

K810508 SONTEK ANGIOGRAPHIC CATHETERApr 17, 1981
52 days to decisionK810508 · Product code: **DQO** · CardiovascularSource: <https://www.510kdatabase.net/k810508/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Feb 24, 1981
Decision date	Apr 17, 1981
Days to decision	52 days
Third-party review	No

APPLICANT

Company	Sontek Corp.
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1981-1983

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Device record: <https://www.510kdatabase.net/k810508/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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