FDA 510(k) Premarket Notification

K812975 LANCER COAGENT HUMAN VACTOR VIII PLASMA

CLEARED

Nov 16, 1981
24 days to decision

K812975 · Product code: **GGP** · Hematology
Source: https://www.510kdatabase.net/k812975/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Test, Qualitative And Quantitative Factor Deficiency (GGP)
Date received	Oct 23, 1981
Decision date	Nov 16, 1981
Days to decision	24 days
Third-party review	No

APPLICANT

Company	**Sherwood Medical Co.**
Location	Mchenry, IL, US
510(k) history	191 submissions · 177 cleared · 1976-1998

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k812975/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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