

**K033593 REPROCESSED SINGLE USE DEVICES BIOPSY
FORCEPS**Nov 20, 2003
7 days to decisionK033593 · Product code: **NLU** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k033593/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Forceps, Biopsy, Electric, Reprocessed (NLU)
Date received	Nov 13, 2003
Decision date	Nov 20, 2003
Days to decision	7 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Clearmedical, Inc.
Location	Bellevue, WA, US
Contact	MIKE KOVACS
510(k) history	14 submissions · 14 cleared · 2002-2005

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k033593/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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