FDA 510(k) Premarket Notification

K092930 NEXUS INTRAVASCULAR TUBING SETS-PRE-PIERCED

CLEARED

Mar 10, 2010
169 days to decision

K092930 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k092930/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Set, Administration, Intravascular (FPA)
Date received	Sep 22, 2009
Decision date	Mar 10, 2010
Days to decision	169 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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