FDA 510(k) Premarket Notification

K841366 DIA FIBRIN

CLEARED

Jun 1, 1984
60 days to decision

K841366 · 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