FDA 510(k) Premarket Notification

K780079 CANNULA, ARTERY, RENAL

CLEARED

Feb 13, 1978
28 days to decision

K780079 · Product code: **DQR** · Cardiovascular
Source: 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