

**K110406 CELLTRACKS AUTOPREP SYSTEM**

Jan 20, 2012  
 340 days to decision

K110406 · Product code: **NQI** · Pathology  
 Source: <https://www.510kdatabase.net/k110406/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Immunomagnetic, Circulating Cancer Cell, Enumeration (NQI)
Date received	Feb 14, 2011
Decision date	Jan 20, 2012
Days to decision	340 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Veridex, LLC</b>
Location	Raritan, NJ, US
Contact	JESSICA SHELFFO
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/k110406/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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