FDA 510(k) Premarket Notification

K132220 PJ'S SERENITY

CLEARED

Feb 20, 2015
583 days to decision

K132220 · Product code: **HGX** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k132220/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Breast, Powered (HGX)
Date received	Jul 17, 2013
Decision date	Feb 20, 2015
Days to decision	583 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Limerick, Inc.**
Location	Burbank, CA, US
Contact	PATRICIA KELLY
510(k) history	4 submissions · 4 cleared · 2002-2015