FDA 510(k) Premarket Notification

K151793 InMode RF System

CLEARED

Feb 19, 2016
233 days to decision

K151793 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k151793/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jul 1, 2015
Decision date	Feb 19, 2016
Days to decision	233 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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k151793/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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