

K172059 LinkQuest Diagnostic Ultrasound System model SQ860

Oct 5, 2017
90 days to decision

K172059 · Product code: **IYN** · Radiology
Source: <https://www.510kdatabase.net/k172059/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Doppler, Ultrasonic (IYN)
Date received	Jul 7, 2017
Decision date	Oct 5, 2017
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Linkquest, Inc.
Location	San Diego, CA, US
Contact	Xiaolong Yu
510(k) history	1 submissions · 1 cleared · 2017-2017

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k172059/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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