

**K812932 ANERIOD GAUZE**Dec 2, 1981  
43 days to decisionK812932 · Product code: **DXG** · CardiovascularSource: <https://www.510kdatabase.net/k812932/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Computer, Diagnostic, Pre-programmed, Single-function (DXG)
Date received	Oct 20, 1981
Decision date	Dec 2, 1981
Days to decision	43 days
Third-party review	No

**APPLICANT**

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Company	<b>Abco Dealers, Inc.</b>
Location	Mchenry, IL, US
510(k) history	127 submissions · 127 cleared · 1976-1991

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

Device record: <https://www.510kdatabase.net/k812932/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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