

K080247 QUIKLOT NOSEBLEEDFeb 27, 2008
27 days to decisionK080247 · Product code: **LYA** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k080247/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Splint, Intranasal Septal (LYA)
Date received	Jan 31, 2008
Decision date	Feb 27, 2008
Days to decision	27 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Z-Medica Corporation
Location	Wallingford, CT, US
Contact	RONALD E PETERSON
510(k) history	8 submissions · 8 cleared · 2005-2013

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k080247/> 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 9, 2026