

K810664 ISOLATOR SYSTEMMar 31, 1981
20 days to decisionK810664 · Product code: **JKA** · Hematology
Source: <https://www.510kdatabase.net/k810664/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubes, Vials, Systems, Serum Separators, Blood Collection (JKA)
Date received	Mar 11, 1981
Decision date	Mar 31, 1981
Days to decision	20 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/k810664/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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