888 · Product code: **IYE** · Radiology
Source: https://www.510kdatabase.net/k050888/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accelerator, Linear, Medical (IYE)
Date received	Apr 7, 2005
Decision date	May 23, 2005
Days to decision	46 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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