

K010342 M2376A DEVICELINK SYSTEM, MODEL M2376AFeb 14, 2001
9 days to decisionK010342 · Product code: **MWI** · Cardiovascular
Source: <https://www.510kdatabase.net/k010342/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Monitor, Physiological, Patient (without Arrhythmia Detection Or Alarms) (MWI)
Date received	Feb 5, 2001
Decision date	Feb 14, 2001
Days to decision	9 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Agilent Technologies, Inc.
Location	Pittsburgh, PA, US
Contact	MIKE HUDON
Website	http://www.agilent.com
510(k) history	30 submissions · 30 cleared · 1985-2017

Agilent Technologies, Inc. is an American global company that provides instruments, software, services, and consumables for laboratories. Headquartered in Santa Clara, California, Agilent was established in 1999 as a spin-off from Hewlett-Packard. The company has received FDA 510(k) clearances from total submissions. Cardiovascular devices represent the dominant focus, accounting for approximately 80% of regulatory submissions. Agilent's FDA 510(k) clearance history spans from 1985 to 2017, establishing a long track record in medical device regulation. Notable cleared dev...