

K803199 SUTURE REMOVAL KITJan 15, 1981
29 days to decisionK803199 · Product code: **KDD** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k803199/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Kit, Surgical Instrument, Disposable (KDD)
Date received	Dec 17, 1980
Decision date	Jan 15, 1981
Days to decision	29 days
Third-party review	No

APPLICANT

Company	Walpak Co.
Location	Mchenry, IL, US
510(k) history	11 submissions · 10 cleared · 1976-1981

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k803199/> 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 July 4, 2026