

K830804 LABILE REMOVING HEMOLYZING REAGENTMay 4, 1983
51 days to decisionK830804 · Product code: **JBD** · Hematology
Source: <https://www.510kdatabase.net/k830804/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Analysis, Electrophoretic Hemoglobin (JBD)
Date received	Mar 14, 1983
Decision date	May 4, 1983
Days to decision	51 days
Third-party review	No

APPLICANT

Company	Corning Medical & Scientific
Location	Mchenry, IL, US
Website	https://www.corning.com
510(k) history	111 submissions · 111 cleared · 1976-1988

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k830804/>; Data sourced from FDA 510(k) public records (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. - Generated May 14, 2026