FDA 510(k) Premarket Notification

K070859 REPROCESSED ETHICON ETS ENDOSCOPIC LINEAR CUTTERS

CLEARED

Sep 12, 2007
168 days to decision

K070859 · Product code: **NLL** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k070859/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Staple, Implantable, Reprocessed (NLL)
Date received	Mar 28, 2007
Decision date	Sep 12, 2007
Days to decision	168 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Sterilmed, Inc.**
Location	Plymouth, MN, US
Contact	CAROLINE BUTTERFIELD
510(k) history	64 submissions · 64 cleared · 2001-2024