FDA 510(k) Premarket Notification

K012660 GIMMI'ALPHA' ENDOSCOPES & ACCESSORIES

CLEARED

Dec 20, 2001
129 days to decision

K012660 · Product code: **FET** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k012660/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Endoscopic Video Imaging System/component, Gastroenterology-urology (FET)
Date received	Aug 13, 2001
Decision date	Dec 20, 2001
Days to decision	129 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Gimmi GmbH**
Location	Amsterdam, Nh, NL
Contact	DAGMAR MASER
510(k) history	6 submissions · 6 cleared · 2001-2010