

**K981721 C0-AXIAL INTRODUCER NEEDLE**Jul 28, 1998  
74 days to decisionK981721 · Product code: **KNW** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k981721/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Biopsy (KNW)
Date received	May 15, 1998
Decision date	Jul 28, 1998
Days to decision	74 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Inrad</b>
Location	Kentwood, MI, US
Contact	ANNE ARMSTRONG
510(k) history	11 submissions · 11 cleared · 1998-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k981721/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 22, 2026