FDA 510(k) Premarket Notification

K840339 UMBILICAL VESSEL CATHETER

CLEARED

Jul 25, 1984
182 days to decision

K840339 · 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