

**K890817 OMNI-TRAK (PATIENT MONITOR)**Aug 10, 1989  
174 days to decisionK890817 · Product code: **CCK** · Anesthesiology  
Source: <https://www.510kdatabase.net/k890817/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Gas, Carbon-dioxide, Gaseous-phase (CCK)
Date received	Feb 17, 1989
Decision date	Aug 10, 1989
Days to decision	174 days
Third-party review	No

**APPLICANT**

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Company	<b>Invivo Research, Inc.</b>
Location	Orlando, FL, US
Contact	TOM W FOSHEE
510(k) history	14 submissions · 14 cleared · 1989-2005

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k890817/>; 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 27, 2026