FDA 510(k) Premarket Notification

K123630 FRESENIUS LIBERTY CYCLER

CLEARED

Sep 9, 2013
290 days to decision

K123630 · Product code: **FKX** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k123630/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Peritoneal, Automatic Delivery (FKX)
Date received	Nov 23, 2012
Decision date	Sep 9, 2013
Days to decision	290 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Fresenius Medical Care North America, Design Cente**
Location	Waltham,, MA, US
Contact	DENISE OPPERMANN
510(k) history	5 submissions · 5 cleared · 2010-2013