

K791750 COUNTER-CHECK PLUSOct 11, 1979
36 days to decisionK791750 · Product code: **JPK** · Hematology
Source: <https://www.510kdatabase.net/k791750/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mixture, Hematology Quality Control (JPK)
Date received	Sep 5, 1979
Decision date	Oct 11, 1979
Days to decision	36 days
Third-party review	No

APPLICANT

Company	Diagnostic Technology, Inc.
Location	Mchenry, IL, US
510(k) history	28 submissions · 28 cleared · 1979-1991

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Device record: <https://www.510kdatabase.net/k791750/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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