FDA 510(k) Premarket Notification

K831330 UNIVERSAL CORONARY CANNULA

CLEARED

Aug 10, 1983
107 days to decision

K831330 · Product code: **DWF** · Cardiovascular
Source: https://www.510kdatabase.net/k831330/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Apr 25, 1983
Decision date	Aug 10, 1983
Days to decision	107 days
Third-party review	No

APPLICANT

Company	**Possis Medical, Inc.**
Location	Walker, MI, US
510(k) history	34 submissions · 34 cleared · 1983-2014

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k831330/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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