

**K981939 AUTOCORR PLUS**Aug 20, 1998  
79 days to decisionK981939 · Product code: **DQA** · AnesthesiologySource: <https://www.510kdatabase.net/k981939/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Jun 2, 1998
Decision date	Aug 20, 1998
Days to decision	79 days
Third-party review	No
Summary / Statement	Summary
Other names	3404 OXIMETER/ECG MONITOR

**APPLICANT**

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Company	<b>Bci Intl., Inc.</b>
Location	Waukesha, WI, US
Contact	DONALD J ALEXANDER
510(k) history	20 submissions · 20 cleared · 1994-1999

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k981939/> 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 26, 2026