

**K953810 FLUFF GAUZE BANDAGE, STERILE & NON-STERILE**Sep 14, 1995  
37 days to decisionK953810 · Product code: **EFQ** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k953810/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Gauze/sponge, Internal (EFQ)
Date received	Aug 8, 1995
Decision date	Sep 14, 1995
Days to decision	37 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>K. W. Griffen Co.</b>
Location	Stamford, CT, US
Contact	DAVID M PIERATOS
510(k) history	4 submissions · 3 cleared · 1987-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k953810/> 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 5, 2026