

**K180526 EIDON FA**Jul 16, 2018  
139 days to decisionK180526 · Product code: **MYC** · Ophthalmic  
Source: <https://www.510kdatabase.net/k180526/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ophthalmoscope, Laser, Scanning (MYC)
Date received	Feb 27, 2018
Decision date	Jul 16, 2018
Days to decision	139 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Centervue S.P.A.</b>
Location	Padova, IT
Contact	Roberto Gabriotti
510(k) history	13 submissions · 13 cleared · 2010-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k180526/> 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