

**K073444 TRITON MEIDCAL TMED IV-SET**Aug 18, 2008  
255 days to decisionK073444 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k073444/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Dec 7, 2007
Decision date	Aug 18, 2008
Days to decision	255 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Triton Medical, LLC</b>
Location	Austin, TX, US
Contact	IAN P GORDON
510(k) history	1 submissions · 1 cleared · 2008-2008

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k073444/> 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 24, 2026