

**K230382 3 Series NeoLux**Mar 24, 2023  
39 days to decisionK230382 · Product code: **FTC** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k230382/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Light, Ultraviolet, Dermatological (FTC)
Date received	Feb 13, 2023
Decision date	Mar 24, 2023
Days to decision	39 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/k230382/> 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 14, 2026