

K924700 FISK HEADRESTJan 15, 1993
120 days to decisionK924700 · Product code: **FQO** · General & Plastic SurgerySource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k924700/> 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 6, 2026