

**K922396 FUJINON FORCEPS -- MODIFICATION**Dec 10, 1992  
205 days to decisionK922396 · Product code: **FCL** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k922396/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Forceps, Biopsy, Non-electric (FCL)
Date received	May 19, 1992
Decision date	Dec 10, 1992
Days to decision	205 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Hobbs Medical, Inc.</b>
Location	Mchenry, IL, US
Contact	ROB WHALEN
510(k) history	21 submissions · 21 cleared · 1983-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k922396/> 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 20, 2026