

K082276 SYNTHEMED DEVICENov 18, 2008
99 days to decisionK082276 · Product code: **LYA** · Ear, Nose, Throat
Source: <https://www.510kdatabase.net/k082276/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Splint, Intranasal Septal (LYA)
Date received	Aug 11, 2008
Decision date	Nov 18, 2008
Days to decision	99 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Synthemed, Inc.
Location	Iselin, NJ, US
Contact	ELI PINES
510(k) history	1 submissions · 1 cleared · 2008-2008

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k082276/> 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 24, 2026