

K870210 CLEARVIEW CW-10Apr 28, 1987
98 days to decisionK870210 · Product code: **GKR** · Hematology
Source: <https://www.510kdatabase.net/k870210/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Hemoglobin, Automated (GKR)
Date received	Jan 20, 1987
Decision date	Apr 28, 1987
Days to decision	98 days
Third-party review	No

APPLICANT

Company	Clearview Instruments, Inc.
Location	Stow, MA, US
Contact	MARIO LESLIE
510(k) history	1 submissions · 1 cleared · 1987-1987

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k870210/>, 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 29, 2026