FDA 510(k) Premarket Notification

K912502 LAPAROSCOPIC LASER FIBER DIVERTER

CLEARED

Jul 25, 1991
50 days to decision

K912502 · Product code: **GCJ** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k912502/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laparoscope, General & Plastic Surgery (GCJ)
Date received	Jun 5, 1991
Decision date	Jul 25, 1991
Days to decision	50 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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