FDA 510(k) Premarket Notification

K070792 QLAB QUANTIFICATION

CLEARED

Apr 6, 2007
15 days to decision

K070792 · Product code: **LLZ** · Radiology
Source: https://www.510kdatabase.net/k070792/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Mar 22, 2007
Decision date	Apr 6, 2007
Days to decision	15 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

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