

**K010770 BCI ADVISOR VITAL SIGNS MONITOR (9200)**Jun 12, 2001  
90 days to decisionK010770 · Product code: **MWI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k010770/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient (without Arrhythmia Detection Or Alarms) (MWI)
Date received	Mar 14, 2001
Decision date	Jun 12, 2001
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Bci, Inc.</b>
Location	Mchenry, IL, US
Contact	DONALD ALEXANDER
510(k) history	13 submissions · 13 cleared · 1979-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k010770/> 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 May 25, 2026