FDA 510(k) Premarket Notification

K203019 LF900

CLEARED

Jun 22, 2021
264 days to decision

K203019 · Product code: **IRP** · Physical Medicine
Source: https://www.510kdatabase.net/k203019/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Massager, Powered Inflatable Tube (IRP)
Date received	Oct 1, 2020
Decision date	Jun 22, 2021
Days to decision	264 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Daesung Maref Co., Ltd.**
Location	Buena Park, CA, US
Contact	So Su Hyeon
510(k) history	14 submissions · 14 cleared · 2011-2023