

**K141790 AQANIFE**Mar 6, 2015  
247 days to decisionK141790 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k141790/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jul 2, 2014
Decision date	Mar 6, 2015
Days to decision	247 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Ovesco Endoscopy AG</b>
Location	Karlsruhe, DE
Contact	TANJA R LINZER
510(k) history	14 submissions · 13 cleared · 2010-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k141790/> 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 24, 2026