

**K861285 SKIN MESH DEVICE**Apr 17, 1986  
10 days to decisionK861285 · Product code: **GFD** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k861285/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dermatome (GFD)
Date received	Apr 7, 1986
Decision date	Apr 17, 1986
Days to decision	10 days
Third-party review	No

**APPLICANT**

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Company	<b>Precision Modified Devices</b>
Location	Salt Lake City, UT, US
Contact	KATHY M SMITH
510(k) history	1 submissions · 1 cleared · 1986-1986

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k861285/>; 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 8, 2026