

**K022645 CURAD**Nov 5, 2002  
89 days to decisionK022645 · Product code: **KMF** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k022645/>**SUBMISSION DETAILS**

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|                       |                                    |
|-----------------------|------------------------------------|
| Decision              | Substantially Equivalent (Cleared) |
| Submission type       | Traditional                        |
| Device classification | Bandage, Liquid (KMF)              |
| Date received         | Aug 8, 2002                        |
| Decision date         | Nov 5, 2002                        |
| Days to decision      | 89 days                            |
| Third-party review    | No                                 |
| Summary / Statement   | Summary                            |

**APPLICANT**

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|----------------|---|
| Company        | <b>Beiersdorf AG</b>  |
| Location       | North Attleboro, MA, US   |
| Contact        | DANIEL J DILLON   |
| Website        | <a href="http://www.beiersdorf.com/">http://www.beiersdorf.com/</a> |
| 510(k) history | 3 submissions · 3 cleared · 2002-2004                               |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k022645/> 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 3, 2026