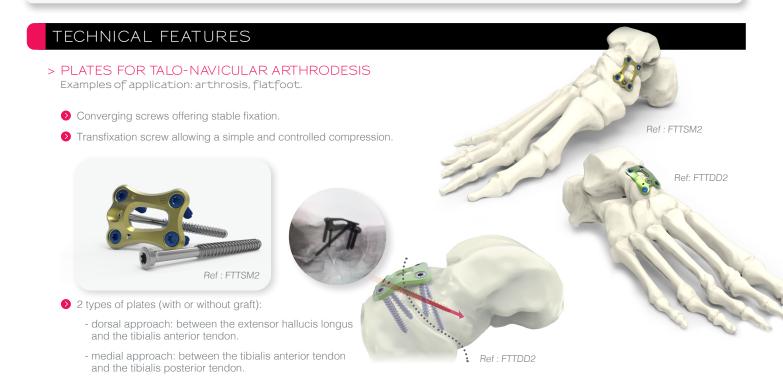
## FOOTMOTION PLATING SYSTEM TALO-NAVICULAR ARTHRODESIS

Indications: The implants of the Footmotion Plating System are intended for arthrodeses, fractures and osteotomies fixation and revision surgeries of the foot in adults.

# N E W C L I P T E C H N I C S

#### Contraindications :

- Serious vascular deterioration, bone devitalization.
- Pregnancy.
- Acute or chronic local or systemic infections.
- · Lack of musculo-cutaneous cover, severe vascular deficiency affecting the concerned area.
- Insufficient bone quality preventing a good fixation of the implants into the bone.
- Muscular deficit, neurological deficiency or behavioral disorders, which could submit the implant to abnormal mechanical strains.
- Allergy to one of the materials used or sensitivity to foreign bodies.
- Serious problems of non-compliance, mental or neurological disorders, failure to follow post-operative care recommendations.
- Unstable physical and/or mental condition.



### REFERENCES

Ø2.8 MM SCREWS		
Ref.	Description	
SLT2.8Lxx	Locking screw - Ø2.8 mm - L10 mm to L34 mm (2 mm incrementation)	
RLT2.8Lxx	Non locking screw - Ø2.8 mm - L10 mm to L34 mm (2 mm incrementation)	

#### Ø3.5 MM SCREWS

Ref.	Description
SLT3.5Lxx	Locking screw - Ø3.5 mm - L10 mm to L40 mm (2 mm incrementation)
RLT3.5Lxx	Non locking screw - Ø3.5 mm - L10 mm to L40 mm (2 mm incrementation)
QLT3.5Lxx-ST	Lag screw - Ø3.5 mm - L20 mm to L40 mm (2 mm incrementation) - STERILE

	TALO-NAVICULAR ARTHRODESIS	
Ref.	Description	Leng
FTTGD1	Talo-Navicular Arthrodesis plate - Left - Size 1	22.3 n
FTTDD1	Talo-Navicular Arthrodesis plate - Right - Size 1	22.3 n
FTTGD2	Talo-Navicular Arthrodesis plate - Left - Size 2	28.3 n
ETTDD2	Talo-Navicular Arthrodesis plate - Bight - Size 2	28.3 n



	MEDIAL PLATES FOR TALO-NAVICULAR ARTHRODESIS	
Ref.	Description	Length
FTTSM1	Medial Talo-Navicular Arthrodesis plate - Size 1	21 mm
FTTSM2	Medial Talo-Navicular Arthrodesis plate - Size 2	26 mm



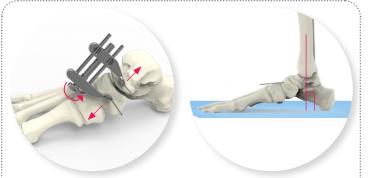
The associated instrumentation and screws are available in the **Footmotion Plating System** set.

#### Remark :

Please note that all implants are also available in sterile packaging. An 'ST' code is added at the end of the reference. Ex : «SLT2.8L10-ST»

## SURGICAL TECHNIQUE

Example using the medial size 2 talo-navicular arthrodesis plate (FTTSM2)



1. The incision is made approximately 5 mm above the tibialis posterior tendon.

The incision for the dorsal talonavicular arthrodesis plate is made between the tibialis anterior tendon and the extensor hallucis longus\*. 2. TN joint exposure and preparation:

Using the opening wedge instrument (ANC642), distract the surface. Then, prepare the joint (a curved curette or a curved lambotte is advised). It is recommended to position the foot support normal to the tibial axis.

Temporarily fix the talo-navicular joint using a Ø1.6 mm pin (33.0216.150).



 Position the plate on the medial side of the talo-navicular joint and insert two Ø1.2 mm pins (33.0212.070) in the navicular part.



- 4. Fix the Ø2.7 mm threaded guide gauge for Ø3.5 mm screws (ANC577) in the two locking holes in the navicular part. Drill using the Ø2.7 mm quick coupling drill bit (ANC591) and measure the necessary screw length.
- **NB**: the depth can also be determined by using the length gauge (ANC589).



5. Insert two Ø3.5 mm locking screws (SLT3.5Lxx) using the T8 screwdriver (ANC575).



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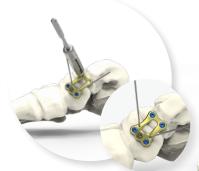
6. Fix the Ø2.7 mm threaded guide gauge for Ø3.5 mm screws (ANC577) in the transfixation hole. Drill and measure.



7. Insert one Ø3.5 mm lag screw (QLT3.5Lxx-ST) using the T8 screwdriver (ANC575).

NB: remove the Ø1.6 mm pin before achieving the compression.

IDENTICAL



 Fix the Ø2.7 mm threaded guide gauge for Ø3.5 mm screws (ANC577) in the two locking holes in the talus part. Drill and measure. Then, insert the two last screws (SLT3.5Lxx) and remove the pins.

orders@newcliptechnics.com

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A Final tightening of the screws must be performed by hand.

The information presented in this brochure is intended to demonstrate a NEWCLIP TECHNICS product. Always refer to the package insert, product label and/or user instructions before using any NEWCLIP TECHNICS product. Surgeons must always rely on their own clinical judgment when deciding which products and techniques to use with their patients. Products may not be available in all markets. Product availability is subject to the regulatory or medical practices that govern individual markets. Please contact your NEWCLIP TECHNICS representative if you have questions about the availability of NEWCLIP TECHNICS products in your area.

NEWCLIP TECHNICS (HQ)NEWCLIP TECHNICS IBERIAPA de la Lande Saint Martin<br/>45 rue des GarottièresCalle Frederic Mompou, 4b<br/>Sant just Desvern44115 Haute-Goulaine, France<br/>+33 (0)2 28 21 23 2508960 Barcelona, España

contact@newclipiberia.com www.newcliptechnics.com NEWCLIP TECHNICS USA Newclip USA 642 Larkfield Center Santa Rosa CA 95403, USA +1 707 230 5078 customerservice@newclipusa.com

tomerservice@newclipusa.co www.newclipusa.com NEWCLIP TECHNICS GERMANY

+49 (0)821 650 749 40

info@newclipgmbh.com

www.newclipgmbh.de

Newclip GmbH Pröllstraße 11, D-86157 Augsburg, KKI Deutschland Ta

#### NEWCLIP TECHNICS JAPAN

Newclip Technics Japan K.K. KKK Bldg. 502, 3-18-1 Asakusabashi Taito-Ku, Tokyo, 111-0053, Japan +81 (0)3 58 25 49 81

www.newcliptechnics.com

NEWCLIP TECHNICS AUSTRALIA

Newclip Australia 3B/11 Donkin Street West End 4101, Australia +61 (0)2 81 886 110 solutions@newclipaustralia.com

www.newcliptechnics.com