

**K100378 1 SERIES MODEL 1 SERIES T2, 1 SERIES X2, 1 SERIES T4, 1 SERIES X4**Apr 28, 2010  
71 days to decisionK100378 · Product code: **FTC** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k100378/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Ultraviolet, Dermatological (FTC)
Date received	Feb 16, 2010
Decision date	Apr 28, 2010
Days to decision	71 days
Third-party review	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/k100378/>; 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