

**K041791 5.2F DUAL LUMEN EXTENDED LENGTH CATHETER  
(DELC)**Sep 10, 2004  
70 days to decisionK041791 · Product code: **NQJ** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k041791/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Hemodialysis, Non-implanted, Ultrafiltration, For Peripheral Use (NQJ)
Date received	Jul 2, 2004
Decision date	Sep 10, 2004
Days to decision	70 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Chf Solutions, Inc.</b>
Location	Washington, DC, US
Contact	AMY PETERSON
510(k) history	13 submissions · 13 cleared · 2002-2020

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k041791/>, Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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