

K092905 AUTOSTART BURETTEMar 4, 2010
164 days to decisionK092905 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k092905/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 21, 2026