

K211585 Bodyport Cardiac ScaleJul 29, 2022
431 days to decisionK211585 · Product code: **DSB** · Cardiovascular
Source: <https://www.510kdatabase.net/k211585/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plethysmograph, Impedance (DSB)
Date received	May 24, 2021
Decision date	Jul 29, 2022
Days to decision	431 days
Third-party review	Yes
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Bodyport, Inc.
Location	San Francisco, CA, US
Contact	Mara Korsunsky
510(k) history	1 submissions · 1 cleared · 2022-2022

REGULATORY CONSULTANT

Consulting firm	Regulatory Technology Services, LLC
Contact	Prithul Bom

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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