

**K113398 NEXUS TKO-6, LUER-ACTIVATED DEVICE**Apr 18, 2012  
153 days to decisionK113398 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k113398/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k113398/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 5, 2026