

K900015 NON-ELECTRIC BIOPSY FORCEPSMar 15, 1990
72 days to decisionK900015 · Product code: **FCL** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k900015/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Forceps, Biopsy, Non-electric (FCL)
Date received	Jan 2, 1990
Decision date	Mar 15, 1990
Days to decision	72 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k900015/>; Data sourced from FDA 510(k) public records (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. - Generated July 2, 2026