CLEARED

K122404 MODEL NUMBER: RMS1-2604, RMS1-2614, RMS1-2404, RMS1-2414, RMS2-2604, RMS2-2614, RMS2-2404, RMS2-2414, RMS3-2604, RMS3-26

May 6, 2013
272 days to decision

K122404 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k122404/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Aug 7, 2012
Decision date	May 6, 2013
Days to decision	272 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Repro-Med Systems, Inc. Dba Rms Medical Products**
Location	New York, NY, US
Contact	ANDREW I SEALFON
510(k) history	3 submissions · 3 cleared · 2013-2019