

**K890182 PROTEGRAFT DV 1500, 1900 & LOW POR, UNIGRAFT
DV**Feb 28, 1990
406 days to decisionK890182 · Product code: **DSY** · Cardiovascular
Source: <https://www.510kdatabase.net/k890182/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Vascular Graft, Of 6mm And Greater Diameter (DSY)
Date received	Jan 18, 1989
Decision date	Feb 28, 1990
Days to decision	406 days
Third-party review	No

APPLICANT

Company	Aesculap Instruments Corp.
Location	Mchenry, IL, US
Contact	DEAN A KNIGHT
510(k) history	13 submissions · 13 cleared · 1977-1990

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k890182/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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