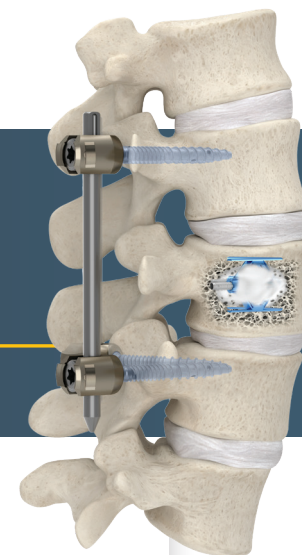


Clinical cases collection

SpineJack system



Pre-op images

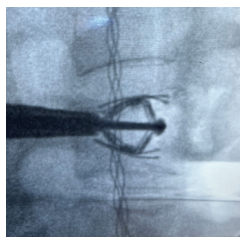


Pre-op MRI (lateral view)

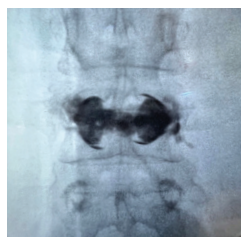


Pre-op CT scan (axial view)

Surgical procedure



Intra-op fluoro (lateral view)



Post-op fluoro (AP view)



AIRO intra-op CT scan (lateral view)



AIRO intra-op CT scan (lateral view)

Physician

Jose Valerio, MD, MBA, MBS, FAANS, FACS

Assisted by: **Yeliana Mayor Pinchevski, PA-C, MPAP**
Miami, Florida

Clinical case

Patient: Female, 71

Level: L3, L2-L4

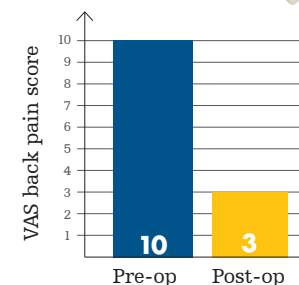
Reason: Traumatic vertebral compression fracture

Case presentation

The patient fell down the stairs and experienced immediate pain in her back after she heard a popping sound. A CT scan confirmed that an A3.2 traumatic fracture occurred at the L3 vertebrae. The vertebra was split in the coronal, sagittal and axial planes. A further CT scan and MRI showed retropulsed fragments that affected the anterior longitudinal ligament (ALL) and posterior longitudinal ligament (PLL). While the ALL and PLL were intact, they were both buckling. The patient presented with 10/10 pain and was unable to walk after the incident.

In this case, the patient was treated using posterior instrumentation along with the SpineJack system at the affected level. During the procedure, two 5.8 mm SpineJack implants were expanded to the maximum height of 20 mm and cemented in place with 4.5 ccs of VertaPlex HV bone cement. Stryker screws were placed percutaneously at L2 and L4 for segment stabilization. CT imaging from Stryker's Airo TruCT showed instrumented ligamentotaxis, stabilization

Visual analog scale



of the burst fracture and anatomical restoration of 17.4 mm. Following the surgery, the patient stated she had 3/10 pain.

According to Dr. Valerio, a younger patient may potentially have had a more invasive procedure, including a corpectomy and a longer construct. Midline vertebral body height restoration was achieved through the use of the SpineJack implants.

Interventional Spine

Bone cement: Serious adverse events, some with fatal outcome, associated with the use of bone cements for vertebroplasty, kyphoplasty and sacroplasty include myocardial infarction, cardiac arrest, cerebrovascular accident, pulmonary embolism and cardiac embolism. Although it is rare, some adverse events have been known to occur beyond one year post-operatively. Additional risks exist with the use of bone cement. Please see the IFU for a complete list of potential risks.

This document is intended solely for the use of healthcare professionals. A physician must always rely on his or her own professional clinical judgment when deciding whether to use a particular product when treating a particular patient. We do not dispense medical advice and recommend that physicians be trained in the use of any particular product before using it.

The information presented is intended to demonstrate Stryker's products. A physician must always refer to the package insert, product label and/or instructions for use, including the instructions for cleaning and sterilization (if applicable), before using any of Stryker's products. Products may not be available in all markets because product availability is subject to the regulatory and/or medical practices in individual markets. Please contact your representative if you have questions about the availability of Stryker's products in your area.

Stryker or its affiliated entities own, use, or have applied for the following trademarks or service marks: SpineJack, Stryker and VertaPlex HV. All other trademarks are trademarks of their respective owners or holders.

The absence of a product, feature, or service name, or logo from this list does not constitute a waiver of Stryker's trademark or other intellectual property rights concerning that name or logo.

D0000114714

Copyright © 2022 Stryker

Stryker Instruments
1941 Stryker Way
Portage, MI 49002