

**K900065 NON-ELECTRIC BIOPSY FORCEPS**Mar 15, 1990  
70 days to decisionK900065 · Product code: **FCL** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k900065/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Forceps, Biopsy, Non-electric (FCL)
Date received	Jan 4, 1990
Decision date	Mar 15, 1990
Days to decision	70 days
Third-party review	No

**APPLICANT**

---

Company	<b>Esco Precision, Inc.</b>
Location	Stony Brook, NY, US
Contact	TED ESSER
510(k) history	3 submissions · 3 cleared · 1990-1991

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k900065/>; 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