FDA 510(k) Premarket Notification

CLEARED

K023668 JMS EXTENSION SET WITH PLANECTA, PLANECTA (STAND ALONE UNIT), PLANECTA LOCK & PLANECTA ADAPTOR

Jul 1, 2003
243 days to decision

K023668 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k023668/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Oct 31, 2002
Decision date	Jul 1, 2003
Days to decision	243 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Jms North America Corp.**
Location	Hayward, CA, US
Contact	SWEE CHEAU CHONG
510(k) history	10 submissions · 10 cleared · 2003-2010

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k023668/ 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