

K812629 ACA FIBRINOGEN TEST PACKOct 8, 1981
24 days to decisionK812629 · Product code: **KQJ** · Hematology
Source: <https://www.510kdatabase.net/k812629/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Fibrinogen Determination (KQJ)
Date received	Sep 14, 1981
Decision date	Oct 8, 1981
Days to decision	24 days
Third-party review	No

APPLICANT

Company	E.I. Dupont DE Nemours & Co., Inc.
Location	Mchenry, IL, US
510(k) history	253 submissions · 252 cleared · 1976-1996

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Device record: <https://www.510kdatabase.net/k812629/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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