

**K960615 GAUZE COMPRESS**Nov 25, 1996  
286 days to decisionK960615 · Product code: **EFQ** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k960615/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Gauze/sponge, Internal (EFQ)
Date received	Feb 13, 1996
Decision date	Nov 25, 1996
Days to decision	286 days
Third-party review	No

**APPLICANT**

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Company	<b>Certified Safety Mfg., Inc.</b>
Location	Kansas City, MO, US
Contact	HOWARD GERSON
510(k) history	17 submissions · 14 cleared · 1994-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k960615/>; 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 May 25, 2026