FDA 510(k) Premarket Notification

K122690 ULTRAQUIK PERIPHERAL NERVE BLOCK NEEDLE

CLEARED

Dec 21, 2012
108 days to decision

K122690 · Product code: **CAZ** · Anesthesiology
Source: https://www.510kdatabase.net/k122690/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Anesthesia Conduction Kit (CAZ)
Date received	Sep 4, 2012
Decision date	Dec 21, 2012
Days to decision	108 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Arrow International, Inc.**
Location	Reading, PA, US
Contact	PAUL AMUDALA
510(k) history	19 submissions · 17 cleared · 2003-2020

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k122690/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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