FDA 510(k) Premarket Notification

K130282 REMINGTON MEDICAL CENTREFIRE 22 BIOPSY INSTRUMENT

CLEARED

May 13, 2013
97 days to decision

K130282 · Product code: **KNW** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k130282/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Biopsy (KNW)
Date received	Feb 5, 2013
Decision date	May 13, 2013
Days to decision	97 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	**Remington Medical, Inc.**
Location	Great Neck, NY, US
Contact	CAITLIN SENTER, MS, RAC
510(k) history	19 submissions · 19 cleared · 1993-2025