

K892932 BIOPRIME TM KITJun 22, 1989
62 days to decisionK892932 · Product code: **FJK** · Gastroenterology & UrologySource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k892932/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 30, 2026