

# Spider Monkey™ Bone Transport System

## PATIENT QUICK REFERENCE GUIDE



**SPIDER  
MONKEY™ TTT**

**Paragon<sup>20</sup>**

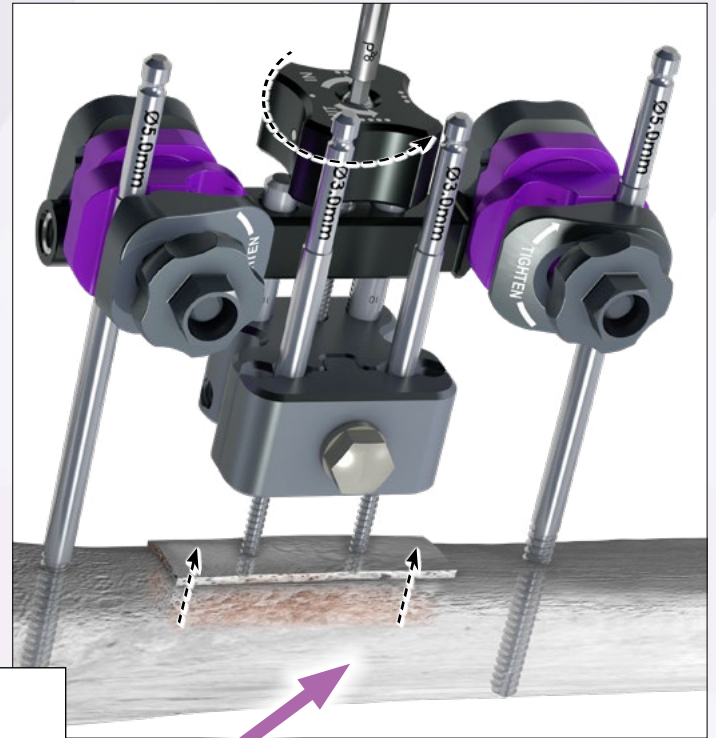
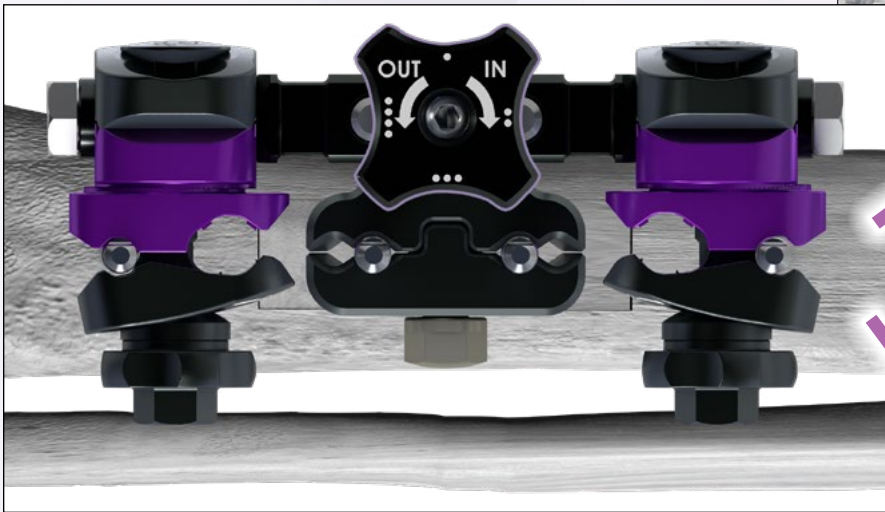
a  ZIMMER BIOMET company



### BONE TRANSPORTER

- Patient Adjustment Knob
- 4 Positions
- Audible click each turn
- OUT/IN Direction

Position One



OUT (turn left)

IN (turn right)



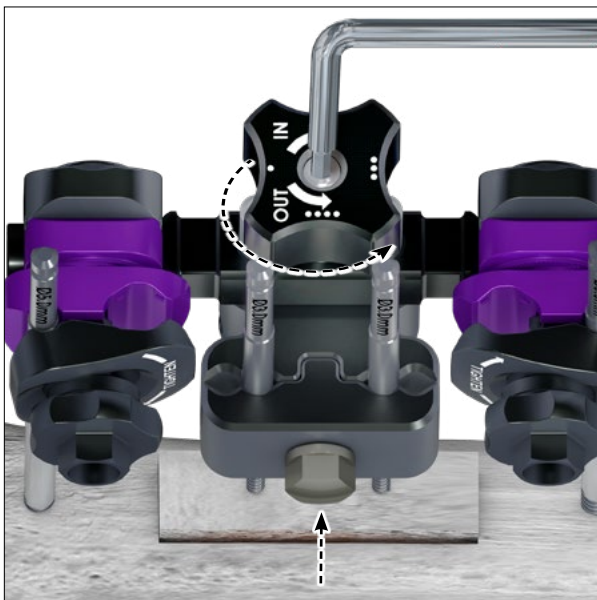
### OUT TURNS

- Number of dots increases
- Segment moves out
- Knob can be adjusted by hand or with the hex driver

NOTE: Your device may be at a different position when you start adjustment. Start the process at the position of your device. Position 1 is shown for illustrative purposes only.

Position One

3 mm Hex Driver



Position Two



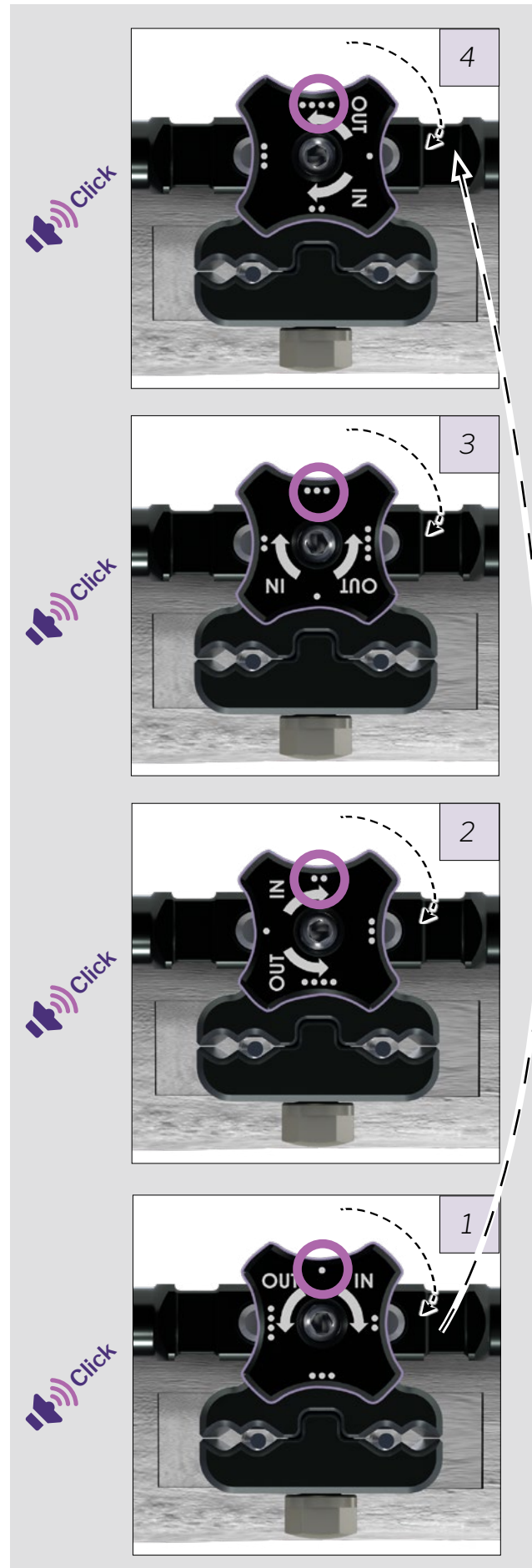
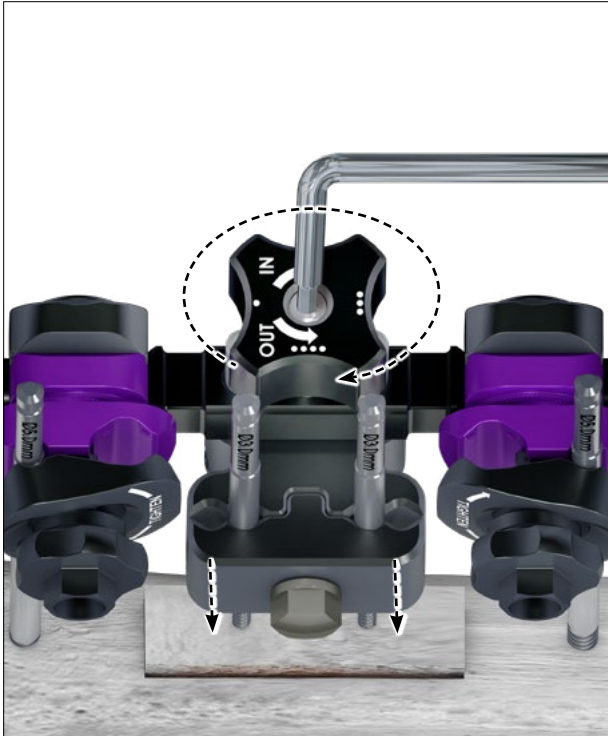
Position Three

Position Four

**IN TURNS**

- Number of dots decreases
- Segment moves in
- Knob can be adjusted by hand or with the hex driver

NOTE: Your device may be at a different position when you start adjustment. Start the process at the position of your device. Position 4 is shown for illustrative purposes only.

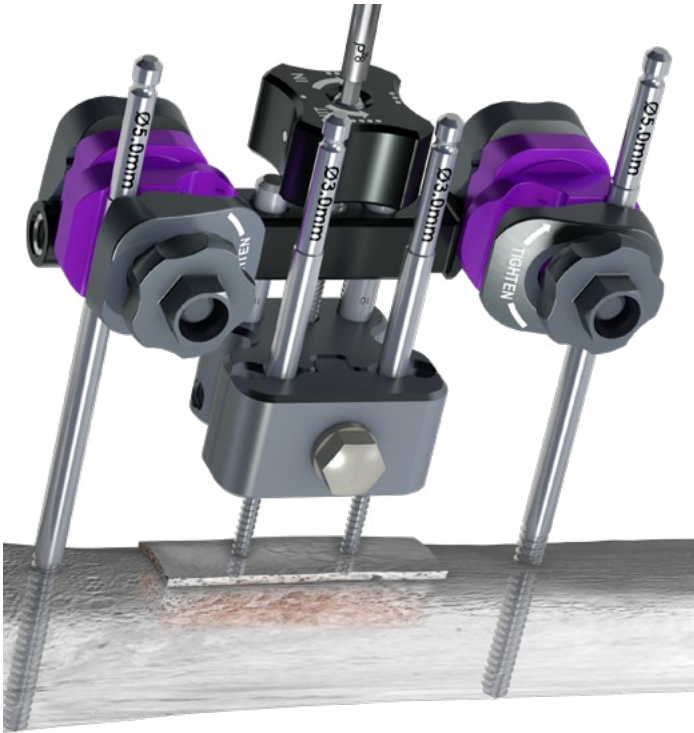


## PATIENT USE

You will follow the post-operative instructions provided by your health care provider.

Your healthcare provider will instruct you or your caregiver on the correct use of the device.

OUT (turn left)



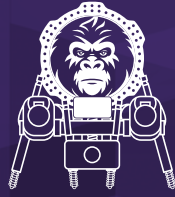
IN (turn right)



**DO NOT** perform any adjustments not included in your instructions.

## DEVICE INFORMATION/PURPOSE

- Information about Indications, Contraindications, Warnings and Precautions, and Potential Adverse Events relevant to the Spider Monkey™ Bone Transport System are contained in the Instructions for Use document of the Spider Monkey™ Bone Transport System P38-IFU-0001.
- Refer to [www.paragon28.com/ifus](http://www.paragon28.com/ifus) for the complete and most current instructions for use document.
- If you experience any difficulties with adjustment of the device, contact your health care provider.
- You will not perform any maintenance or cleaning of the device. The health care provider shall inform you about the need for periodic medical follow-up and of the removal of the medical device in the future.
- The health care provider shall warn you about the surgical and residual risks and make you aware of possible adverse events. A successful result is not achieved in every surgical case. Additional complications may develop at any time due to improper use, medical reasons or device failure that require surgical reintervention to remove or replace the medical device.



# SPIDER MONKEY™ TTT

## Patient Quick Reference Guide

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