FDA 510(k) Premarket Notification

K883696 FLO-RESTER

CLEARED

Nov 30, 1988
92 days to decision

K883696 · Product code: **DXC** · Cardiovascular
Source: https://www.510kdatabase.net/k883696/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Clamp, Vascular (DXC)
Date received	Aug 30, 1988
Decision date	Nov 30, 1988
Days to decision	92 days
Third-party review	No

APPLICANT

Company	**Bio-Vascular, Inc.**
Location	St. Paul, MN, US
Contact	SHERRY A KRATTLEY
510(k) history	26 submissions · 25 cleared · 1986-2000