FDA 510(k) Premarket Notification

K113398 NEXUS TKO-6, LUER-ACTIVATED DEVICE

CLEARED

Apr 18, 2012
153 days to decision

K113398 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k113398/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Nov 17, 2011
Decision date	Apr 18, 2012
Days to decision	153 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