FDA 510(k) Premarket Notification

K040961 INJECTOR NEEDLE/SNARE

CLEARED

Jul 8, 2004
86 days to decision

K040961 · Product code: **FDI** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k040961/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Snare, Flexible (FDI)
Date received	Apr 13, 2004
Decision date	Jul 8, 2004
Days to decision	86 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**United States Endoscopy Group, Inc.**
Location	Mentor, OH, US
Contact	GRETCHEN Y COHEN
510(k) history	94 submissions · 92 cleared · 1991-2020