

**K020739 BCI 3178 PEDIATRIC PULSE OXIMETRY SENSOR**Jun 3, 2002  
89 days to decisionK020739 · Product code: **DQA** · AnesthesiologySource: <https://www.510kdatabase.net/k020739/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Mar 6, 2002
Decision date	Jun 3, 2002
Days to decision	89 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Bci, Inc.</b>
Location	Mchenry, IL, US
Contact	DONALD J 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/k020739/> 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