FDA 510(k) Premarket Notification

K831759 INTRACARDIA SUCKERS

CLEARED

Aug 12, 1983
72 days to decision

K831759 · Product code: **DWF** · Cardiovascular
Source: https://www.510kdatabase.net/k831759/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Jun 1, 1983
Decision date	Aug 12, 1983
Days to decision	72 days
Third-party review	No

APPLICANT

Company	**Research Industries Corp.**
Location	Walker, MI, US
510(k) history	5 submissions · 5 cleared · 1983-1995