

Akilink[®]

MINI INVASIVE ACHILLES TENDON REPAIR SYSTEM



Single use system



90° handle to ease introduction and positioning



Multidirectional sutures targeting jig



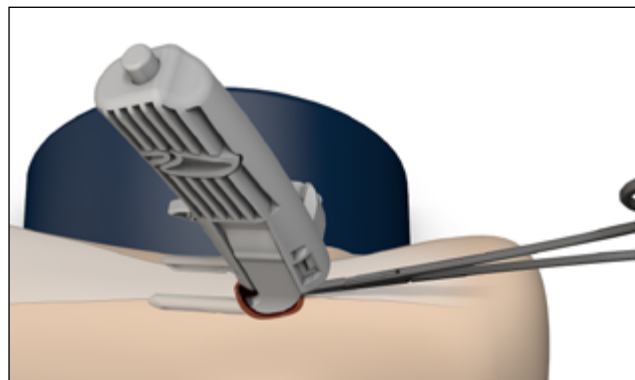
A GLOBAL EXTREMITY COMPANY

Akilink®

PRODUCT DESCRIPTION

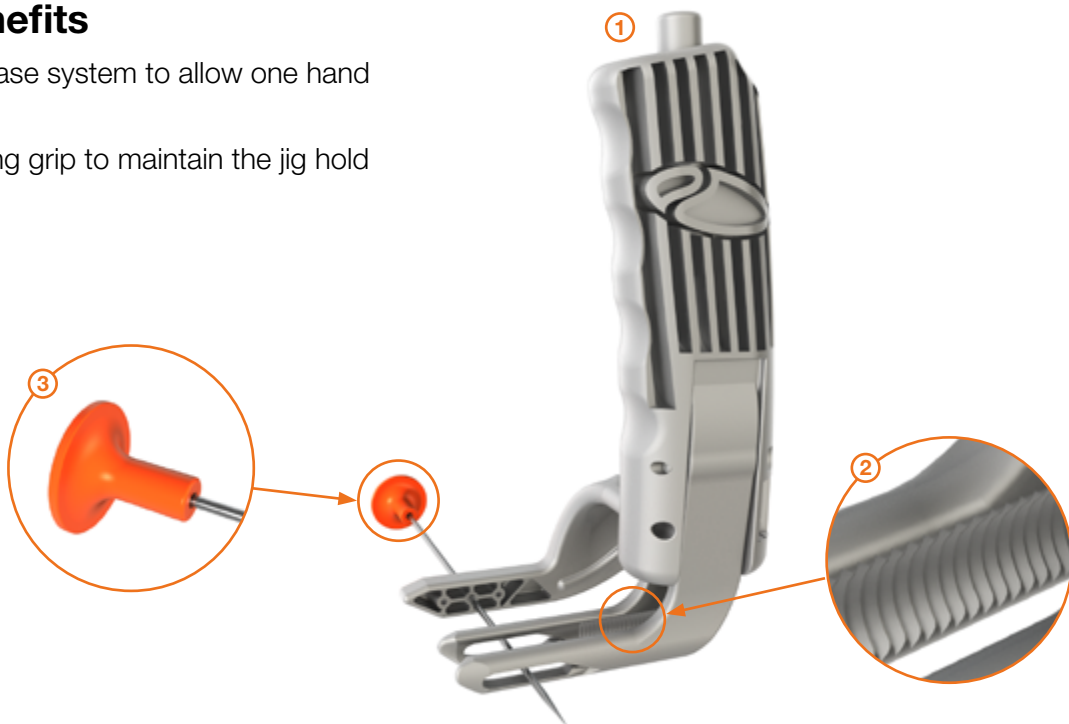
Easy introduction

- 90° handle « laryngoscope like », to ease introduction of the Akilink and to hold the Achilles tendon stump while sliding the Akilink inside the tendon sheath.



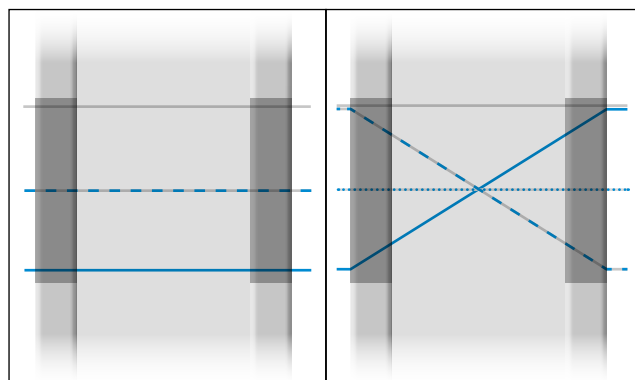
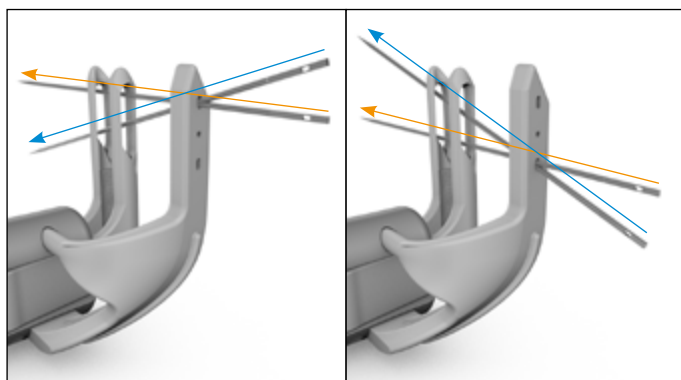
Instrument benefits

- Push/Slide & Release system to allow one hand only ①
- Inner leg anti-sliding grip to maintain the jig hold on the tendon ②
- Needle pusher ③



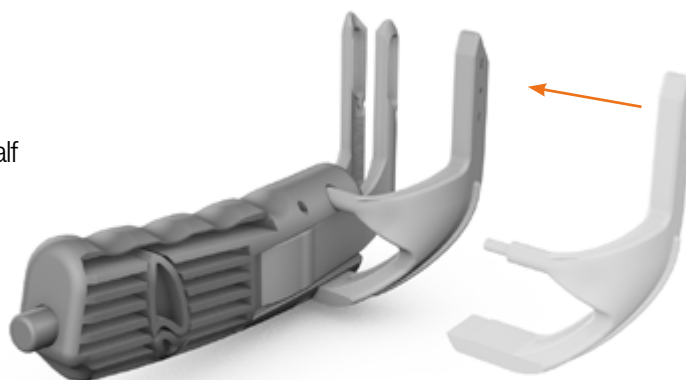
Sutures

- Double cross suturing
- Versatile sutures directions



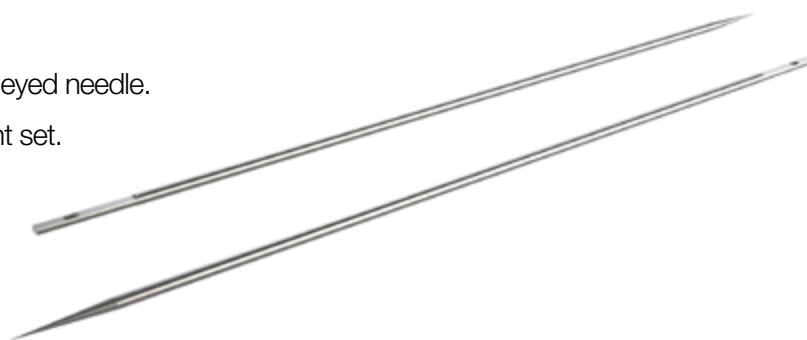
External guide

- Removable external guide for extra wide Calf



Needles

- 1.6 mm diameter and 150 mm length eyed needle.
- 2 needles in each single use instrument set.

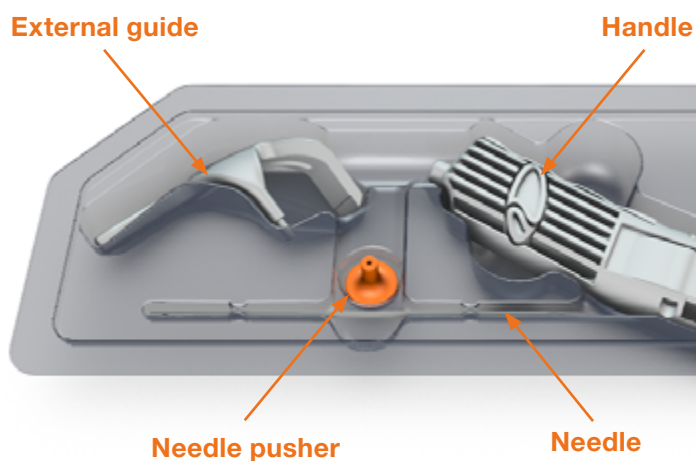


Akilink®

ORDERING INFORMATION

Single use instrument set / Always ready to use

G03 10001 Akilink® Mini Invasive Achilles Tendon Repair System



INDICATIONS

The Akilink® system is indicated for the mini invasive repair of acute (less than 10 days) closed ruptures of the Achilles tendon, located between 2 cm and 8 cm above the tuberosity of the calcaneum.

Note: 2 cm is the lowest limit, in this case, please choose an oblique suture on the calcaneus side.

CONTRA-INDICATIONS

The device should not be used in a patient who has currently, or who has history of:

- Acute or chronic, systemic inflammations,
- Active infections,
- Chronic rupture,
- Open rupture,
- Complex open rupture with skin defect,
- Rupture located between 0 and 2 cm above the tuberosity of the calcaneum or higher than 8 cm above the tuberosity of the calcaneum,
- Previous local surgery,
- Sensitivity/allergies to the medical device materials,
- Patient under steroid, quinolones,
- Diabetic patient.

RECOMMANDATION

It is recommended to carefully read the instructions for use available in the package insert.

DEVICES

Instruments: Single Use Instruments: Class IIa - CE2797

REIMBURSEMENT

Reimbursement may vary from countries to countries. Check with local authorities.

MANUFACTURER

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DOCUMENT

Reference : BR-DIG-Akilink-EN-032020

Availability of these products might vary from a given country or region to another, as a result of specific local regulatory approval or clearance requirements for sale in such country or region.

Always refer to the appropriate instructions for use for complete clinical instructions.

Non contractual document. The manufacturer reserves the right, without prior notice, to modify the products in order to improve their quality.

CAUTION: Federal law (USA) restricts this device to sale and use by, or on the order of a physician.

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