FDA 510(k) Premarket Notification

K842921 REES & ECKER DILUTION FLUID

CLEARED

Sep 7, 1984
44 days to decision

K842921 · Product code: **GLG** · Hematology
Source: https://www.510kdatabase.net/k842921/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Platelet Counting, Manual (GLG)
Date received	Jul 25, 1984
Decision date	Sep 7, 1984
Days to decision	44 days
Third-party review	No

APPLICANT

Company	**E K Ind., Inc.**
Location	Mchenry, IL, US
510(k) history	50 submissions · 49 cleared · 1984-1984