FDA 510(k) Premarket Notification

K922396 FUJINON FORCEPS -- MODIFICATION

CLEARED

Dec 10, 1992
205 days to decision

K922396 · Product code: **FCL** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k922396/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Forceps, Biopsy, Non-electric (FCL)
Date received	May 19, 1992
Decision date	Dec 10, 1992
Days to decision	205 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Hobbs Medical, Inc.**
Location	Mchenry, IL, US
Contact	ROB WHALEN
510(k) history	21 submissions · 21 cleared · 1983-2024