FDA 510(k) Premarket Notification

K092382 NEXUS PRESSURE RATED EXTENSION SETS

CLEARED

Apr 26, 2010
264 days to decision

K092382 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k092382/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Aug 5, 2009
Decision date	Apr 26, 2010
Days to decision	264 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Nexus Medical, LLC**
Location	Raleigh, NC, US
Contact	SUSAN CURRY
510(k) history	16 submissions · 16 cleared · 2003-2015