

**K173034 FibroScan® Family of Products (Models: 502 Touch, 530 Compact, and 430 Mini+)**Nov 14, 2017  
47 days to decisionK173034 · Product code: IYO · Radiology  
Source: <https://www.510kdatabase.net/k173034/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Sep 28, 2017
Decision date	Nov 14, 2017
Days to decision	47 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Echosens</b>
Location	Dedham, MA, US
Contact	Karine Bonenfant
510(k) history	11 submissions · 11 cleared · 2013-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k173034/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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