

**K130879 NEXUS IV TUBING SET WITH LUER-ACCESSBILE ONE-WAY CHECK VALVE - IV PLUS**

Jul 9, 2013  
102 days to decision

K130879 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k130879/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Mar 29, 2013
Decision date	Jul 9, 2013
Days to decision	102 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/k130879/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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