FDA 510(k) Premarket Notification

K080247 QUIKCLOT NOSEBLEED

CLEARED

Feb 27, 2008
27 days to decision

K080247 · 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