

**K213855 Arrow Pressure Injectable Midline Catheter**Sep 2, 2022  
266 days to decisionK213855 · Product code: **PND** · General Hospital  
Source: <https://www.510kdatabase.net/k213855/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k213855/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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