FDA 510(k) Premarket Notification

K121929 ZEPHYR 'X-SERIES PATIENT TRANSFER SLED

CLEARED

Jan 17, 2013
199 days to decision

K121929 · 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