FDA 510(k) Premarket Notification

K092172 EXEL I.V. EXTENSION SET

CLEARED

Jan 4, 2010
167 days to decision

K092172 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k092172/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Jul 21, 2009
Decision date	Jan 4, 2010
Days to decision	167 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Exelint Intl. Co.**
Location	Culver City, CA, US
Contact	ARMAND HAMID
510(k) history	12 submissions · 12 cleared · 2000-2010