FDA 510(k) Premarket Notification

K132690 UP600 DIAGNOSTIC DOPPLER ULTRASOUND SYSTEM

CLEARED

Feb 25, 2014
181 days to decision

K132690 · Product code: **IYN** · Radiology
Source: https://www.510kdatabase.net/k132690/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Doppler, Ultrasonic (IYN)
Date received	Aug 28, 2013
Decision date	Feb 25, 2014
Days to decision	181 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Qisda Corporation**
Location	Dublin, CA, US
Contact	BOB LEIKER
510(k) history	4 submissions · 4 cleared · 2014-2019

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k132690/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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