FDA 510(k) Premarket Notification

K011208 BIOVER

CLEARED

Oct 2, 2002
531 days to decision

K011208 · Product code: **DXC** · Cardiovascular
Source: https://www.510kdatabase.net/k011208/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Clamp, Vascular (DXC)
Date received	Apr 19, 2001
Decision date	Oct 2, 2002
Days to decision	531 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Biover AG**
Location	Ridley Park, PA, US
Contact	HEINZ E WICK
510(k) history	1 submissions · 1 cleared · 2002-2002