

**K063466 TOP FINE**Mar 2, 2007  
106 days to decisionK063466 · Product code: **FMI** · General Hospital  
Source: <https://www.510kdatabase.net/k063466/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Nov 16, 2006
Decision date	Mar 2, 2007
Days to decision	106 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Daejin Tech Medical Manufacturing Co., Ltd.</b>
Location	Deer Field, IL, US
Contact	DANIEL KAMM
510(k) history	2 submissions · 2 cleared · 2007-2007

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k063466/> 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 28, 2026