

**K896531 NOVADERM**Apr 13, 1990  
148 days to decisionK896531 · Product code: **KMF** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k896531/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bandage, Liquid (KMF)
Date received	Nov 16, 1989
Decision date	Apr 13, 1990
Days to decision	148 days
Third-party review	No

**APPLICANT**

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Company	<b>Syncare Corp.</b>
Location	Charleston, SC, US
Contact	TODD DERBIN
510(k) history	1 submissions · 1 cleared · 1990-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k896531/>, 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 7, 2026