FDA 510(k) Premarket Notification

K910502 VESSEL DILATORS

CLEARED

Apr 29, 1991
83 days to decision

K910502 · Product code: **FKA** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k910502/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Vessel (FKA)
Date received	Feb 5, 1991
Decision date	Apr 29, 1991
Days to decision	83 days
Third-party review	No

APPLICANT

Company	**Akcess Medical Products, Inc.**
Location	New Brunswick, NJ, US
Contact	BALBIR B KAPANY
510(k) history	36 submissions · 28 cleared · 1989-1994