

**K212510 7 Series Phototherapy Device**Nov 4, 2021  
86 days to decisionK212510 · Product code: **FTC** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k212510/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Ultraviolet, Dermatological (FTC)
Date received	Aug 10, 2021
Decision date	Nov 4, 2021
Days to decision	86 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Daavlin Distributing Co.</b>
Location	Bryan, OH, US
Contact	Michele Thiel
510(k) history	15 submissions · 15 cleared · 2002-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k212510/> 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 25, 2026