

K920198 GLYCOMAT HAEMOGLOBIN ANALYSERJul 21, 1992
188 days to decisionK920198 · Product code: **LCP** · Hematology
Source: <https://www.510kdatabase.net/k920198/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Assay, Glycosylated Hemoglobin (LCP)
Date received	Jan 15, 1992
Decision date	Jul 21, 1992
Days to decision	188 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Drew Scientific , Ltd.
Location	London W4 4ph Uk, GB
Contact	DREW
510(k) history	7 submissions · 7 cleared · 1992-2006

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k920198/> Data sourced from FDA 510(k) public records (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. - Generated June 28, 2026