FDA 510(k) Premarket Notification

K130173 SONOSITE MAXX SERIES ULTRASOUND SYSTEM

CLEARED

Mar 25, 2013
60 days to decision

K130173 · Product code: **IYN** · Radiology
Source: https://www.510kdatabase.net/k130173/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Doppler, Ultrasonic (IYN)
Date received	Jan 24, 2013
Decision date	Mar 25, 2013
Days to decision	60 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	**FUJIFILM Sonosite, Inc.**
Location	Bothell, WA, US
Contact	SCOTT PAULSON
Website	https://www.sonosite.com
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...