

K931935 ENDPOINT 500 PERACETIC ACID TEST INDICATORSFeb 7, 1994
294 days to decisionK931935 · Product code: **JOJ** · General HospitalSource: <https://www.510kdatabase.net/k931935/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Indicator, Physical/chemical Sterilization Process (JOJ)
Date received	Apr 19, 1993
Decision date	Feb 7, 1994
Days to decision	294 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Minntech Corp.
Location	Minneapolis, MN, US
Contact	MARK ALDANA
510(k) history	33 submissions · 33 cleared · 1987-2012

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k931935/> 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 June 13, 2026