FDA 510(k) Premarket Notification

K950112 CENTRALITE(R)-DLL SERIES

CLEARED

Aug 16, 1995
217 days to decision

K950112 · Product code: **IWE** · Radiology
Source: https://www.510kdatabase.net/k950112/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Patient Position, Light-beam (IWE)
Date received	Jan 11, 1995
Decision date	Aug 16, 1995
Days to decision	217 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