

# 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

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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

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k180499/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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