# PATIENT SPECIFIC TALUS SPACER

#### THE ONLY FDA APPROVED 3D PRINTED TOTAL TALUS IMPLANT

- The FIRST and ONLY Patient Specific Total Talus Replacement Implant approved by the FDA
- Expected to provide PAIN RELIEF and PRESERVE MOTION of the ankle joint



Indicated for Avascular Necrosis of the ankle joint\*

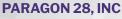
 Patient specific technique guide provided for PRE-PLANNED anatomy correction

 MULTIPLE IMPLANT SIZES provided for intra-operative selection

Surgeon choice of implant material

- Cobalt Chromium

- Titanium with Titanium Nitride coating



44 Riverdale Avenue, Monmouth Beach, NJ 07750

#### PSI@Paragon28.com

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



\*The Paragon 28® Patient Specific Talus Spacer is indicated for avascular necrosis of the ankle joint. The anatomical landmarks necessary for the design and creation of the Additive Orthopaedics Patient Specific Talus Spacer must be present and identifiable on computed tomography scan.

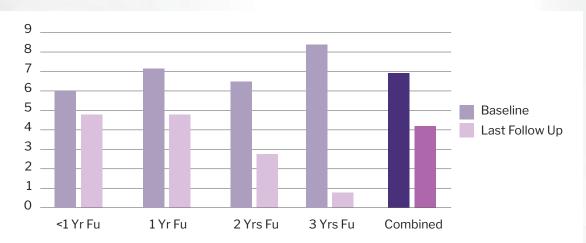
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# REDUCTION IN VAS PAIN SCORES

### **IMPROVEMENT IN QUALITY OF LIFE**

### VAS Pain (cm)

Mean Baseline and Last Follow-Up by Duration of Follow-Up

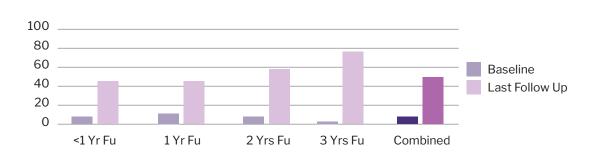




Patients who received a Paragon 28® Patient Specific Talus Spacer in the clinical study showed a reduction in baseline VAS Pain Scores, improvement in ROM, and improvement of functional outcomes based on FAOS subscales, including pain, symptom, activities of daily living, ability to perform sports and recreational activities, and foot/ankle-related quality of life. Please refer to the IFU for complete Clinical Data information.

## **Quality of Life**

Mean Baseline and Last Follow-Up by Duration of Follow-Up



TO WORK WITH US, PLEASE REACH OUT TO YOUR LOCAL PARAGON 28° REPRESENTATIVE OR CONTACT US DIRECTLY: PSI@PARAGON28.COM

Part # AOPSTSPB-01 RevA [2021-12-14]

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Paragon 28, Inc. 44 44 Riverdale Avenue Monmouth Beach, NJ 07750 (855)786-2828