

**K031689 DUAL LUMEN EXTENDED LENGTH CATHETER
(DELIC)**Dec 23, 2003
204 days to decisionK031689 · Product code: **NQJ** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k031689/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Hemodialysis, Non-implanted, Ultrafiltration, For Peripheral Use (NQJ)
Date received	Jun 2, 2003
Decision date	Dec 23, 2003
Days to decision	204 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k031689/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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