

**K110282 LIGHTY DD INFUSION SET**Apr 27, 2011  
86 days to decisionK110282 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k110282/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Jan 31, 2011
Decision date	Apr 27, 2011
Days to decision	86 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Medx-Set, Ltd.</b>
Location	Hudson, NH, US
Contact	RAYMOND KELLY
510(k) history	1 submissions · 1 cleared · 2011-2011

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k110282/> 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 19, 2026