

K801815 AUTOLETOct 10, 1980
72 days to decisionK801815 · 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

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Device record: <https://www.510kdatabase.net/k801815/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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