

K171304 Maxxi Rip SensorJun 24, 2019
782 days to decisionK171304 · Product code: **MNR** · AnesthesiologySource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k171304/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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