

**K811391 OXIMETRIX ACCU SAT CPB**Jul 20, 1981  
63 days to decisionK811391 · Product code: **DRY** · CardiovascularSource: <https://www.510kdatabase.net/k811391/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Blood-gas, On-line, Cardiopulmonary Bypass (DRY)
Date received	May 18, 1981
Decision date	Jul 20, 1981
Days to decision	63 days
Third-party review	No

**APPLICANT**

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Company	<b>Oximetrix, Inc.</b>
Location	Mchenry, IL, US
510(k) history	12 submissions · 12 cleared · 1976-1986

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

Device record: <https://www.510kdatabase.net/k811391/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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