

**K101838 VTS1000**Feb 18, 2011  
232 days to decisionK101838 · Product code: **KZA** · General Hospital  
Source: <https://www.510kdatabase.net/k101838/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Vein Location, Liquid Crystal (KZA)
Date received	Jul 1, 2010
Decision date	Feb 18, 2011
Days to decision	232 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Vuetek Scientific, LLC</b>
Location	Marietta, GA, US
Contact	PAUL SUMNER
510(k) history	1 submissions · 1 cleared · 2011-2011

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k101838/> 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 June 19, 2026