

**K954918 COMBIDERM ACD**Dec 26, 1995  
61 days to decisionK954918 · Product code: **KMF** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k954918/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bandage, Liquid (KMF)
Date received	Oct 26, 1995
Decision date	Dec 26, 1995
Days to decision	61 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Convatec, A Bristol-Myers Squibb Co.</b>
Location	Skillman, NJ, US
Contact	MCFORD ZAMORE
510(k) history	10 submissions · 5 cleared · 1994-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k954918/> 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 30, 2026