FDA 510(k) Premarket Notification

K892932 BIOPRIME TM KIT

CLEARED

Jun 22, 1989
62 days to decision

K892932 · Product code: **FJK** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k892932/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Tubing, Blood, With And Without Anti-regurgitation Valve (FJK)
Date received	Apr 21, 1989
Decision date	Jun 22, 1989
Days to decision	62 days
Third-party review	No

APPLICANT

Company	**Hospal**
Location	Williamsburg, VA, US
Contact	JO LANG
510(k) history	1 submissions · 1 cleared · 1989-1989