

K820037 SICKLE-CHECKFeb 4, 1982
28 days to decisionK820037 · 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

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Device record: <https://www.510kdatabase.net/k820037/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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