

**K020430 DIATEK CONNECTOR ASSEMBLY REPLACEMENT KIT
(CARK)**May 9, 2002
90 days to decisionK020430 · Product code: **NFK** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k020430/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Kit, Repair, Catheter, Hemodialysis (NFK)
Date received	Feb 8, 2002
Decision date	May 9, 2002
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Diatek, Inc.
Location	Mchenry, IL, US
Contact	JON WILSON
510(k) history	15 submissions · 15 cleared · 1979-2002

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k020430/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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