FDA 510(k) Premarket Notification

K920895 DADE(R) PROTHROMBIN FRAGMENT F1.2 ELISA

CLEARED

Jun 4, 1992
99 days to decision

K920895 · Product code: **MIF** · Hematology
Source: https://www.510kdatabase.net/k920895/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prothrombin Fragment 1.2 (MIF)
Date received	Feb 26, 1992
Decision date	Jun 4, 1992
Days to decision	99 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Baxter Diagnostics, Inc.**
Location	Miami, FL, US
Contact	ALODIA M RUIZ
Website	https://www.baxter.com/
510(k) history	72 submissions · 72 cleared · 1988-1995

Baxter Diagnostics, Inc. is a diagnostic device manufacturer based in Miami, US. The company specialized in microbiology and chemistry diagnostic solutions. Baxter Diagnostics received FDA 510(k) clearances from total submissions between 1988 and 1995. The company's regulatory focus centered on microbiology devices, particularly dried antimicrobial susceptibility testing panels and related diagnostic assays. This historical record reflects the company's core expertise in microbial identification and resistance testing. The company is no longer active in FDA 510(k) submiss...