

**K800826 FIBRINOGEN SCREEN**May 30, 1980  
46 days to decisionK800826 · Product code: **KQJ** · Hematology  
Source: <https://www.510kdatabase.net/k800826/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)     |
| Submission type       | Traditional                            |
| Device classification | System, Fibrinogen Determination (KQJ) |
| Date received         | Apr 14, 1980                           |
| Decision date         | May 30, 1980                           |
| Days to decision      | 46 days                                |
| Third-party review    | No                                     |

**APPLICANT**

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| Company        | <b>Pacific Hemostasis</b>               |
| Location       | Mchenry, IL, US                         |
| 510(k) history | 29 submissions · 29 cleared · 1978-2000 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k800826/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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