

**K964299 N-ASSAY TIA MULTI V-NL**Jul 14, 1997  
258 days to decisionK964299 · Product code: **LKL** · Immunology  
Source: <https://www.510kdatabase.net/k964299/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Alpha-1-acid-glycoprotein, Antigen, Antiserum, Control (LKL)
Date received	Oct 29, 1996
Decision date	Jul 14, 1997
Days to decision	258 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Crestat Diagnostics, Inc.</b>
Location	Portland, OR, US
510(k) history	29 submissions · 29 cleared · 1990-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k964299/>; 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 28, 2026