

**K220557 MoistureMeterD Compact, LymphScanner**May 27, 2022  
88 days to decisionK220557 · Product code: **OBH** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k220557/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Extracellular Fluid, Lymphedema, Extremity (OBH)
Date received	Feb 28, 2022
Decision date	May 27, 2022
Days to decision	88 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Delfin Technologies, Ltd.</b>
Location	Kuopio, FI
Contact	Jouni Nuutinen
510(k) history	2 submissions · 2 cleared · 2015-2022

**REGULATORY CONSULTANT**

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Consulting firm	<b>F Care Systems USA (Db: Delfin USA, LLC)</b>
Contact	Patrick Danciu

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k220557/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated May 27, 2026