FDA 510(k) Premarket Notification

K820533 B4182-1

CLEARED

Apr 8, 1982
37 days to decision

K820533 · Product code: **GIE** · Hematology
Source: https://www.510kdatabase.net/k820533/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Fibrometer (GIE)
Date received	Mar 2, 1982
Decision date	Apr 8, 1982
Days to decision	37 days
Third-party review	No

APPLICANT

Company	**American Scientific Products**
Location	Mchenry, IL, US
510(k) history	28 submissions · 28 cleared · 1981-1986