FDA 510(k) Premarket Notification

K780835 ARANTROL

CLEARED

Jun 22, 1978
30 days to decision

K780835 · 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