

K251904 VascuChek Kit, VascuChek Transceiver, VascuChek Clinical Probe, VascuChek Speaker Dock (VC-KIT-02, VC-TRX-02, VC-CP-01, VC-SD-01)

Nov 20, 2025
153 days to decision

K251904 · Product code: **DPW** · Cardiovascular
Source: <https://www.510kdatabase.net/k251904/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Flowmeter, Blood, Cardiovascular (DPW)
Date received	Jun 20, 2025
Decision date	Nov 20, 2025
Days to decision	153 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary
Other names	VascuChek Kit, VascuChek Transceiver, VascuChek Surgical Probe, VascuChek Speaker Dock (VC-KIT-02, VC-TRX-02, VC-SP-01, VC-SD-01)

APPLICANT

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

REGULATORY CONSULTANT

Consulting firm	Secure BioMed Evaluations
Contact	Justin Gracyalny

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)

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Device record: <https://www.510kdatabase.net/k251904/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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