

**K790204 PLASMA, STANDARDIZED NORMAL**Mar 12, 1979  
41 days to decisionK790204 · Product code: **JPA** · Hematology  
Source: <https://www.510kdatabase.net/k790204/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Multipurpose For In Vitro Coagulation Studies (JPA)
Date received	Jan 30, 1979
Decision date	Mar 12, 1979
Days to decision	41 days
Third-party review	No

**APPLICANT**

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Company	<b>Dade, Baxter Travenol Diagnostics, Inc.</b>
Location	Mchenry, IL, US
510(k) history	44 submissions · 44 cleared · 1976-1980

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k790204/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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