FDA 510(k) Premarket Notification

CLEARED

K180499 Reprocessed LigaSure Maryland Jaw Sealer/Divider

Apr 23, 2018
56 days to decision

K180499 · Product code: **NUJ** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k180499/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation Accessories, Laparoscopic & Endoscopic, Reprocessed (NUJ)
Date received	Feb 26, 2018
Decision date	Apr 23, 2018
Days to decision	56 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Stryker Sustainability Solutions**
Location	Tempe, AZ, US
Contact	Scott English
510(k) history	31 submissions · 31 cleared · 2011-2025