FDA 510(k) Premarket Notification

K860342 MICROSAMPLE COAGULATION ANALYSER, MODEL 110WP

CLEARED

Mar 4, 1986
34 days to decision

K860342 · Product code: **JPA** · Hematology
Source: https://www.510kdatabase.net/k860342/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Multipurpose For In Vitro Coagulation Studies (JPA)
Date received	Jan 29, 1986
Decision date	Mar 4, 1986
Days to decision	34 days
Third-party review	No

APPLICANT

Company	**Bio/Data Corp.**
Location	Mchenry, IL, US
Contact	JAMES EICHELBERGER
510(k) history	37 submissions · 37 cleared · 1977-2000

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k860342/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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