FDA 510(k) Premarket Notification

K820596 CYANMETHEMOGLOBIN TEST

CLEARED

Mar 26, 1982
22 days to decision

K820596 · Product code: **KHG** · Hematology
Source: https://www.510kdatabase.net/k820596/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Whole Blood Hemoglobin Determination (KHG)
Date received	Mar 4, 1982
Decision date	Mar 26, 1982
Days to decision	22 days
Third-party review	No

APPLICANT

Company	**Boehringer Mannheim Corp.**
Location	Mchenry, IL, US
510(k) history	340 submissions · 340 cleared · 1976-1999

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k820596/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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