

K790323 COAG-A-MATE 2001Mar 21, 1979
34 days to decisionK790323 · Product code: **JPA** · Hematology
Source: <https://www.510kdatabase.net/k790323/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Multipurpose For In Vitro Coagulation Studies (JPA)
Date received	Feb 15, 1979
Decision date	Mar 21, 1979
Days to decision	34 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/k790323/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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