

**K121433 GOODLIFE CS-200 (SINGLE PATIENT USE), CS-200 PROFESSIONAL (MULTIPLE PATIENT USE) BLOOD GLUCOSE MONITORING SYSTEMS**Feb 14, 2013  
276 days to decisionK121433 · Product code: LFR · Chemistry  
Source: <https://www.510kdatabase.net/k121433/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Glucose Dehydrogenase, Glucose (LFR)
Date received	May 14, 2012
Decision date	Feb 14, 2013
Days to decision	276 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Hmd Biomedical, Inc.</b>
Location	Ocean, NJ, US
Contact	JESSICA TUNG
510(k) history	10 submissions · 10 cleared · 2003-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k121433/> 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 May 27, 2026