

K864919 SCANLAN (TM) VEIN/GRAFT HOLDERJan 12, 1987
27 days to decisionK864919 · Product code: **KPK** · CardiovascularSource: <https://www.510kdatabase.net/k864919/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Marker, Ostia, Aorto-saphenous Vein (KPK)
Date received	Dec 16, 1986
Decision date	Jan 12, 1987
Days to decision	27 days
Third-party review	No

APPLICANT

Company	Scanlan Intl., Inc.
Location	Mchenry, IL, US
Contact	KENNETH R BLAKE
510(k) history	19 submissions · 19 cleared · 1978-1996

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