

K121223 QLAB WITH FHN AND VPQ PLUG-INMay 15, 2012
22 days to decisionK121223 · Product code: **LLZ** · Radiology
Source: <https://www.510kdatabase.net/k121223/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Apr 23, 2012
Decision date	May 15, 2012
Days to decision	22 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	Philips Ultrasound, Inc.
Location	Santa Ana, CA, US
Contact	PENNY GRECO
510(k) history	46 submissions · 46 cleared · 1985-2021

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k121223/> 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 26, 2026