FDA 510(k) Premarket Notification

K900065 NON-ELECTRIC BIOPSY FORCEPS

CLEARED

Mar 15, 1990
70 days to decision

K900065 · Product code: **FCL** · Gastroenterology & Urology
Source: 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	**Esco Precision, Inc.**
Location	Stony Brook, NY, US
Contact	TED ESSER
510(k) history	3 submissions · 3 cleared · 1990-1991