

K863570 LE VEEN WOUND SPLINTSep 29, 1986
17 days to decisionK863570 · Product code: **KGS** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k863570/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Retention Device, Suture (KGS)
Date received	Sep 12, 1986
Decision date	Sep 29, 1986
Days to decision	17 days
Third-party review	No

APPLICANT

Company	Tek-Aid, Inc.
Location	Charleston, SC, US
Contact	SUSAN J SMOAK
510(k) history	1 submissions · 1 cleared · 1986-1986

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k863570/>, Data sourced from FDA 510(k) public records (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. - Generated July 8, 2026