FDA 510(k) Premarket Notification

K881516 COMPRESAID

CLEARED

Apr 29, 1988
18 days to decision

K881516 · Product code: **FZF** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k881516/

SUBMISSION DETAILS

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

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