

K023141 MODIFICATION TO STANDARD HUMAN PLASMAJan 16, 2003
118 days to decisionK023141 · Product code: **JPA** · Hematology
Source: <https://www.510kdatabase.net/k023141/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Multipurpose For In Vitro Coagulation Studies (JPA)
Date received	Sep 20, 2002
Decision date	Jan 16, 2003
Days to decision	118 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Dade Behring, Inc.
Location	Newark,, DE, US
Contact	DONNA A WOLF
510(k) history	343 submissions · 343 cleared · 1978-2010

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