

**K980382 NON-WOVEN COMPRESS**Mar 9, 1998  
35 days to decisionK980382 · Product code: **KMF** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k980382/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Bandage, Liquid (KMF)
Date received	Feb 2, 1998
Decision date	Mar 9, 1998
Days to decision	35 days
Third-party review	No

**APPLICANT**

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Company	<b>Convatec, A Division of E.R. Squibb &amp; Sons</b>
Location	Walker, MI, US
Contact	ADRIENNE MCNALLY
510(k) history	81 submissions · 68 cleared · 1982-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k980382/>; 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 1, 2026