

**K093271 EXTENSION SET (STANDARD BORE TUBING),
EXTENSION SET (MINIBORE TUBING), IV ADMINISTRATION
SET (STANDARD**

Jan 27, 2010
100 days to decision

K093271 · Product code: **FPA** · General Hospital
Source: <https://www.510kdatabase.net/k093271/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Oct 19, 2009
Decision date	Jan 27, 2010
Days to decision	100 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Victus, Inc.
Location	Miami, FL, US
Contact	ILEANA YANES
510(k) history	3 submissions · 3 cleared · 2002-2010

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k093271/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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