

K780835 ARANTROLJun 22, 1978
30 days to decisionK780835 · Product code: **JPK** · Hematology
Source: <https://www.510kdatabase.net/k780835/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mixture, Hematology Quality Control (JPK)
Date received	May 23, 1978
Decision date	Jun 22, 1978
Days to decision	30 days
Third-party review	No

APPLICANT

Company	Aranco Diagnostic Laboratories, Inc.
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1978-1978

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Device record: <https://www.510kdatabase.net/k780835/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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