FDA 510(k) Premarket Notification

K980688 INTEGRITI PATIENT MONITOR

CLEARED

Mar 16, 1999
386 days to decision

K980688 · Product code: **MWI** · Cardiovascular
Source: https://www.510kdatabase.net/k980688/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient (without Arrhythmia Detection Or Alarms) (MWI)
Date received	Feb 23, 1998
Decision date	Mar 16, 1999
Days to decision	386 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Alliance Instruments**
Location	Vancouver, WA, US
Contact	JIM ROOKS
510(k) history	1 submissions · 1 cleared · 1999-1999

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k980688/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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