

K221616 Patient Monitor, models LMPLUS-12, LMPLUS-15 and LMPLUS-17Aug 1, 2022
59 days to decisionK221616 · Product code: **MHX** · Cardiovascular
Source: <https://www.510kdatabase.net/k221616/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	Jun 3, 2022
Decision date	Aug 1, 2022
Days to decision	59 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Caf Medical Solutions, Inc.
Location	Hockley, TX, US
Contact	Oscar Arriaga Yamin
510(k) history	3 submissions · 3 cleared · 2022-2022

REGULATORY CONSULTANT

Consulting firm	Compliance4Devices
Contact	Juan Tezak

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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