K960400 DIAMOND-TOUCH AND MICRO DIAMOND-TOUCH INSTRUMENTS/DIAMOND-LINE INSTRUMENTS/DIAMOND-PORT(ACCESS PARTS)

CLEARED

Mar 12, 1996
43 days to decision

K960400 · Product code: **FBM** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k960400/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula And Trocar, Suprapubic, Non-disposable (FBM)
Date received	Jan 29, 1996
Decision date	Mar 12, 1996
Days to decision	43 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Snowden-Pencer**
Location	Norcross, GA, US
Contact	JULIE A STEPHENS
510(k) history	10 submissions · 10 cleared · 1986-1996

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k960400/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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