

K823522 DILULAB 230Dec 28, 1982
29 days to decisionK823522 · Product code: **GKH** · Hematology
Source: <https://www.510kdatabase.net/k823522/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Apparatus, Automated Blood Cell Diluting (GKH)
Date received	Nov 29, 1982
Decision date	Dec 28, 1982
Days to decision	29 days
Third-party review	No

APPLICANT

Company	Innovative Medical Systems, Inc.
Location	Mchenry, IL, US
510(k) history	22 submissions · 22 cleared · 1978-1997

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Device record: <https://www.510kdatabase.net/k823522/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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