FDA 510(k) Premarket Notification

K823930 HEMATOLOGY QUALITY CONTROL MIXTURE

CLEARED

Feb 1, 1983
34 days to decision

K823930 · Product code: **GJP** · Hematology
Source: https://www.510kdatabase.net/k823930/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Control, Platelet (GJP)
Date received	Dec 29, 1982
Decision date	Feb 1, 1983
Days to decision	34 days
Third-party review	No

APPLICANT

Company	**Fisher Scientific Co., LLC**
Location	Mchenry, IL, US
510(k) history	89 submissions · 89 cleared · 1976-1993