FDA 510(k) Premarket Notification

K790204 PLASMA, STANDARDIZED NORMAL

CLEARED

Mar 12, 1979
41 days to decision

K790204 · Product code: **JPA** · Hematology
Source: https://www.510kdatabase.net/k790204/

SUBMISSION DETAILS

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

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