

**K921022 DISPOSABLE CURETTE**Dec 3, 1992  
275 days to decisionK921022 · Product code: **MDM** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k921022/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Manual, Surgical, General Use (MDM)
Date received	Mar 3, 1992
Decision date	Dec 3, 1992
Days to decision	275 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Dermtec, Inc.</b>
Location	Kershaw, SC, US
Contact	BREWER, JR.
510(k) history	1 submissions · 1 cleared · 1992-1992

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k921022/> 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 6, 2026