

K811069 FIBRINOGEN CONTROL PLASMAJul 10, 1981
81 days to decisionK811069 · Product code: **KQJ** · Hematology
Source: <https://www.510kdatabase.net/k811069/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Fibrinogen Determination (KQJ)
Date received	Apr 20, 1981
Decision date	Jul 10, 1981
Days to decision	81 days
Third-party review	No

APPLICANT

Company	American Dade
Location	Mchenry, IL, US
510(k) history	149 submissions · 149 cleared · 1980-1987

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Device record: <https://www.510kdatabase.net/k811069/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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