

**K923593 HEMOCHRON/FACTOR VI FIBRINOGEN ASSAY AND
CONT PLAS**Sep 22, 1992
64 days to decisionK923593 · Product code: **KQJ** · Hematology
Source: <https://www.510kdatabase.net/k923593/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Fibrinogen Determination (KQJ)
Date received	Jul 20, 1992
Decision date	Sep 22, 1992
Days to decision	64 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	International Technidyne Corp.
Location	Walker, MI, US
Contact	MATT LESNIESKI
510(k) history	47 submissions · 47 cleared · 1983-2013

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k923593/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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