

**K971840 OMNI-TRAK MODEL 3150 SERIES PATIENT MONITORING SYSTEM**

Mar 4, 1998  
289 days to decision

K971840 · Product code: **MHX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k971840/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	May 19, 1997
Decision date	Mar 4, 1998
Days to decision	289 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Invivo Research, Inc.</b>
Location	Orlando, FL, US
Contact	FRANCIS X CASEY
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/k971840/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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