

**K881151 MICROSRAPE**Apr 6, 1988  
20 days to decisionK881151 · Product code: **LXK** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k881151/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Scrapper, Skin Specimen (LXK)
Date received	Mar 17, 1988
Decision date	Apr 6, 1988
Days to decision	20 days
Third-party review	No

**APPLICANT**

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Company	<b>Ortho Pharmaceutical Corp.</b>
Location	Walker, MI, US
Contact	RUSSELL J HUME
510(k) history	9 submissions · 9 cleared · 1979-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k881151/>, 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 29, 2026