

K841521 SCANLAN ENDARSECTORAug 3, 1984
113 days to decisionK841521 · Product code: **DWX** · CardiovascularSource: <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

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Device record: <https://www.510kdatabase.net/k841521/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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