

K233326 Merge Hemo, Model RCSV2Jun 21, 2024
266 days to decisionK233326 · Product code: **DQK** · CardiovascularSource: <https://www.510kdatabase.net/k233326/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Computer, Diagnostic, Programmable (DQK)
Date received	Sep 29, 2023
Decision date	Jun 21, 2024
Days to decision	266 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Merge Healthcare Incorporated
Location	Hartland, WI, US
Contact	Carol Nakagawa
510(k) history	8 submissions · 8 cleared · 2016-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k233326/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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