

K834372 PHA-1 PROGRAMMABLE HEMATOLOGY ANALYZFeb 10, 1984
71 days to decisionK834372 · Product code: **GKL** · Hematology
Source: <https://www.510kdatabase.net/k834372/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Counter, Cell, Automated (particle Counter) (GKL)
Date received	Dec 1, 1983
Decision date	Feb 10, 1984
Days to decision	71 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/k834372/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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