FDA 510(k) Premarket Notification

K213855 Arrow Pressure Injectable Midline Catheter

CLEARED

Sep 2, 2022
266 days to decision

K213855 · Product code: **PND** · General Hospital
Source: https://www.510kdatabase.net/k213855/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Midline Catheter (PND)
Date received	Dec 10, 2021
Decision date	Sep 2, 2022
Days to decision	266 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Arrow International, LLC Subsidiary of Teleflex Incorporated**
Location	Morrisville, NC, US
Contact	Kim Pennington
510(k) history	1 submissions · 1 cleared · 2022-2022