

K802117 FIBRI QUIKSep 26, 1980
22 days to decisionK802117 · Product code: **GJA** · Hematology
Source: <https://www.510kdatabase.net/k802117/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Test, Thrombin Time (GJA)
Date received	Sep 4, 1980
Decision date	Sep 26, 1980
Days to decision	22 days
Third-party review	No

APPLICANT

Company	General Diagnostics
Location	Mchenry, IL, US
510(k) history	39 submissions · 39 cleared · 1976-1988

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Device record: <https://www.510kdatabase.net/k802117/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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