

K251507 INTEGRE LIOFeb 6, 2026
266 days to decisionK251507 · Product code: **HQF** · Ophthalmic
Source: <https://www.510kdatabase.net/k251507/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Ophthalmic (HQF)
Date received	May 16, 2025
Decision date	Feb 6, 2026
Days to decision	266 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Quantel Medical
Location	Cournon D'auvergne-Cedex, FR
Contact	Bruno Pages
Website	https://www.quantelmedical.com
510(k) history	30 submissions · 30 cleared · 2000-2026

REGULATORY CONSULTANT

Consulting firm	O'Connell Regulatory Consultants, Inc.
Contact	Maureen O'Connell

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)**510k Database** - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k251507/> Data sourced from FDA 510(k) public records (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. - Generated May 13, 2026