

K780079 CANNULA, ARTERY, RENALFeb 13, 1978
28 days to decisionK780079 · Product code: **DQR** · CardiovascularSource: <https://www.510kdatabase.net/k780079/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Catheter (DQR)
Date received	Jan 16, 1978
Decision date	Feb 13, 1978
Days to decision	28 days
Third-party review	No

APPLICANT

Company	Dravon Medical, Inc.
Location	Mchenry, IL, US
510(k) history	16 submissions · 16 cleared · 1977-1987

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Device record: <https://www.510kdatabase.net/k780079/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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