FDA 510(k) Premarket Notification

K823764 SCANLAN SINGLE-USE AORTA PUNCH

CLEARED

Mar 24, 1983
100 days to decision

K823764 · Product code: **DWS** · Cardiovascular
Source: https://www.510kdatabase.net/k823764/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instruments, Surgical, Cardiovascular (DWS)
Date received	Dec 14, 1982
Decision date	Mar 24, 1983
Days to decision	100 days
Third-party review	No

APPLICANT

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