FDA 510(k) Premarket Notification

K173530 Indego(R)

CLEARED

Jan 31, 2018
77 days to decision

K173530 · Product code: **PHL** · Neurology
Source: https://www.510kdatabase.net/k173530/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Exoskeleton (PHL)
Date received	Nov 15, 2017
Decision date	Jan 31, 2018
Days to decision	77 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