

**K960949 PROSYS BEDSIDE DRAINAGE BAG (STERILE) MODEL
650167**Apr 19, 1996
39 days to decisionK960949 · Product code: **KNX** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k960949/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Collector, Urine, (and Accessories) For Indwelling Catheter (KNX)
Date received	Mar 11, 1996
Decision date	Apr 19, 1996
Days to decision	39 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Bristol-Myers Squibb
Location	Skillman, NJ, US
Contact	SUSIE MERCHANT
510(k) history	1 submissions · 1 cleared · 1996-1996

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k960949/>; 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 July 1, 2026