

**K991513 HOMEPUMP C-SERIES AND HOMEPUMP C-SERIES
ONE-STEP KVO**Jun 21, 1999
52 days to decisionK991513 · Product code: **FPA** · General Hospital
Source: <https://www.510kdatabase.net/k991513/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Apr 30, 1999
Decision date	Jun 21, 1999
Days to decision	52 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	I-Flow Corp.
Location	Irvine, CA, US
Contact	ROBERT J BARD
510(k) history	41 submissions · 41 cleared · 1988-2007

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k991513/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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