FDA 510(k) Premarket Notification

K840500 SEDIPLAST

CLEARED

Feb 21, 1984
15 days to decision

K840500 · 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