

K923176 SURGICAL INSTRUMENT KIT, DISPOSABLEJun 17, 1993
370 days to decisionK923176 · Product code: **KDD** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k923176/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Kit, Surgical Instrument, Disposable (KDD)
Date received	Jun 12, 1992
Decision date	Jun 17, 1993
Days to decision	370 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Ulti-Med Intl., Inc.
Location	Mchenry, IL, US
Contact	DAVID INSCO
510(k) history	21 submissions · 18 cleared · 1983-2011

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k923176/> 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 19, 2026