

**K955045 OMNI-TRAK PATIENT MONITORING SYSTEM**Jul 11, 1996  
251 days to decisionK955045 · Product code: **DPS** · CardiovascularSource: <https://www.510kdatabase.net/k955045/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	Nov 3, 1995
Decision date	Jul 11, 1996
Days to decision	251 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k955045/> 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