

K830022 S.M.P. CONTRAST MEDIA SETJan 21, 1983
17 days to decisionK830022 · Product code: **DQO** · CardiovascularSource: <https://www.510kdatabase.net/k830022/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Jan 4, 1983
Decision date	Jan 21, 1983
Days to decision	17 days
Third-party review	No

APPLICANT

Company	Omega Technologies, Inc.
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1983-1983

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Device record: <https://www.510kdatabase.net/k830022/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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