

K811777 HELENA PLASMINOGEN QUIPLATE PROCEDUREJul 2, 1981
9 days to decisionK811777 · Product code: **DDX** · Immunology
Source: <https://www.510kdatabase.net/k811777/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plasminogen, Antigen, Antiserum, Control (DDX)
Date received	Jun 23, 1981
Decision date	Jul 2, 1981
Days to decision	9 days
Third-party review	No

APPLICANT

Company	Helena Laboratories
Location	Mchenry, IL, US
510(k) history	280 submissions · 280 cleared · 1978-2013

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Device record: <https://www.510kdatabase.net/k811777/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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