

K212382 Ambu® aScope™ Gastro, Ambu® aBox™2Feb 3, 2022
185 days to decisionK212382 · Product code: **FDS** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k212382/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Gastroscope And Accessories, Flexible/rigid (FDS)
Date received	Aug 2, 2021
Decision date	Feb 3, 2022
Days to decision	185 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Ambu A/S
Location	Glen Burnie, MD, US
Contact	Mette Anderson
Website	https://www.ambu.com
510(k) history	38 submissions · 38 cleared · 2005-2026

Ambu A/S is a global medical device company specializing in single-use endoscopy and airway management solutions. The company operates with a manufacturing facility in Glen Burnie, Maryland, and serves hospitals and emergency care settings worldwide. Ambu created the single-use endoscopy market in 2009 and remains the market leader in this category. Ambu has received FDA 510(k) clearances from total submissions, with no denied submissions on record. The company's regulatory activity spans from 2005 to 2026, demonstrating sustained innovation and market presence. Recent cl...

REGULATORY CONSULTANT

Consulting firm	Ambu, Inc.
Contact	Sanjay Parikh

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)

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Device record: <https://www.510kdatabase.net/k212382/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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