

**K833826 BTI BIOX III OXIMETER CRITICAL CAREUNI**Dec 30, 1983  
65 days to decisionK833826 · Product code: **DQA** · Anesthesiology  
Source: <https://www.510kdatabase.net/k833826/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Oct 26, 1983
Decision date	Dec 30, 1983
Days to decision	65 days
Third-party review	No

**APPLICANT**

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Company	<b>Biox Technology, Inc.</b>
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1982-1983

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k833826/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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