FDA 510(k) Premarket Notification

K870210 CLEARVIEW CW-10

CLEARED

Apr 28, 1987
98 days to decision

K870210 · 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