

K801623 CAVITRON ULTRASONIC SURG. ASPIR. CUSANov 18, 1980
126 days to decisionK801623 · Product code: **LFL** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k801623/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Ultrasonic Surgical (LFL)
Date received	Jul 15, 1980
Decision date	Nov 18, 1980
Days to decision	126 days
Third-party review	No

APPLICANT

Company	Cavitron Corp.
Location	Mchenry, IL, US
510(k) history	32 submissions · 32 cleared · 1976-1981

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k801623/> 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 28, 2026