FDA 510(k) Premarket Notification

CLEARED

K093256 SELECTCORE VARIABLE THROW BIOPSY DEVICE MODEL 991014, 991514, 991018, 991518, 992018

Nov 17, 2009
29 days to decision

K093256 · Product code: **KNW** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k093256/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Biopsy (KNW)
Date received	Oct 19, 2009
Decision date	Nov 17, 2009
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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