FDA 510(k) Premarket Notification

K801815 AUTOLET

CLEARED

Oct 10, 1980
72 days to decision

K801815 · Product code: **JCA** · Hematology
Source: https://www.510kdatabase.net/k801815/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Bleeding Time (JCA)
Date received	Jul 30, 1980
Decision date	Oct 10, 1980
Days to decision	72 days
Third-party review	No

APPLICANT

Company	**Ulster Scientific, Inc.**
Location	Mchenry, IL, US
510(k) history	20 submissions · 20 cleared · 1980-1994