FDA 510(k) Premarket Notification

K153568 InMode Plus System

CLEARED

Jul 12, 2016
211 days to decision

K153568 · Product code: **PBX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k153568/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Massager, Vacuum, Radio Frequency Induced Heat (PBX)
Date received	Dec 14, 2015
Decision date	Jul 12, 2016
Days to decision	211 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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