

**K013731 GIMMI ALPHA SINUSCOPES MODEL # E.8282.01/31/71
+ E.8284.01/31/71**

Nov 27, 2001
18 days to decision

K013731 · Product code: **EOQ** · Ear, Nose, Throat
Source: <https://www.510kdatabase.net/k013731/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bronchoscope (flexible Or Rigid) (EOQ)
Date received	Nov 9, 2001
Decision date	Nov 27, 2001
Days to decision	18 days
Third-party review	No
Summary / Statement	Summary
Other names	GIMMI ALPHA BRONCHOSCOPES MODEL # E.8189.00/01/02/03

APPLICANT

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k013731/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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