

K821963 EIA-F8 ENZYMEIMMUNOASSAY KIT FOR QUANT.Jul 27, 1982
26 days to decisionK821963 · Product code: **GGP** · Hematology
Source: <https://www.510kdatabase.net/k821963/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Test, Qualitative And Quantitative Factor Deficiency (GGP)
Date received	Jul 1, 1982
Decision date	Jul 27, 1982
Days to decision	26 days
Third-party review	No

APPLICANT

Company	Intl. Immunoassay Laboratories, Inc.
Location	Mchenry, IL, US
510(k) history	27 submissions · 27 cleared · 1980-1992

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Device record: <https://www.510kdatabase.net/k821963/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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