

**K920989 SURGICAL INSTRUMENT KIT, DISPOSABLE**Oct 19, 1992  
231 days to decisionK920989 · Product code: **KDD** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k920989/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - KD
Submission type	Traditional
Device classification	Kit, Surgical Instrument, Disposable (KDD)
Date received	Mar 2, 1992
Decision date	Oct 19, 1992
Days to decision	231 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k920989/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 19, 2026