

K040266 CORE NEEDLE BIOPSYJul 9, 2004
155 days to decisionK040266 · Product code: **KNW** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k040266/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Biopsy (KNW)
Date received	Feb 5, 2004
Decision date	Jul 9, 2004
Days to decision	155 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Remington Medical, Inc.
Location	Great Neck, NY, US
Contact	MARIA P GRANITO
510(k) history	19 submissions · 19 cleared · 1993-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k040266/> Data sourced from FDA 510(k) public records (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. - Generated May 25, 2026