

**K092963 GLOBAL BLASTOCYST VITRIFICATION KIT &  
VITRIFICATION THAWING KIT - BASED ON S3**Jan 14, 2011  
476 days to decisionK092963 · Product code: **MQL** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k092963/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Media, Reproductive (MQL)
Date received	Sep 25, 2009
Decision date	Jan 14, 2011
Days to decision	476 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>&lt;Genx&gt; Intl., Inc.</b>
Location	Madison, CT, US
Contact	MICHAEL D CECCHI
510(k) history	32 submissions · 32 cleared · 1996-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k092963/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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