FDA 510(k) Premarket Notification

K033593 REPROCESSED SINGLE USE DEVICES BIOPSY FORCEPS

CLEARED

Nov 20, 2003
7 days to decision

K033593 · 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