

**K140691 IG\_PLEX CELIAC DGP PANEL**Nov 6, 2014  
232 days to decisionK140691 · Product code: **MVM** · Immunology  
Source: <https://www.510kdatabase.net/k140691/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Autoantibodies, Endomysial(tissue Transglutaminase) (MVM)
Date received	Mar 19, 2014
Decision date	Nov 6, 2014
Days to decision	232 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Sqi Diagnostics Systems, Inc.</b>
Location	Toronto, On, CA
Contact	Kate Smith
510(k) history	1 submissions · 1 cleared · 2014-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k140691/> 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