

K760234 OXIMETER, IN VIVO CATHETERAug 23, 1976
39 days to decisionK760234 · Product code: **DQE** · CardiovascularSource: <https://www.510kdatabase.net/k760234/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Oximeter, Fiber-optic (DQE)
Date received	Jul 15, 1976
Decision date	Aug 23, 1976
Days to decision	39 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k760234/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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