FDA 510(k) Premarket Notification

K896680 PROBE SHEATH FOR ADMS & CAPISTRANO LABS TRANSREC.

CLEARED

Feb 14, 1990
79 days to decision

K896680 · Product code: **FED** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k896680/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Endoscopic Access Overtube, Gastroenterology-urology (FED)
Date received	Nov 27, 1989
Decision date	Feb 14, 1990
Days to decision	79 days
Third-party review	No

APPLICANT

Company	**Interson Corp.**
Location	Pleasanton, CA, US
Contact	MARK F HAYWARD
510(k) history	9 submissions · 9 cleared · 1989-1991