

**K241833 V-GRAD**Mar 21, 2025  
269 days to decisionK241833 · Product code: **MQL** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k241833/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Media, Reproductive (MQL)
Date received	Jun 25, 2024
Decision date	Mar 21, 2025
Days to decision	269 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Vitromed GmbH</b>
Location	Jena, DE
Contact	Greg Holland
510(k) history	7 submissions · 7 cleared · 2020-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k241833/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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