FDA 510(k) Premarket Notification

K241140 ViroZap Indoor Air Purifier, In Duct Model 1008

CLEARED

Aug 1, 2024
99 days to decision

K241140 · Product code: **FRA** · General Hospital
Source: https://www.510kdatabase.net/k241140/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Purifier, Air, Ultraviolet, Medical (FRA)
Date received	Apr 24, 2024
Decision date	Aug 1, 2024
Days to decision	99 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Applied Photonix, LLC**
Location	Tampa, FL, US
Contact	D. Yogi Goswami
510(k) history	1 submissions · 1 cleared · 2024-2024

REGULATORY CONSULTANT

Consulting firm	**King & Spalding Llp**
Contact	Jeffrey Shapiro

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