

**K760868 VEST RESTRAINT**Nov 1, 1976  
14 days to decisionK760868 · Product code: **KNW** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k760868/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Biopsy (KNW)
Date received	Oct 18, 1976
Decision date	Nov 1, 1976
Days to decision	14 days
Third-party review	No

**APPLICANT**

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Company	<b>Orthopedic Equipment Co., Inc.</b>
Location	Mchenry, IL, US
510(k) history	45 submissions · 45 cleared · 1976-1984

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

Device record: <https://www.510kdatabase.net/k760868/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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