

K914818 DADE(R) ALPHA 2 ANTIPLASMIN(2 2-AP) CHROMO ASSAYJan 15, 1992
82 days to decisionK914818 · Product code: **GGP** · Hematology
Source: <https://www.510kdatabase.net/k914818/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Test, Qualitative And Quantitative Factor Deficiency (GGP)
Date received	Oct 25, 1991
Decision date	Jan 15, 1992
Days to decision	82 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Baxter Diagnostics, Inc.
Location	Miami, FL, US
Contact	NANCY A HORNBAKER
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 The company is no longer active in FDA 510(k) submiss...

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