

**K946238 SYMBIOSIS GASTROINTESTINAL BIOPSY FORCEPS**Jan 20, 1995  
29 days to decisionK946238 · Product code: **FCL** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k946238/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Forceps, Biopsy, Non-electric (FCL)
Date received	Dec 22, 1994
Decision date	Jan 20, 1995
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Symbiosis Corp.</b>
Location	Miami, FL, US
Contact	KEVIN W SMITH
510(k) history	34 submissions · 32 cleared · 1989-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k946238/> 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 May 22, 2026