FDA 510(k) Premarket Notification

K172302 InMode PLUS System

CLEARED

Dec 8, 2017
130 days to decision

K172302 · Product code: **PBX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k172302/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Massager, Vacuum, Radio Frequency Induced Heat (PBX)
Date received	Jul 31, 2017
Decision date	Dec 8, 2017
Days to decision	130 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Inmode MD , Ltd.**
Location	Kfar Saba, IL
Contact	Amit Goren
510(k) history	21 submissions · 21 cleared · 2013-2021