FDA 510(k) Premarket Notification

K800925 VERIFY S PROD. #34141

CLEARED

Jul 21, 1980
91 days to decision

K800925 · Product code: **GGW** · Hematology
Source: https://www.510kdatabase.net/k800925/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Test, Time, Partial Thromboplastin (GGW)
Date received	Apr 21, 1980
Decision date	Jul 21, 1980
Days to decision	91 days
Third-party review	No

APPLICANT

Company	**General Diagnostics**
Location	Mchenry, IL, US
510(k) history	39 submissions · 39 cleared · 1976-1988

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k800925/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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