

**K062308 LMT-KATH-S**Mar 20, 2007  
224 days to decisionK062308 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k062308/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Set, Administration, Intravascular (FPA)
Date received	Aug 8, 2006
Decision date	Mar 20, 2007
Days to decision	224 days
Third-party review	No
Summary / Statement	Summary
Other names	LMT-KATH-I

**APPLICANT**

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Company	<b>Licher Medizintechnologie GmbH</b>
Location	Wedemark, DE
Contact	DIETER MUSHOFF
510(k) history	1 submissions · 1 cleared · 2007-2007

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k062308/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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