FDA 510(k) Premarket Notification

K230205 Veloxion System

CLEARED

Feb 24, 2023
30 days to decision

K230205 · Product code: **FJL** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k230205/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Resectoscope (FJL)
Date received	Jan 25, 2023
Decision date	Feb 24, 2023
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Corinth Medtech, Inc.**
Location	Cupertino, CA, US
Contact	George Hsu
510(k) history	7 submissions · 7 cleared · 2017-2023