PRESS RELEASE

Axis Spine Gains FDA Clearance for Axis-ALIF System

May 27, 2020

Axis Spine Technologies have received FDA 510(k) clearance for their first implant system, the Axis-ALIF.

Axis has developed a platform technology of build in-situ, modular cages which offer numerous advantages over conventional cage designs. The modular endplates are inserted in a closed position, atraumatically helping to preserve the structural integrity of the vertebral endplate.

Once *in-situ*, the appropriate modular core is inserted and locked. The multiple combinations of endplates and cores provide surgeons with unmatched sagittal as well as coronal correction capabilities.

Jon Arcos, CEO & Founder, said "We are really pleased and grateful to have received such timely clearance from The FDA given the current situation. We now have the opportunity to bring this technology and its benefits to American patients ahead of plan. The next few months will be a very exciting time for the company."

Contact: Jon Arcos
CEO & Founder

Mobile : +44 (0) 7399 499 579
Office : +44 (0) 3301 230 535
Email : jarcos@axisspinetech.com
Web : www.axisspinetech.com

Axis Spine Technologies Ltd. is a company registered in England and Wales. Registered Number: 10287664. Registered Office address: 153-155 London Road, Hemel Hempstead, Herts HP3 9SQ. UK.