

K921143 PNEUMO-WEDGESep 11, 1992
184 days to decisionK921143 · Product code: **FQO** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k921143/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Table, Operating-room, Ac-powered (FQO)
Date received	Mar 11, 1992
Decision date	Sep 11, 1992
Days to decision	184 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Kansas Creative Devices, Inc.
Location	Laewood, KS, US
Contact	JEFFREY S HOWELL
510(k) history	1 submissions · 1 cleared · 1992-1992

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k921143/> 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 7, 2026