

**K121738 MYVISIONTRACK(TM)**Feb 22, 2013  
254 days to decisionK121738 · Product code: **HOQ** · Ophthalmic  
Source: <https://www.510kdatabase.net/k121738/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Grid, Amsler (HOQ)
Date received	Jun 13, 2012
Decision date	Feb 22, 2013
Days to decision	254 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Vital Art and Science Incorporated</b>
Location	Richardson, TX, US
Contact	MICHAEL BARTLETT
510(k) history	2 submissions · 2 cleared · 2013-2015

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k121738/> 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 5, 2026