

K841366 DIA FIBRINJun 1, 1984
60 days to decisionK841366 · Product code: **KQJ** · Hematology
Source: <https://www.510kdatabase.net/k841366/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Fibrinogen Determination (KQJ)
Date received	Apr 2, 1984
Decision date	Jun 1, 1984
Days to decision	60 days
Third-party review	No

APPLICANT

Company	Diatech, Inc.
Location	Mchenry, IL, US
510(k) history	17 submissions · 17 cleared · 1984-1988

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Device record: <https://www.510kdatabase.net/k841366/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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