

K840339 UMBILICAL VESSEL CATHETERJul 25, 1984
182 days to decisionK840339 · Product code: **FOS** · General Hospital
Source: <https://www.510kdatabase.net/k840339/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Umbilical Artery (FOS)
Date received	Jan 25, 1984
Decision date	Jul 25, 1984
Days to decision	182 days
Third-party review	No

APPLICANT

Company	Catheter Technology Corp.
Location	Walker, MI, US
510(k) history	13 submissions · 13 cleared · 1983-1988

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Device record: <https://www.510kdatabase.net/k840339/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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