

**K073503 VERSA LITE**Jan 23, 2008  
41 days to decisionK073503 · Product code: **FTC** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k073503/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Ultraviolet, Dermatological (FTC)
Date received	Dec 13, 2007
Decision date	Jan 23, 2008
Days to decision	41 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Diomedics, Inc.</b>
Location	The Woodlands, TX, US
Contact	GARY MOCNIK
510(k) history	3 submissions · 3 cleared · 1993-2008

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k073503/> 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 25, 2026