FDA 510(k) Premarket Notification

K173740 Reprocessed DePuy Mitek Ablation Wand

CLEARED

Mar 19, 2018
102 days to decision

K173740 · Product code: **NUJ** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k173740/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation Accessories, Laparoscopic & Endoscopic, Reprocessed (NUJ)
Date received	Dec 7, 2017
Decision date	Mar 19, 2018
Days to decision	102 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Renovo, Inc.**
Location	Bend, OR, US
Contact	Mark K. Wells
510(k) history	6 submissions · 6 cleared · 2017-2018