

K800622 EXTENSION LINEApr 29, 1980
42 days to decisionK800622 · Product code: **DQO** · Cardiovascular
Source: <https://www.510kdatabase.net/k800622/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Mar 18, 1980
Decision date	Apr 29, 1980
Days to decision	42 days
Third-party review	No

APPLICANT

Company	Norton Performance Plastics Corp.
Location	Mchenry, IL, US
510(k) history	39 submissions · 39 cleared · 1977-1985

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Device record: <https://www.510kdatabase.net/k800622/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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