FDA 510(k) Premarket Notification

CLEARED

K042780 NEOSCOPE - ENDOSCOPIC DIAGNOSTIC & TREATMENT SYSTEM

Feb 3, 2005
120 days to decision

K042780 · 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