FDA 510(k) Premarket Notification

K896806 CATHETER PLUG/PROTECTOR

CLEARED

Feb 26, 1990
87 days to decision

K896806 · Product code: **KNY** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k896806/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessories, Catheter, G-u (KNY)
Date received	Dec 1, 1989
Decision date	Feb 26, 1990
Days to decision	87 days
Third-party review	No

APPLICANT

Company	**Westmark, Sterile Packing Systems, Inc. (Sps)**
Location	Grand Rapids, MI, US
Contact	SURY SHONEY
510(k) history	28 submissions · 26 cleared · 1988-1990

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k896806/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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