FDA 510(k) Premarket Notification

K803095 COPPER SULFATE SOLUTION SP. GR.1.055

CLEARED

Mar 6, 1981
88 days to decision

K803095 · Product code: **KSL** · Hematology
Source: https://www.510kdatabase.net/k803095/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Solution, Copper Sulfate For Specific Gravity Determinations (KSL)
Date received	Dec 8, 1980
Decision date	Mar 6, 1981
Days to decision	88 days
Third-party review	No

APPLICANT

Company	**Accra Laboratories, Inc.**
Location	Mchenry, IL, US
510(k) history	20 submissions · 20 cleared · 1981-1985

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k803095/ 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