FDA 510(k) Premarket Notification

CLEARED

K173265 UltraVision 2 Diagnostic Ultrasound System

Nov 7, 2017
27 days to decision

K173265 · Product code: **IYN** · Radiology
Source: https://www.510kdatabase.net/k173265/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	System, Imaging, Pulsed Doppler, Ultrasonic (IYN)
Date received	Oct 11, 2017
Decision date	Nov 7, 2017
Days to decision	27 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	**Winprobe Corporation**
Location	Palm Beach Gradens, FL, US
Contact	Guy Scott
510(k) history	2 submissions · 2 cleared · 2015-2017