FDA 510(k) Premarket Notification

K141200 PADI-LOCK

CLEARED

Oct 2, 2014
146 days to decision

K141200 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k141200/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	May 9, 2014
Decision date	Oct 2, 2014
Days to decision	146 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Padi-Lock, LLC**
Location	Great Neck, NY, US
Contact	MARIA GRIFFIN
510(k) history	1 submissions · 1 cleared · 2014-2014