

K791915 WHITE SEALING CAPDec 19, 1979
85 days to decisionK791915 · Product code: **DQO** · CardiovascularSource: <https://www.510kdatabase.net/k791915/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Sep 25, 1979
Decision date	Dec 19, 1979
Days to decision	85 days
Third-party review	No

APPLICANT

Company	Quinton, Inc.
Location	Mchenry, IL, US
510(k) history	164 submissions · 160 cleared · 1976-2003

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Device record: <https://www.510kdatabase.net/k791915/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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