

K980592 NTECH CYSTOSCOPESApr 28, 1998
70 days to decisionK980592 · Product code: **FAJ** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k980592/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cystoscope And Accessories, Flexible/rigid (FAJ)
Date received	Feb 17, 1998
Decision date	Apr 28, 1998
Days to decision	70 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	N-Tech Endoscopy, Inc.
Location	Mundelein, IL, US
Contact	NICK SURO
510(k) history	3 submissions · 3 cleared · 1998-1998

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k980592/> 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