FDA 510(k) Premarket Notification

K120195 SAFETY SUBCUTANEOUS TISSUE INFUSION SET

CLEARED

Jul 6, 2012
165 days to decision

K120195 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k120195/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Jan 23, 2012
Decision date	Jul 6, 2012
Days to decision	165 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Multi Med**
Location	Apollo Beach, FL, US
Contact	TANYA O'BRIEN
510(k) history	1 submissions · 1 cleared · 2012-2012