

K812628 ACA FIBRINOGEN CALIBRATORNov 24, 1981
70 days to decisionK812628 · Product code: **GFX** · Hematology
Source: <https://www.510kdatabase.net/k812628/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Fibrinogen Standard (GFX)
Date received	Sep 15, 1981
Decision date	Nov 24, 1981
Days to decision	70 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/k812628/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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