

K241140 ViroZap Indoor Air Purifier, In Duct Model 1008Aug 1, 2024
99 days to decisionK241140 · 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](https://accessdata.fda.gov)510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k241140/> 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 13, 2026