

**K163120 EPIQ 5 Diagnostic Ultrasound System**Jan 10, 2017  
63 days to decisionK163120 · Product code: **IYN** · Radiology  
Source: <https://www.510kdatabase.net/k163120/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Doppler, Ultrasonic (IYN)
Date received	Nov 8, 2016
Decision date	Jan 10, 2017
Days to decision	63 days
Third-party review	No
Summary / Statement	Summary
Other names	EPIQ 7 Diagnostic Ultrasound System; Affiniti 50 Diagnostic Ultrasound System; Affiniti 70 Diagnostic Ultrasound System

**APPLICANT**

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Company	<b>Philips Ultrasound, Inc.</b>
Location	Santa Ana, CA, US
Contact	Peng Cui
510(k) history	46 submissions · 46 cleared · 1985-2021

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k163120/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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