

**K042666 LUTH-ALL SUB-Q NEEDLE SET**Dec 21, 2004  
83 days to decisionK042666 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k042666/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Sep 29, 2004
Decision date	Dec 21, 2004
Days to decision	83 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Luth-All Medical Products, LLC</b>
Location	Laguna Niguel, CA, US
Contact	SHEPARD G BENTLEY
510(k) history	1 submissions · 1 cleared · 2004-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k042666/> 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 July 2, 2026