

K820533 B4182-1Apr 8, 1982
37 days to decisionK820533 · 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

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Device record: <https://www.510kdatabase.net/k820533/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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