

**K093489 IN VISION-PLUS NANO WITH NEUTRAL ADVANTAGE TECHNOLOGY**Jul 29, 2010  
261 days to decisionK093489 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k093489/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Nov 10, 2009
Decision date	Jul 29, 2010
Days to decision	261 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Rymed Technologies, Inc.</b>
Location	Shelton, CT, US
Contact	JOSEPH AZARY
510(k) history	3 submissions · 3 cleared · 1996-2010

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k093489/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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