

K042780 NEOSCOPE - ENDOSCOPIC DIAGNOSTIC & TREATMENT SYSTEMFeb 3, 2005
120 days to decisionK042780 · Product code: **FAJ** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k042780/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cystoscope And Accessories, Flexible/rigid (FAJ)
Date received	Oct 6, 2004
Decision date	Feb 3, 2005
Days to decision	120 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Prosurg, Inc.
Location	San Jose, CA, US
Contact	ASHVIN DESAI
510(k) history	16 submissions · 16 cleared · 2000-2016

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k042780/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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