

**K982409 VES 1501-M**Aug 21, 1998  
42 days to decisionK982409 · Product code: **FWF** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k982409/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Camera, Television, Endoscopic, Without Audio (FWF)
Date received	Jul 10, 1998
Decision date	Aug 21, 1998
Days to decision	42 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Angiolaz, Inc.</b>
Location	Rockingham, VT, US
Contact	JOHN D PLUMADORE
510(k) history	10 submissions · 10 cleared · 1990-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k982409/> 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 29, 2026