FDA 510(k) Premarket Notification

K191220 Pure Vu System

CLEARED

Jun 6, 2019
30 days to decision

K191220 · Product code: **FDF** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k191220/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Colonoscope And Accessories, Flexible/rigid (FDF)
Date received	May 7, 2019
Decision date	Jun 6, 2019
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Motus GI Medical Technologies , Ltd.**
Location	Tirat Carmel, IL
Contact	Ravit Peled
510(k) history	7 submissions · 7 cleared · 2016-2023

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k191220/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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