

K803156 SURGITEK MEDI-LOOPSJan 15, 1981
34 days to decisionK803156 · Product code: **KDC** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k803156/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Surgical, Disposable (KDC)
Date received	Dec 12, 1980
Decision date	Jan 15, 1981
Days to decision	34 days
Third-party review	No

APPLICANT

Company	Medical Engineering Corp.
Location	Mchenry, IL, US
510(k) history	28 submissions · 28 cleared · 1977-1993

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k803156/> 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 27, 2026