FDA 510(k) Premarket Notification

K092306 SECURACATH UNIVERSAL

CLEARED

Jul 9, 2010
345 days to decision

K092306 · Product code: **LJS** · General Hospital
Source: https://www.510kdatabase.net/k092306/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Long-term Greater Than 30 Days (LJS)
Date received	Jul 29, 2009
Decision date	Jul 9, 2010
Days to decision	345 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Interrad Medical, Inc.**
Location	Plymouth, MN, US
Contact	JONATHAN KAHAN
510(k) history	7 submissions · 7 cleared · 2008-2021

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k092306/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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