

**K060756 MEDLOG, MODEL 1-0-10**Oct 30, 2006  
223 days to decisionK060756 · Product code: **MWI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k060756/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient (without Arrhythmia Detection Or Alarms) (MWI)
Date received	Mar 21, 2006
Decision date	Oct 30, 2006
Days to decision	223 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Bmc Gesellschaft Fur Medizintechnik Mbh</b>
Location	Hamburg, DE
Contact	ANGELIKA M BERGER
510(k) history	1 submissions · 1 cleared · 2006-2006

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k060756/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 29, 2026