

**K823764 SCANLAN SINGLE-USE AORTA PUNCH**Mar 24, 1983  
100 days to decisionK823764 · Product code: **DWS** · CardiovascularSource: <https://www.510kdatabase.net/k823764/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k823764/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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