FDA 510(k) Premarket Notification

K841521 SCANLAN ENDARSECTOR

CLEARED

Aug 3, 1984
113 days to decision

K841521 · Product code: **DWX** · Cardiovascular
Source: https://www.510kdatabase.net/k841521/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stripper, Artery, Intraluminal (DWX)
Date received	Apr 12, 1984
Decision date	Aug 3, 1984
Days to decision	113 days
Third-party review	No

APPLICANT

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