

**K133912 ORIGIO SEQUENTIAL FERT, ORIGIO SEQUENTIAL FERT WITH PHENOL RED, ORIGIO SEQUENTIAL CLEAV, ORIGIO SEQUENTIAL CLEAV, ORIGIO**May 14, 2014  
142 days to decisionK133912 · Product code: **MQL** · Obstetrics & Gynecology  
Source: <https://www.510kdatabase.net/k133912/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Media, Reproductive (MQL)
Date received	Dec 23, 2013
Decision date	May 14, 2014
Days to decision	142 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Origio A/S</b>
Location	San Diego, CA, US
Contact	TOVE KJAER
510(k) history	14 submissions · 14 cleared · 2010-2020

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k133912/>; 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 20, 2026