FDA 510(k) Premarket Notification

K222416 RIA Safeguard

CLEARED

May 1, 2023
264 days to decision

K222416 · Product code: **FRF** · General Hospital
Source: https://www.510kdatabase.net/k222416/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cleaner, Air, Medical Recirculating (FRF)
Date received	Aug 10, 2022
Decision date	May 1, 2023
Days to decision	264 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Ria Tech Co., Ltd.**
Location	Taipei City, TW
Contact	David Chao
510(k) history	1 submissions · 1 cleared · 2023-2023

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k222416/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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