K002470 M2424A SONOS 5500/4500 DIAGNOSTIC ULTRASOUND SYSTEM, VERSION B.2. 21330A TRANSDUCER

CLEARED

Sep 8, 2000
28 days to decision

K002470 · Product code: **IYN** · Radiology
Source: https://www.510kdatabase.net/k002470/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Doppler, Ultrasonic (IYN)
Date received	Aug 11, 2000
Decision date	Sep 8, 2000
Days to decision	28 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	**Agilent Technologies, Inc.**
Location	Pittsburgh, PA, US
Contact	ANN F MICHAELS
Website	http://www.agilent.com
510(k) history	30 submissions · 30 cleared · 1985-2017

Agilent Technologies, Inc. is an American global company that provides instruments, software, services, and consumables for laboratories. Headquartered in Santa Clara, California, Agilent was established in 1999 as a spin-off from Hewlett-Packard. The company has received FDA 510(k) clearances from total submissions. Cardiovascular devices represent the dominant focus, accounting for approximately 80% of regulatory submissions. Agilent's FDA 510(k) clearance history spans from 1985 to 2017, establishing a long track record in medical device regulation. Notable cleared dev...