FDA 510(k) Premarket Notification

K171334 Indego

CLEARED

Sep 8, 2017
123 days to decision

K171334 · Product code: **PHL** · Neurology
Source: https://www.510kdatabase.net/k171334/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Exoskeleton (PHL)
Date received	May 8, 2017
Decision date	Sep 8, 2017
Days to decision	123 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Parker Hannifin Corporation**
Location	Macedonia, OH, US
Contact	Achilleas Dorotheou
510(k) history	5 submissions · 5 cleared · 2016-2024