FDA 510(k) Premarket Notification

K790678 INTRAUTERINE CATHETER TIP

CLEARED

May 11, 1979
32 days to decision

K790678 · 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