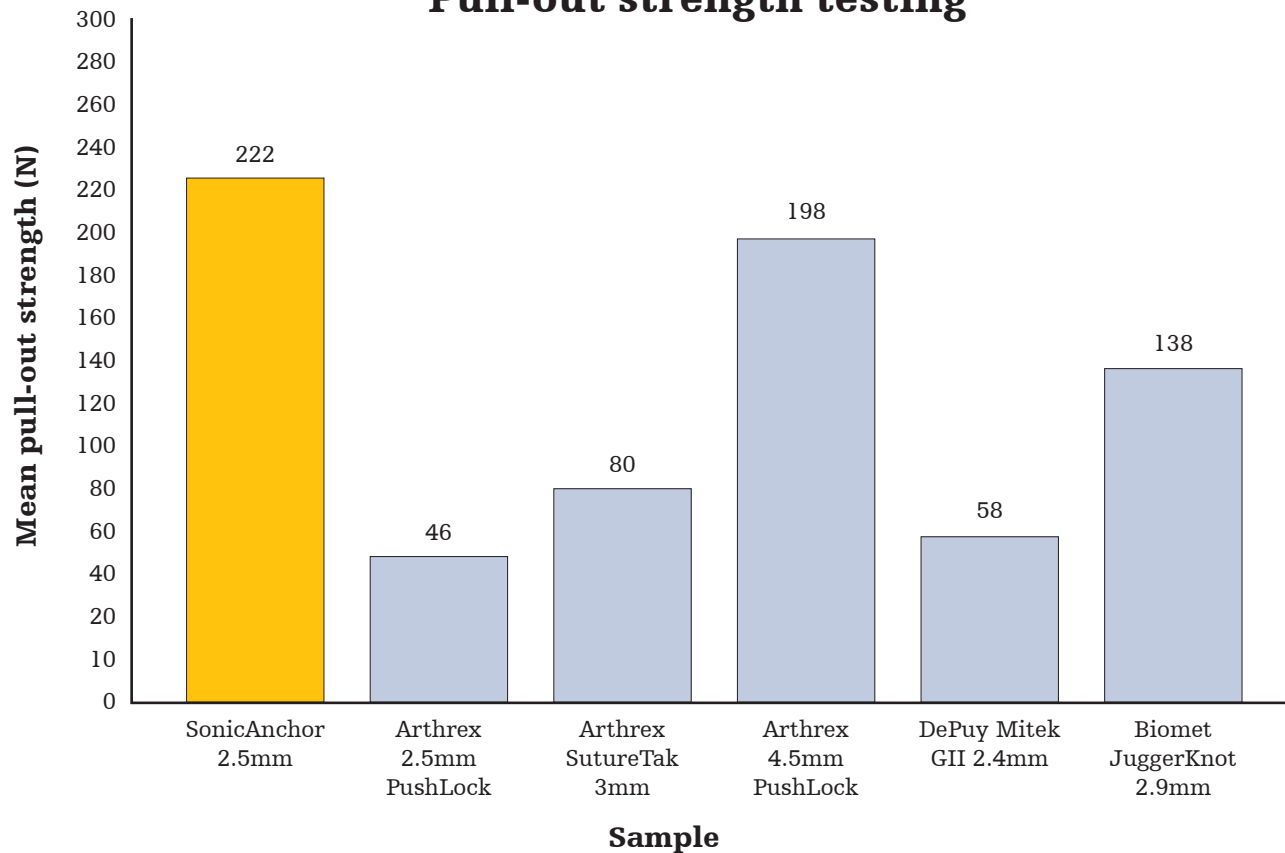


SonicAnchor™

Pull-out strength testing



Pull-out strength testing

According to 250815TW1¹, pull-out strength testing was performed within a water bath (filled with deionized water) at 37±1°C at a displacement rate of 60 mm/min. The test samples were conditioned to the temperature for 10 min before application of the tensile loads. During testing, the force-displacement curve over time was recorded and the test was stopped after implant failure. Based on this data, the pull-out strength (maximum force during testing) was determined for each sample.

1. Test report 140416TW1: SonicAnchor: internal document, Stryker, Trauma & Extremities, 2016.

This document is intended solely for the use of healthcare professionals. A surgeon must always rely on his or her own professional clinical judgment when deciding whether to use a particular product when treating a particular patient. Stryker does not dispense medical advice and recommends that surgeons be trained in the use of any particular product before using it in surgery.

The information presented is intended to demonstrate a Stryker product. A surgeon must always refer to the package insert, product label and/or instructions for use, including the instructions for Cleaning and Sterilization (if applicable), before using any Stryker product. Products may not be available in all markets because product availability is subject to the regulatory and/or medical practices in individual markets. Please contact your Stryker representative if you have questions about the availability of Stryker products in your area.

Stryker Corporation or its divisions or other corporate affiliated entities own, use or have applied for the following trademarks or service marks: SonicFusion, SonicAnchor, Stryker. All other trademarks are trademarks of their respective owners or holders.

Content ID: SON-EM-2, 01-2017

Copyright© 2017 Stryker