

K980221 SILIMED NASAL RETAINERApr 2, 1998
70 days to decisionK980221 · Product code: **LYA** · Ear, Nose, Throat
Source: <https://www.510kdatabase.net/k980221/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Splint, Intranasal Septal (LYA)
Date received	Jan 22, 1998
Decision date	Apr 2, 1998
Days to decision	70 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Silimed, LLC
Location	Crofton, MD, US
Contact	E.J. Smith
510(k) history	9 submissions · 9 cleared · 1998-1998

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k980221/> 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 29, 2026