FDA 510(k) Premarket Notification

K022242 INTERVASCULAR INTERGARD KNITTED VASCULAR GRAFT SIZER

CLEARED

Aug 14, 2002
34 days to decision

K022242 · Product code: **DSY** · Cardiovascular
Source: https://www.510kdatabase.net/k022242/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Vascular Graft, Of 6mm And Greater Diameter (DSY)
Date received	Jul 11, 2002
Decision date	Aug 14, 2002
Days to decision	34 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Intervascular**
Location	Wellesley, MA, US
Contact	Amy Aulwes
510(k) history	3 submissions · 3 cleared · 2002-2005

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k022242/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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