

**K781814 SULFATE, FIBRIQUIK PROTAMINE**Nov 8, 1978  
13 days to decisionK781814 · Product code: **JBN** · Hematology  
Source: <https://www.510kdatabase.net/k781814/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Fibrin Monomer Paracoagulation (JBN)
Date received	Oct 26, 1978
Decision date	Nov 8, 1978
Days to decision	13 days
Third-party review	No

**APPLICANT**

---

Company	<b>General Diagnostics</b>
Location	Mchenry, IL, US
510(k) history	39 submissions · 39 cleared · 1976-1988

---

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

Device record: <https://www.510kdatabase.net/k781814/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 29, 2026