FDA 510(k) Premarket Notification

K911448 MULTIPLE BIOPSY DEVICE-MBX

CLEARED

Apr 16, 1991
32 days to decision

K911448 · Product code: **FCK** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k911448/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Biopsy, Suction (FCK)
Date received	Mar 15, 1991
Decision date	Apr 16, 1991
Days to decision	32 days
Third-party review	No

APPLICANT

Company	**Triton Technology, Inc.**
Location	San Diego, CA, US
Contact	WILLIAM Z KOLOZSI
510(k) history	4 submissions · 4 cleared · 1988-1993