

**K050245 CELLSEARCH CIRCULATING TUMOR CELL KIT**Mar 15, 2005  
41 days to decisionK050245 · Product code: **NQI** · Pathology  
Source: <https://www.510kdatabase.net/k050245/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Immunomagnetic, Circulating Cancer Cell, Enumeration (NQI)
Date received	Feb 2, 2005
Decision date	Mar 15, 2005
Days to decision	41 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/k050245/> 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