FDA 510(k) Premarket Notification

K820037 SICKLE-CHECK

CLEARED

Feb 4, 1982
28 days to decision

K820037 · Product code: **JCM** · Hematology
Source: https://www.510kdatabase.net/k820037/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Control, Hemoglobin, Abnormal (JCM)
Date received	Jan 7, 1982
Decision date	Feb 4, 1982
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**Diagnostic Technology, Inc.**
Location	Mchenry, IL, US
510(k) history	28 submissions · 28 cleared · 1979-1991