FDA 510(k) Premarket Notification

K981721 C0-AXIAL INTRODUCER NEEDLE

CLEARED

Jul 28, 1998
74 days to decision

K981721 · Product code: **KNW** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k981721/

SUBMISSION DETAILS

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

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