

**K051200 HEMOGLOBIN A1C REAGENT SET**Aug 8, 2005  
90 days to decisionK051200 · Product code: **LCP** · Toxicology  
Source: <https://www.510kdatabase.net/k051200/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Assay, Glycosylated Hemoglobin (LCP)
Date received	May 10, 2005
Decision date	Aug 8, 2005
Days to decision	90 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Teco Diagnostics</b>
Location	Placentia, CA, US
Contact	JIAN VAECHES
510(k) history	51 submissions · 51 cleared · 1993-2017

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k051200/> 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 31, 2026