

Instructions for L1322 Polycentric Pneumatic 4 Bar Knee With Manual Lock



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1. Description and purpose

These instructions are for use by the practitioner.

- The L1322 knee is to be used exclusively as part of a lower limb prosthesis.
- Recommended for amputees K2 to K3.
- Weight limit for a user is up to 125kg / 275lbs•
- Only ST&G Pylons are recommended with ST&G Knees due to tolerance variability among different manufacturers

Contra-indications

- Residual muscular weakness, contractures or proprioceptive dysfunction including poor balance.
- Contra lateral joint instabilities or pathology
- Complicated conditions involving multiple disabilities



Ensure that the user has understood any IFU (Instructions For Use), drawing particular attention to the safety information.

Product Code

•L1322

Polycentric Pneumatic 4-Bar Knee (Aluminum)



125kg
275lb

2. Construction

Principal Parts

- Frame Aluminum Alloy, Brass, Stainless Steel, Steel
- Knee Aluminum Alloy, Stainless Steel
- Knee control Various materials principally Aluminum Alloy, Stainless Steel, Poly Urethane, Pneumatic Cylinder

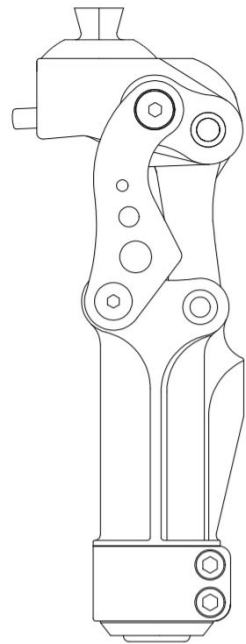
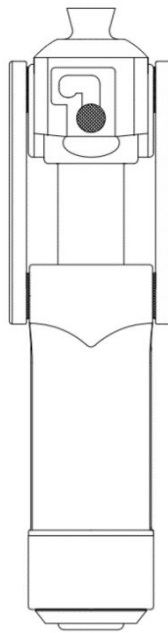
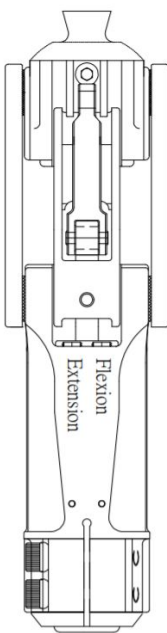


Fig. 1 (a) Posterior View

(b) Anterior View

c) Lateral View

3. Function

- Adjustable extension assist
 - Independently adjustable pneumatic flexion and extension resistances
 - Pyramid mount
 - 30mm Distal Tube Clamp
 - Built-in bearings which allow for ultra smooth walking movement
 - Frame construction of superlight aluminum alloy
 - Anterior mounted manual lock for full extension position
-

4 Safety Information



The Caution symbol highlights safety information which must be followed carefully.



Be aware of finger trap hazard at all times



Any changes in performance of the knee e.g. any change in how the knee functions should be immediately reported to the Clinician / Practitioner



Any excessive changes in heel height may adversely affect the ability to lock the knee, and the stability of the prosthesis



The user should be advised to contact their Clinician / Practitioner if their condition changes.



Clinician / Practitioner is responsible for assessing that patient is appropriate for this device.



Clinician / Practitioner is responsible for instructing patient of relevant content related to operation and safety of this device.



ST&G products are manufactured and tested for a particular weight and activity impact level. Use by another user for whom it was not originally manufactured may cause injury and voids any written or implied warranty or liability. The liability shall fall upon those that resale/provide the product to anyone other than the original user the foot was delivered to. Not to be resold by private party.



Manual Lock feature only intended for temporary circumstances in fully extended position!

5 Maintenance

- Maintenance must be carried out by qualified personnel.
- Bi-Annual inspection to be sure lock and knee function is satisfactory.
- Check for visual defects that may affect proper function.
- A loaner system is available should servicing be required.

The wearer should be advised:

Any changes in performance of this device must be reported to the Clinician / Practitioner.

Changes in performance may include:

- Inability to lock during weight activation
- Any unusual noises

Cleaning:

- Use a damp cloth and mild soap to clean the outside surfaces.
- DO NOT use aggressive cleaning agents.
- DO NOT use any petroleum lubrication on pivots or lock mechanism as this will void the warranty, and could attract excessive dust or aggregate which can accelerate wear and tear on the component.
- If the limb/knee comes into contact with salt or chlorinated water, it should be rinsed with fresh water and dried.

6 Limitations on use

Intended Life:

- Service life of the product is covered by the warranty period (1 year)
- This product is recommended for use with other ST&G Products.

Lifting Loads:

Amputee weight and activity is governed by the stated limits.
Combined amputee, and carrying load, should not be at, or exceed stated weight limit.

Environment:

Store at room temperature, and avoid extremely hot, cold, or moist storage environments Storage Temperature Range: between temperatures of -10°C to 50°C [14°F and 122°F]

Do not store in wet or very hot environment.

Allowable environmental conditions

Use – relative humidity: 0 % to 90 %, non-condensing

Storage/transportation – relative humidity: 20 % to 90 %, non-condensing

Splash - Fresh water, rain, salt water, urine, dust, sand, particles of foam cosmetics

cleaning required after contact with salt-laden air, salt water, chlorinated water, urine, sand

Unallowable environmental conditions

Storage/transportation: high vibrations, impacts

Cleaning agents containing solvents



7. Alignment and Set-Up

 Users be aware of potential finger trap hazard

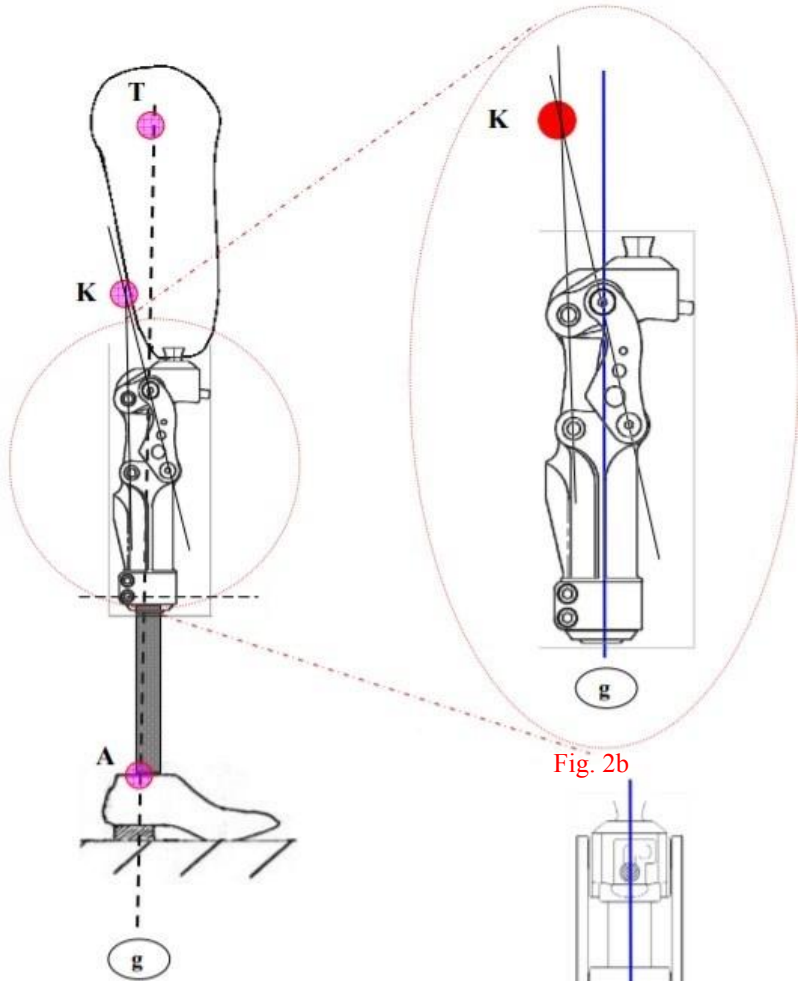


Fig. 2a

Fig. 2b

Fig. 2c

The line of gravity (LOG) shall pass through the 1st axis (upper-front axis) initially.

7.1 BENCH ALIGNMENT:

Note: 4-bar knees inherently are very stable due to the geometry built into each design. This is commonly referred to as the Instant Knee Center (IKC). The IKC point when doing bench alignment, will fall behind the traditional TKA line that we will reference. (Fig. 2a,2b) Tg line in Fig. 3 is ideal placement, but in certain instances, it may be necessary to accommodate placement anteriorly (0 to 10mm). The Tg line is referencing a moving A/P weight bearing line, so it could be slightly anterior or neutral.

a) With prosthesis assembled, taking into account hip flexion contractures, abduction, Line Of Progression, and toe out (Fig.2a), the TKA plumb line should pass through the knee center (center of proximal/anterior pivot Fig.2a, 2b) and in front of the K point (IKC).

NOTE: Take into account shoe heel height, and add 3mm safety factor.

b) Ideally, the pylon connecting the knee and foot should end up vertical. There may be a variance due to the foot alignment recommendations. In this case, the maximum anterior tilt of the pylon should not exceed 3 degrees, and it may be necessary to utilize 1222T offset tube clamp adapter.

c) With prosthesis donned, the weight line should pass through the centerline of the knee in the Coronal or M/L plane (Fig. 2c). Excessive outset or inset will put undue stress on the knee joint.

d) With prosthesis donned, the weight line for Sagittal or A/P plane should have the plumb line passing ideally through the knee center (proximal anterior pivot), and be perpendicular to the ground. (Fig. 2a, 2b)



It is not recommended to have alignment posterior to the reference line, as it could cause knee instability!

8 Knee Adjustment

8.1 Flexion / Extension Adjustment

Swing Phase Pneumatic setting is pre-set from the factory. Extension or flexion adjustment is only needed if the clinician finds the wearer shows a need for higher or lower walking speeds.

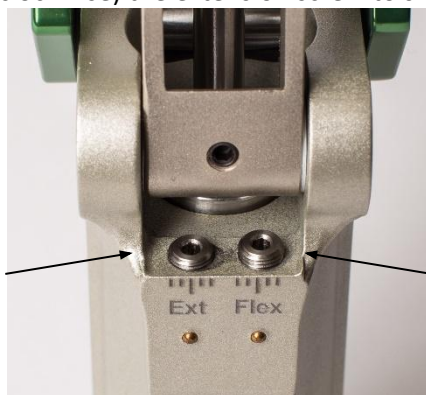
Swing Phase Control Adjustment: It is advisable to adjust **flexion before extension** for optimum walking symmetry. If needed, please follow directions below.

! Ensure full knee extension occurs before performing extension adjustment.

! Use following procedure only if there is a need to adjust extension or flexion:

- 1) Turn extension screw counter-clockwise to lowest resistance - then
- 2) Turn the flexion screw clockwise to set to highest resistance
(Do not over tighten if screw has resistance, or bottoms out – damage may occur and void warranty!)
- Incrementally loosen (counter-clockwise) the flexion screw to adjust heel lift;
- Incrementally tighten (clockwise) the extension screw to smoothly stop extension.

Adjust Extension
screw with
2.5mm hex
wrench



Adjust Flexion
screw with
2.5mm hex
wrench

Extension Resistance Adjustment: (Figure above right)
Using a 2.5mm hex wrench, turn Extension adjustment screw:
Clockwise increases knee extension resistance.
Anti-clockwise reduces knee extension resistance.

Flexion Resistance Adjustment: (Figure above left)
Using a 2.5mm hex wrench, turn Flexion adjustment screw:
Clockwise increases knee flexion resistance.
Anti-clockwise reduces knee flexion resistance.

For more info on Pneumatic adjustments, refer to STNGCorporation on youtube for relevant video material

8.2 Manual Lock Lever

Locked position is in the upper most resting notch position to the left



Unlocked position is in the lower most resting notch position to the right



Note: Manual lock only works by pulling out the lock button detent, then sliding to either locked or unlocked position!



9 Maintenance of Knee Unit



9.1 Servicing Flexion and Extension Stop Bumpers

Use a small screw driver to pick out the rubber bumpers on the back of knee head. Apply glue to new ones and insert back into knee head.

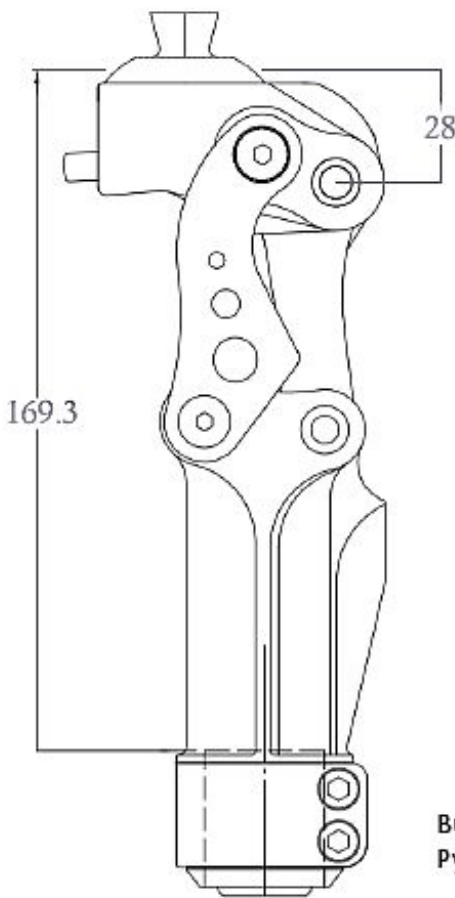
Use a small screw driver to pick out the extension stop rubber bumper on knee level adjusting screw. Insert new one into slot.



10 Technical Specification

- Operating & Storage Temperature Range: -10°C to 50°C (14°F to 122°F)
- Weight: 875g
- Recommended Activity: K2 to K3
- Maximum User Weight: 125kg (275lbs)
- Maximum flexion angle: 135 degrees
- Proximal Alignment attachment: Male Pyramid
- Lock Mechanism: Manually positioned lever
- Distal Alignment attachment: Tube Clamp
- Tube clamp torque setting: 12Nm
- Pyramid Build Height to KC: 28mm
- Build Height: 169.3mm

•Materials: Aluminum Alloy, Stainless Steel, Steel, Rubber



Build Height:
Pyramid = 169.3mm

10 Warranty

Warranted for 2 years from the date of invoice by ST&G.

The user should be aware that changes or modifications not approved will void the warranty.

ST&G products are manufactured and tested for a particular weight and activity impact level. Use by another user for whom it was not originally manufactured may cause injury and shall void any written or implied warranty or liability. The liability shall fall upon those that resale/provide the product to anyone other than the original user the foot was delivered to. Not to be resold by private party.

11 Liability

The manufacturer recommends using the device only under the specified conditions and for the intended purposes. The device must be maintained according to the instructions for use supplied with the device. The manufacturer is not liable for damage caused by the component combinations that were not authorized by the manufacturer.

CE Conformity



This product meets the requirements Council EU 2017/745 (MDR). for medical products. This product has been classified as a class I product according to the classification criteria outlined in appendix IX of the guidelines. Please keep this manual in safe place for future use.

Useful Life

Service life of the product is covered by the warranty period.

Disposal

The device and its packaging must be disposed of in accordance with respective national/local environmental regulations.

L1322IFU REV H (4-30-25)

Specifications subject to change