FDA 510(k) Premarket Notification

K781969 BLOOD, HEMATOLOGY REFERENCE CONTROL

CLEARED

Jan 3, 1979
40 days to decision

K781969 · Product code: **JPK** · Hematology
Source: https://www.510kdatabase.net/k781969/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mixture, Hematology Quality Control (JPK)
Date received	Nov 24, 1978
Decision date	Jan 3, 1979
Days to decision	40 days
Third-party review	No

APPLICANT

Company	**Coulter Electronics, Inc.**
Location	Mchenry, IL, US
510(k) history	101 submissions · 101 cleared · 1976-1994