

**K120830 NAVIGATOR DELIVERY SYSTEM (NAVIGATOR DS)**Apr 16, 2012  
28 days to decisionK120830 · Product code: **FRN** · General Hospital  
Source: <https://www.510kdatabase.net/k120830/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Pump, Infusion (FRN)
Date received	Mar 19, 2012
Decision date	Apr 16, 2012
Days to decision	28 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Carticept Medical, Inc.</b>
Location	Alpharetta, GA, US
Contact	Tanya Eberle
510(k) history	6 submissions · 6 cleared · 2011-2015

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k120830/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 3, 2026