

K011208 BIOVEROct 2, 2002
531 days to decisionK011208 · 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

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