

**K203019 LF900**Jun 22, 2021  
264 days to decisionK203019 · Product code: **IRP** · Physical MedicineSource: <https://www.510kdatabase.net/k203019/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Massager, Powered Inflatable Tube (IRP)
Date received	Oct 1, 2020
Decision date	Jun 22, 2021
Days to decision	264 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Daesung Meref Co., Ltd.</b>
Location	Buena Park, CA, US
Contact	So Su Hyeon
510(k) history	14 submissions · 14 cleared · 2011-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k203019/> 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 26, 2026