FDA 510(k) Premarket Notification

# K834372 PHA-1 PROGRAMMABLE HEMATOLOGY ANALYZ

**CLEARED**

Feb 10, 1984
71 days to decision

K834372 · Product code: **GKL** · Hematology
Source: https://www.510kdatabase.net/k834372/

## SUBMISSION DETAILS

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|---|---|
| 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 |