FDA 510(k) Premarket Notification

K924700 FISK HEADREST

CLEARED

Jan 15, 1993
120 days to decision

K924700 · Product code: **FQO** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k924700/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Table, Operating-room, Ac-powered (FQO)
Date received	Sep 17, 1992
Decision date	Jan 15, 1993
Days to decision	120 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Rhino Mfg.**
Location	Rancho Cordova, CA, US
Contact	SHANE FISK
510(k) history	2 submissions · 2 cleared · 1992-1993