

K944186 SINUSCOPENov 7, 1994
73 days to decisionK944186 · Product code: **EOB** · Ear, Nose, Throat
Source: <https://www.510kdatabase.net/k944186/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Nasopharyngoscope (flexible Or Rigid) (EOB)
Date received	Aug 26, 1994
Decision date	Nov 7, 1994
Days to decision	73 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Stryker Endoscopy
Location	San Jose, CA, US
Contact	MARK ZIEGLER
Website	https://www.stryker.com
510(k) history	101 submissions · 101 cleared · 1993-2026

Stryker Endoscopy is a medical device manufacturer based in San Jose, US. The company specializes in endoscopic and surgical imaging systems for minimally invasive procedures. Stryker Endoscopy has received FDA 510(k) clearances from total submissions since its first clearance in 1993. The company maintains active regulatory status, with its latest clearance in 2026. Its portfolio spans General & Plastic Surgery devices, orthopedic surgical tools, and obstetric and gynecologic instruments. Recent cleared devices include advanced imaging systems such as 4K camera platforms...
