FDA 510(k) Premarket Notification

K920198 GLYCOMAT HAEMOGLOBIN ANALYSER

CLEARED

Jul 21, 1992
188 days to decision

K920198 · 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