

K170874 Eko Model E5 System (EME5), Eko DUOMay 19, 2017
56 days to decisionK170874 · Product code: **DQD** · CardiovascularSource: <https://www.510kdatabase.net/k170874/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stethoscope, Electronic (DQD)
Date received	Mar 24, 2017
Decision date	May 19, 2017
Days to decision	56 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Eko Devices, Inc.
Location	Berkeley, CA, US
Contact	Connor Landgraf
510(k) history	6 submissions · 6 cleared · 2015-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k170874/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 27, 2026