

**K020234 A1CNOW FOR RX HOME USE**Feb 15, 2002  
23 days to decisionK020234 · Product code: **LCP** · Hematology  
Source: <https://www.510kdatabase.net/k020234/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Assay, Glycosylated Hemoglobin (LCP)
Date received	Jan 23, 2002
Decision date	Feb 15, 2002
Days to decision	23 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Metrika, Inc.</b>
Location	Sunnyvale, CA, US
Contact	ERIKA AMMIRATI
510(k) history	11 submissions · 11 cleared · 1999-2005

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k020234/> 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