FDA 510(k) Premarket Notification

K802568 HEMOGLOBIN CONTROL

CLEARED

Nov 12, 1980
26 days to decision

K802568 · Product code: **KHG** · Hematology
Source: https://www.510kdatabase.net/k802568/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Whole Blood Hemoglobin Determination (KHG)
Date received	Oct 17, 1980
Decision date	Nov 12, 1980
Days to decision	26 days
Third-party review	No

APPLICANT

Company	**Marilyn L. Amick**
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1978-1980

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k802568/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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