FDA 510(k) Premarket Notification

K092905 AUTOSTART BURETTE

CLEARED

Mar 4, 2010
164 days to decision

K092905 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k092905/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Sep 21, 2009
Decision date	Mar 4, 2010
Days to decision	164 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Analytica Pty, Ltd.**
Location	Stillwater, MN, US
Contact	ELAINE DUNCAN
510(k) history	1 submissions · 1 cleared · 2010-2010