

K070323 TEMPORARY LIMB SALVAGE SHUNTFeb 15, 2007
13 days to decisionK070323 · Product code: **DXC** · Cardiovascular
Source: <https://www.510kdatabase.net/k070323/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Clamp, Vascular (DXC)
Date received	Feb 2, 2007
Decision date	Feb 15, 2007
Days to decision	13 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Vascutek, Ltd.
Location	Pa49rr Scotland, GB
Contact	KAREN KELSO
Website	http://www.vascutek.com/
510(k) history	26 submissions · 23 cleared · 1991-2025

Vascutek, Ltd. is a global medical device company committed to developing innovative solutions for aortic and peripheral vascular disease. The company operates with a manufacturing facility in Inchinnan, Renfrewshire, Scotland. Now part of Terumo Aortic, the brand continues to advance cardiovascular care worldwide. Vascutek has a strong FDA 510(k) regulatory track record. The company has received FDA 510(k) clearances from total submissions since its first clearance in 1991. All submissions focus on Cardiovascular devices, reflecting the company's specialization in aortic...
