

K121929 ZEPHYR & X-SERIES PATIENT TRANSFER SLEDJan 17, 2013
199 days to decisionK121929 · Product code: **FRZ** · Radiology
Source: <https://www.510kdatabase.net/k121929/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Patient Transfer, Powered (FRZ)
Date received	Jul 2, 2012
Decision date	Jan 17, 2013
Days to decision	199 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Diacor, Inc.
Location	Salt Lake City, UT, US
Contact	KEVIN ANDERSON
510(k) history	10 submissions · 10 cleared · 1986-2013

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k121929/> 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 May 17, 2026