FDA 510(k) Premarket Notification

K980221 SILIMED NASAL RETAINER

CLEARED

Apr 2, 1998
70 days to decision

K980221 · 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