FDA 510(k) Premarket Notification

K893398 CENTRALITE BACKPOINTER

CLEARED

Aug 17, 1989
108 days to decision

K893398 · Product code: **IWE** · Radiology
Source: https://www.510kdatabase.net/k893398/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Patient Position, Light-beam (IWE)
Date received	May 1, 1989
Decision date	Aug 17, 1989
Days to decision	108 days
Third-party review	No

APPLICANT

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