FDA 510(k) Premarket Notification

K192607 LiteWalk

CLEARED

Dec 19, 2019
90 days to decision

K192607 · Product code: **IRP** · Physical Medicine
Source: https://www.510kdatabase.net/k192607/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Massager, Powered Inflatable Tube (IRP)
Date received	Sep 20, 2019
Decision date	Dec 19, 2019
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Viasonix , Ltd.**
Location	Yonkers, NY, US
Contact	Shlomi Deler
510(k) history	6 submissions · 6 cleared · 2011-2024