

**K073338 CELLSEARCH CIRCULATING TUMOR CELL KIT**Feb 26, 2008  
90 days to decisionK073338 · Product code: **NQI** · Pathology  
Source: <https://www.510kdatabase.net/k073338/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Immunomagnetic, Circulating Cancer Cell, Enumeration (NQI)
Date received	Nov 28, 2007
Decision date	Feb 26, 2008
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Veridex, LLC</b>
Location	Raritan, NJ, US
Contact	DEBRA J RASMUSSEN
510(k) history	12 submissions · 12 cleared · 2004-2013

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k073338/> 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