

**K823930 HEMATOLOGY QUALITY CONTROL MIXTURE**Feb 1, 1983  
34 days to decisionK823930 · Product code: **GJP** · Hematology  
Source: <https://www.510kdatabase.net/k823930/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Control, Platelet (GJP)
Date received	Dec 29, 1982
Decision date	Feb 1, 1983
Days to decision	34 days
Third-party review	No

**APPLICANT**

---

Company	<b>Fisher Scientific Co., LLC</b>
Location	Mchenry, IL, US
510(k) history	89 submissions · 89 cleared · 1976-1993

---

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

Device record: <https://www.510kdatabase.net/k823930/> 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 May 14, 2026