FDA 510(k) Premarket Notification

K921143 PNEUMO-WEDGE

CLEARED

Sep 11, 1992
184 days to decision

K921143 · Product code: **FQO** · General & Plastic Surgery
Source: 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