FDA 510(k) Premarket Notification

K880887 NEOCATH

CLEARED

May 24, 1988
83 days to decision

K880887 · Product code: **DWF** · Cardiovascular
Source: https://www.510kdatabase.net/k880887/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Mar 2, 1988
Decision date	May 24, 1988
Days to decision	83 days
Third-party review	No

APPLICANT

Company	**Taris Corp.**
Location	Fernandina Beach, FL, US
Contact	WILLIAM J SCHRODER
510(k) history	3 submissions · 3 cleared · 1988-1988