FDA 510(k) Premarket Notification

K823522 DILULAB 230

CLEARED

Dec 28, 1982
29 days to decision

K823522 · 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