FDA 510(k) Premarket Notification

K946238 SYMBIOSIS GASTROINTESTINAL BIOPSY FORCEPS

CLEARED

Jan 20, 1995
29 days to decision

K946238 · Product code: **FCL** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k946238/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Forceps, Biopsy, Non-electric (FCL)
Date received	Dec 22, 1994
Decision date	Jan 20, 1995
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Symbiosis Corp.**
Location	Miami, FL, US
Contact	KEVIN W SMITH
510(k) history	34 submissions · 32 cleared · 1989-1996