

**K092579 INTRAUTERINE INSEMINATION (IUI) CATHETER**Apr 2, 2010  
228 days to decisionK092579 · Product code: **MQF** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k092579/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Assisted Reproduction (MQF)
Date received	Aug 17, 2009
Decision date	Apr 2, 2010
Days to decision	228 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Fertiligent, Ltd.</b>
Location	Indianapolis, IN, US
Contact	JOHN A STEEN
510(k) history	2 submissions · 2 cleared · 2010-2015

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k092579/> 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 5, 2026