FDA 510(k) Premarket Notification

K030246 VICTUS I.V. ADMINISTRATION SET, MODELS 27071 AND 27072

CLEARED

Feb 26, 2003
33 days to decision

K030246 · 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