

**K112154 FEMTO LDV**Mar 16, 2012  
233 days to decisionK112154 · Product code: **HQF** · Ophthalmic  
Source: <https://www.510kdatabase.net/k112154/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Ophthalmic (HQF)
Date received	Jul 27, 2011
Decision date	Mar 16, 2012
Days to decision	233 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Sie Ag,Surgical Instument Engineering</b>
Location	Greenwood Village, CO, US
Contact	Kevin Walls
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/k112154/> 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 15, 2026