FDA 510(k) Premarket Notification

K171304 Maxxi Rip Sensor

CLEARED

Jun 24, 2019
782 days to decision

K171304 · Product code: **MNR** · Anesthesiology
Source: https://www.510kdatabase.net/k171304/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ventilatory Effort Recorder (MNR)
Date received	May 3, 2017
Decision date	Jun 24, 2019
Days to decision	782 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Neurovirtual USA, Inc.**
Location	Doral, FL, US
Contact	Eduardo Faria
510(k) history	8 submissions · 8 cleared · 2006-2020