

**K920981 RISTOCETIN COFACTOR ASSAY/VON WILLEBRAND
FACT ASSA**Apr 17, 1992
46 days to decisionK920981 · Product code: **GGP** · Hematology
Source: <https://www.510kdatabase.net/k920981/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Test, Qualitative And Quantitative Factor Deficiency (GGP)
Date received	Mar 2, 1992
Decision date	Apr 17, 1992
Days to decision	46 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Medical Diagnostic Technologies, Inc.
Location	Augusta, GA, US
Contact	MICHAEL D BICK
510(k) history	57 submissions · 57 cleared · 1985-2006

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k920981/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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