

**K990941 NEWLIFE DISH (ETCHED OR NON-ETCHED, WITH OR WITHOUT LID), MODELS 14-52, 1452E, 1452EL, 1452L**Jul 14, 1999  
114 days to decisionK990941 · Product code: **MQK** · Obstetrics & Gynecology  
Source: <https://www.510kdatabase.net/k990941/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Labware, Assisted Reproduction (MQK)
Date received	Mar 22, 1999
Decision date	Jul 14, 1999
Days to decision	114 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Humagen Fertility Diagnostics, Inc.</b>
Location	Charlottesville, VA, US
Contact	CINDY SHOWALTER
510(k) history	5 submissions · 5 cleared · 1994-2002

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k990941/> Data sourced from FDA 510(k) public records (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. - Generated June 29, 2026