

**K896680 PROBE SHEATH FOR ADMS & CAPISTRANO LABS  
TRANSREC.**Feb 14, 1990  
79 days to decisionK896680 · 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	<b>Interson Corp.</b>
Location	Pleasanton, CA, US
Contact	MARK F HAYWARD
510(k) history	9 submissions · 9 cleared · 1989-1991

---

**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k896680/>. 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 May 21, 2026