

**K111049 MOBILE VERSA LIGHT**May 25, 2012  
406 days to decisionK111049 · Product code: **FTC** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k111049/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Ultraviolet, Dermatological (FTC)
Date received	Apr 15, 2011
Decision date	May 25, 2012
Days to decision	406 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medtek Lighting Corp</b>
Location	Knoxville, TN, US
Contact	ROBERT WAGNER
510(k) history	1 submissions · 1 cleared · 2012-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k111049/> 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 June 14, 2026