

**K930074 IN-VITRO DIAGNOSTIC REAGENT SET**May 11, 1993  
124 days to decisionK930074 · Product code: **JGJ** · Chemistry  
Source: <https://www.510kdatabase.net/k930074/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Photometric Method, Magnesium (JGJ)
Date received	Jan 7, 1993
Decision date	May 11, 1993
Days to decision	124 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Horizon Diagnostics, Inc.</b>
Location	Jackson, MI, US
Contact	DAVID K GARCIA
510(k) history	7 submissions · 7 cleared · 1993-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k930074/> 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 July 1, 2026