

**K103028 EEVATM PETRI DISH**Aug 11, 2011  
302 days to decisionK103028 · Product code: **MQK** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k103028/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Labware, Assisted Reproduction (MQK)
Date received	Oct 13, 2010
Decision date	Aug 11, 2011
Days to decision	302 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Auxogyn, Inc.</b>
Location	Menlo Park, CA, US
Contact	ROBERT NEWMAN
510(k) history	4 submissions · 3 cleared · 2011-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k103028/> 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 14, 2026