

**K021565 LUTHER SAFETY HUBER NEEDLE SET**Nov 14, 2002  
185 days to decisionK021565 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k021565/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	May 13, 2002
Decision date	Nov 14, 2002
Days to decision	185 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Luther Research Partners</b>
Location	Irvine, CA, US
Contact	GREY HOLLAND
510(k) history	1 submissions · 1 cleared · 2002-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k021565/> 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