FDA 510(k) Premarket Notification

K032142 DATACAPTOR

CLEARED

Aug 8, 2003
25 days to decision

K032142 · Product code: **MWI** · Cardiovascular
Source: https://www.510kdatabase.net/k032142/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Monitor, Physiological, Patient (without Arrhythmia Detection Or Alarms) (MWI)
Date received	Jul 14, 2003
Decision date	Aug 8, 2003
Days to decision	25 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Capsule Technologie**
Location	Deer Field, IL, US
Contact	DANIEL KAMM
510(k) history	5 submissions · 5 cleared · 2001-2019