

K012660 GIMMI'ALPHA' ENDOSCOPES & ACCESSORIESDec 20, 2001
129 days to decisionK012660 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k012660/>, Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 21, 2026