FDA 510(k) Premarket Notification

K013683 MODIFICATION TO: ABBOTT VASCULAR SUTURE ANASTOMOSIS DEVICE

CLEARED

Dec 6, 2001
29 days to decision

K013683 · Product code: **GAW** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k013683/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Suture, Nonabsorbable, Synthetic, Polypropylene (GAW)
Date received	Nov 7, 2001
Decision date	Dec 6, 2001
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Abbott Vascular, Inc.**
Location	Redwood, CA, US
Contact	PATTY HEVEY
510(k) history	20 submissions · 17 cleared · 2000-2014