

**K172302 InMode PLUS System**Dec 8, 2017  
130 days to decisionK172302 · Product code: **PBX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k172302/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Massager, Vacuum, Radio Frequency Induced Heat (PBX)
Date received	Jul 31, 2017
Decision date	Dec 8, 2017
Days to decision	130 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Inmode MD , Ltd.</b>
Location	Kfar Saba, IL
Contact	Amit Goren
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/k172302/> 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