FDA 510(k) Premarket Notification

K121223 QLAB WITH FHN AND VPQ PLUG-IN

CLEARED

May 15, 2012
22 days to decision

K121223 · 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