FDA 510(k) Premarket Notification

K890817 OMNI-TRAK (PATIENT MONITOR)

CLEARED

Aug 10, 1989
174 days to decision

K890817 · Product code: **CCK** · Anesthesiology
Source: https://www.510kdatabase.net/k890817/

SUBMISSION DETAILS

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

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