

K840500 SEDIPLASTFeb 21, 1984
15 days to decisionK840500 · Product code: **JPH** · Hematology
Source: <https://www.510kdatabase.net/k840500/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Test, Erythrocyte Sedimentation Rate (JPH)
Date received	Feb 6, 1984
Decision date	Feb 21, 1984
Days to decision	15 days
Third-party review	No

APPLICANT

Company	Precision Technology, Inc.
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1984-1984

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Device record: <https://www.510kdatabase.net/k840500/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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