

**K030246 VICTUS I.V. ADMINISTRATION SET, MODELS 27071
AND 27072**Feb 26, 2003
33 days to decisionK030246 · Product code: **FPA** · General Hospital
Source: <https://www.510kdatabase.net/k030246/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Jan 24, 2003
Decision date	Feb 26, 2003
Days to decision	33 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k030246/> Data sourced from FDA 510(k) public records (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. - Generated June 21, 2026