

**K981166 ACCUCORE CORE BIOPSY NEEDLE CATALOG
 CODES:581014, 581614, 581618, 582018, 582518, 581620,
 582020, 581214**

Jun 22, 1998
 83 days to decision

K981166 · Product code: **KNW** · Gastroenterology & Urology
 Source: <https://www.510kdatabase.net/k981166/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Biopsy (KNW)
Date received	Mar 31, 1998
Decision date	Jun 22, 1998
Days to decision	83 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k981166/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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