FDA 510(k) Premarket Notification

K904073 COAGULATION REFERENCE PLASMA, ABNORMAL

CLEARED

Oct 22, 1990
47 days to decision

K904073 · Product code: **GGC** · Hematology
Source: https://www.510kdatabase.net/k904073/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Control, Plasma, Abnormal (GGC)
Date received	Sep 5, 1990
Decision date	Oct 22, 1990
Days to decision	47 days
Third-party review	No

APPLICANT

Company	**Medi-Tech, Inc.**
Location	Mchenry, IL, US
Contact	MICHAEL BICK
510(k) history	36 submissions · 35 cleared · 1978-1996

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k904073/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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