

**K220882 Vivid E80, Vivid E90, Vivid E95**Jul 22, 2022  
119 days to decisionK220882 · Product code: **IYN** · Radiology  
Source: <https://www.510kdatabase.net/k220882/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Doppler, Ultrasonic (IYN)
Date received	Mar 25, 2022
Decision date	Jul 22, 2022
Days to decision	119 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Ge Medical Systems Ultrasound And</b>
Location	Wauwatosa, WI, US
Contact	Tracey Ortiz
510(k) history	5 submissions · 5 cleared · 2020-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k220882/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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