FDA 510(k) Premarket Notification

K260673 LOGIQ Vita

CLEARED

Mar 24, 2026
22 days to decision

K260673 · Product code: **IYN** · Radiology
Source: https://www.510kdatabase.net/k260673/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Doppler, Ultrasonic (IYN)
Date received	Mar 2, 2026
Decision date	Mar 24, 2026
Days to decision	22 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary
Other names	LOGIQ Vita Pro; LOGIQ Vita Express; LOGIQ Vita Plus; LOGIQ Vita Power; LOGIQ S20; LOGIQ S20 Pro; LOGIQ S20 Express; LOGIQ S20 Plus; LOGIQ S20 Power

APPLICANT

Company	**GE Medical Systems Ultrasound and Primary Care Diagnostics**
Location	Wauwatosa, WI, US
Contact	Lee Bush
510(k) history	64 submissions · 64 cleared · 2015-2026