FDA 510(k) Premarket Notification

K142952 InMode Diolaze Device

CLEARED

Jan 13, 2015
95 days to decision

K142952 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k142952/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Oct 10, 2014
Decision date	Jan 13, 2015
Days to decision	95 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