FDA 510(k) Premarket Notification

K030999 EPTFE VASCULAR PROSTHESIS

CLEARED

Apr 9, 2003
9 days to decision

K030999 · Product code: **DSY** · Cardiovascular
Source: https://www.510kdatabase.net/k030999/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Prosthesis, Vascular Graft, Of 6mm And Greater Diameter (DSY)
Date received	Mar 31, 2003
Decision date	Apr 9, 2003
Days to decision	9 days
Third-party review	No
Summary / Statement	Statement
Other names	SEALPTFE

APPLICANT

Company	**Vascutek, Ltd.**
Location	Pa49rr Scotland, GB
Contact	STEVEN ARICK
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...