

**K142952 InMode Diolaze Device**Jan 13, 2015  
95 days to decisionK142952 · Product code: **GEX** · General & Plastic SurgerySource: <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	<b>Inmode MD , Ltd.</b>
Location	Kfar Saba, IL
Contact	Ahava Stein
510(k) history	21 submissions · 21 cleared · 2013-2021

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k142952/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 27, 2026