

K020197 MODIFICATION TO DATACAPTORFeb 21, 2002
30 days to decisionK020197 · Product code: **MWI** · Cardiovascular
Source: <https://www.510kdatabase.net/k020197/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Monitor, Physiological, Patient (without Arrhythmia Detection Or Alarms) (MWI)
Date received	Jan 22, 2002
Decision date	Feb 21, 2002
Days to decision	30 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

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