

K202483 MyHomeDocMar 23, 2021
207 days to decisionK202483 · Product code: **DQD** · Cardiovascular
Source: <https://www.510kdatabase.net/k202483/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stethoscope, Electronic (DQD)
Date received	Aug 28, 2020
Decision date	Mar 23, 2021
Days to decision	207 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Myhomedoc , Ltd.
Location	Ra'Anana, IL
Contact	Orly Maor
510(k) history	1 submissions · 1 cleared · 2021-2021

REGULATORY CONSULTANT

Consulting firm	Hogan Lovells US LLP
Contact	Jonathan Kahan

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

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