

K252471 Paptizer 360Apr 29, 2026
266 days to decisionK252471 · Product code: **QXQ** · General Hospital
Source: <https://www.510kdatabase.net/k252471/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Respiratory Accessory Microbial Reduction Device. (QXQ)
Date received	Aug 6, 2025
Decision date	Apr 29, 2026
Days to decision	266 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	LiViliti Health Products Corporation
Location	Lake City, FL, US
Contact	Brian Sharpe
Website	https://www.liviliti.com
510(k) history	1 submissions · 1 cleared · 2026-2026

LiViliti Health Products Corporation develops General Hospital devices with a manufacturing facility in Lake City, US. The company specializes in smart sanitization technology for healthcare environments. LiViliti has received FDA 510(k) clearance from total submission. The company's regulatory activity spans 2026, with its cleared device focused on General Hospital applications. Recent clearance demonstrates active engagement with FDA oversight. The Paptizer 360 represents the company's core offering in hospital-grade sanitization. The device features UVC LED technology ...

REGULATORY CONSULTANT

Consulting firm	Nilo Medical Consulting Group
Contact	Michael Nilo

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