

**K190476 FUJIFILM Sonosite Vevo MD Imaging System**Mar 15, 2019  
16 days to decisionK190476 · Product code: **IYN** · Radiology  
Source: <https://www.510kdatabase.net/k190476/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Doppler, Ultrasonic (IYN)
Date received	Feb 27, 2019
Decision date	Mar 15, 2019
Days to decision	16 days
Third-party review	Yes
Summary / Statement	Summary

**APPLICANT**

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Company	<b>FUJIFILM Sonosite, Inc.</b>
Location	Bothell, WA, US
Contact	Anoush Frankian
Website	<a href="https://www.sonosite.com">https://www.sonosite.com</a>
510(k) history	25 submissions · 25 cleared · 2013-2026

FUJIFILM Sonosite, Inc. is a portable ultrasound manufacturer based in Bothell, US. The company specializes in point-of-care ultrasound systems for clinical imaging. FUJIFILM Sonosite has received FDA 510(k) clearances from total submissions since 2013. The company's portfolio focuses exclusively on Radiology devices. The latest clearance was in 2026, demonstrating continued regulatory activity and product innovation. The company's cleared devices include portable ultrasound systems designed for diverse clinical settings. Products span multiple system lines, each configur...

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