

**K881516 COMPRESAID**Apr 29, 1988  
18 days to decisionK881516 · Product code: **FZF** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k881516/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Splint, Extremity, Inflatable, External (FZF)
Date received	Apr 11, 1988
Decision date	Apr 29, 1988
Days to decision	18 days
Third-party review	No

**APPLICANT**

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Company	<b>R. Evans Corp.</b>
Location	Phoenix, AZ, US
Contact	ROBERT J EVANS
510(k) history	1 submissions · 1 cleared · 1988-1988

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k881516/>; 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 8, 2026