FDA 510(k) Premarket Notification

K191315 Remington Medical, Inc. Automatic Cutting Needle (NAC)

CLEARED

Jul 12, 2019
58 days to decision

K191315 · Product code: **KNW** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k191315/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Biopsy (KNW)
Date received	May 15, 2019
Decision date	Jul 12, 2019
Days to decision	58 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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