

**K063363 FIRMGRIP PERIPHERALLY INSERTED CATHETER
DEVICE**May 7, 2007
181 days to decisionK063363 · Product code: **LJS** · General Hospital
Source: <https://www.510kdatabase.net/k063363/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Long-term Greater Than 30 Days (LJS)
Date received	Nov 7, 2006
Decision date	May 7, 2007
Days to decision	181 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Flexicath , Ltd.
Location	Kfar Saba, IL
Contact	AHAVA STEIN
510(k) history	4 submissions · 4 cleared · 2007-2012

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k063363/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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