

K790678 INTRAUTERINE CATHETER TIPMay 11, 1979
32 days to decisionK790678 · Product code: **KXO** · Radiology
Source: <https://www.510kdatabase.net/k790678/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Pressure, Intrauterine (KXO)
Date received	Apr 9, 1979
Decision date	May 11, 1979
Days to decision	32 days
Third-party review	No

APPLICANT

Company	Sonicaid, Inc.
Location	Mchenry, IL, US
510(k) history	18 submissions · 17 cleared · 1977-1988

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Device record: <https://www.510kdatabase.net/k790678/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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