

# K221468 Citregen™ Tendon Interference Screw (TIS), Citrelock™ Tendon Fixation Device, Citrespline™ and Citrelock™ ACL Implants

Feb 14, 2023  
270 days to decision

K221468 · Product code: MAI · Orthopedic  
Source: <https://www.510kdatabase.net/k221468/>

## SUBMISSION DETAILS

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Fastener, Fixation, Biodegradable, Soft Tissue (MAI)
Date received	May 20, 2022
Decision date	Feb 14, 2023
Days to decision	270 days
Third-party review	No
Summary / Statement	Summary

## APPLICANT

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Company	<b>Acuitive Technologies, Inc.</b>
Location	Alendale, NJ, US
Contact	Matthew Poggie
510(k) history	6 submissions · 6 cleared · 2021-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k221468/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://accessdata.fda.gov)).

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