

**K951983 IMNI-II**Jun 20, 1995  
54 days to decisionK951983 · Product code: **LRJ** · General HospitalSource: <https://www.510kdatabase.net/k951983/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Disinfectant, Medical Devices (LRJ)
Date received	Apr 27, 1995
Decision date	Jun 20, 1995
Days to decision	54 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Cottrell, Ltd.</b>
Location	Englewood, CO, US
Contact	JACK SCOVILLE
510(k) history	9 submissions · 9 cleared · 1990-1999

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k951983/> 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 July 3, 2026