

**K831401 SINGLE NEEDLE SUBCLAVIAN VEIN ACCESS**Jun 3, 1983  
32 days to decisionK831401 · Product code: **FIQ** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k831401/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, A-v Shunt (FIQ)
Date received	May 2, 1983
Decision date	Jun 3, 1983
Days to decision	32 days
Third-party review	No

**APPLICANT**

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Company	<b>Med-West, Inc.</b>
Location	Mchenry, IL, US
510(k) history	11 submissions · 8 cleared · 1983-1988

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

Device record: <https://www.510kdatabase.net/k831401/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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