

**K153568 InMode Plus System**Jul 12, 2016  
211 days to decisionK153568 · Product code: **PBX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k153568/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k153568/> 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