

**K033030 CALYSTO SERIES IV PATIENT CARE MONITORS AND
CENTRAL STATION AND ECG MANAGEMENT SYSTEM**Feb 24, 2004
151 days to decisionK033030 · Product code: **MWI** · Cardiovascular
Source: <https://www.510kdatabase.net/k033030/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient (without Arrhythmia Detection Or Alarms) (MWI)
Date received	Sep 26, 2003
Decision date	Feb 24, 2004
Days to decision	151 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Witt Biomedical Corp
Location	Melbourne, FL, US
Contact	ELFRIEDE PAGAN
510(k) history	5 submissions · 5 cleared · 2004-2010

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k033030/>, Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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