# CONCORDE LIFT™ EXPANDABLE INTERBODY DEVICE

## INTRODUCTION

DePuy Synthes is excited to introduce the latest addition to the clinically established CONCORDE® System: The **CONCORDE LIFT™ Expandable Interbody Device.** Designed as a solution for collapsed interbody spaces, the CONCORDE LIFT Implant provides the flexibility of an expandable cage with continuous expansion mechanism for stability and height restoration during interbody fusion procedures.

The sterile packed implant is offered in Convex and Lordotic configurations with multiple footprints. The streamlined set of instruments is designed for efficient endplate preparation, implant insertion, graft delivery into the disc space and graft delivery into the implant post-expansion.

## VALUE PROPOSITION

**The CONCORDE LIFT Expandable Interbody Device provides a robust expandable titanium cage with instrumentation that delivers control and performance to clinicians through tactile feedback and reliable graft delivery for the TLIF approach. CONCORDE LIFT Expandable Interbody Device is a component of a procedural solution which includes CONCORDE® Clear MIS Discectomy Device and VIPER PRIME™ System.**

## IMPLANT SET (CONLPA)

| | | |
|---|---|---|
| **1978-09-021C** | CONCORDE LIFT™, Convex 9x21 mm, 8-13 mm |  |
| **1978-09-026C** | CONCORDE LIFT™, Convex 9x26 mm, 8-13 mm | |
| **1978-11-021C** | CONCORDE LIFT™, Convex 11x21 mm, 8-13 mm | |
| **1978-11-026C** | CONCORDE LIFT™, Convex 11x26 mm, 8-13 mm | |
| **1978-09-023L** | CONCORDE LIFT™, Lordotic 9x23 mm, 10-15 mm |  |
| **1978-09-027L** | CONCORDE LIFT™, Lordotic 9x27 mm, 10-15 mm | |
| **1978-11-023L** | CONCORDE LIFT™, Lordotic 11x23 mm, 10-15 mm | |
| **1978-11-027L** | CONCORDE LIFT™, Lordotic 11x27 mm, 10-15 mm | |

## INSTRUMENT SET (CONLPB)

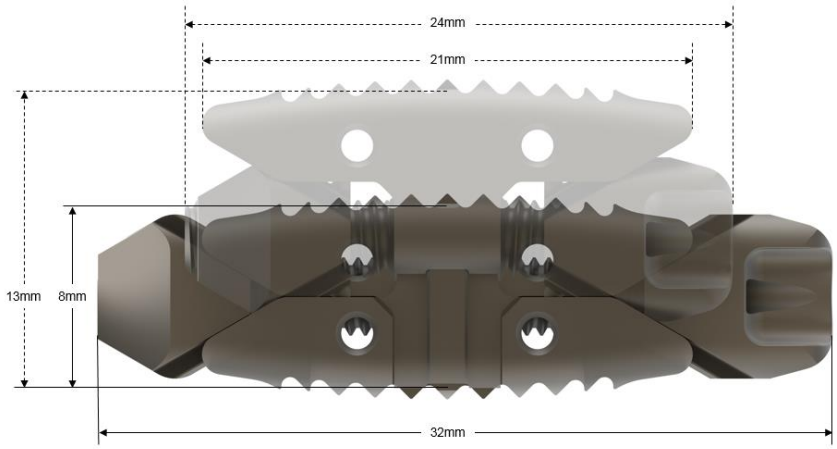
| | |
|---|---|
| **2878-80-117** | CONCORDE LIFT™ Tether |
| **2878-90-004** | CONCORDE LIFT™ Outer Sleeve |
| **2878-04-101** | CONCORDE LIFT™ Driver Shaft |
| **2878-90-010** | CONCORDE LIFT™ Torque Limiting Handle |
| **2878-04-107** | CONCORDE LIFT™ DIST/SHVR 5X7 |
| **2878-04-108** | CONCORDE LIFT™ DIST/SHVR 5X8 |
| **2878-04-109** | CONCORDE LIFT™ DIST/SHVR 5X9 |
| **2878-04-110** | CONCORDE LIFT™ DIST/SHVR 5X10 |
| **2878-04-111** | CONCORDE LIFT™ DIST/SHVR 5X11 |
| **2731-28-001** | Modular T-Handle |
| **2878-04-104** | CONCORDE LIFT™ CNVX TRIAL (9x8) |

| | |
|---|---|
| **2878-04-105** | CONCORDE LIFT™ LRD TRIAL (9x10) |
| **2878-04-114** | CONCORDE LIFT™ Large Graft Delivery |
| **2878-80-017** | CONCORDE LIFT™ Graft Plunger |
| **2878-04-113** | CONCORDE LIFT™ Small Graft Delivery |
| **2878-04-119** | CONCORDE LIFT™ Flex Graft Plunger |
| **2731-13-001** | Small Dilator |
| **2864-10-019** | Leopard Slap Hammer |
| **2797-92-109** | Generic Lid Assembly |
| **2878-04-116** | CONCORDE LIFT™ Implant Case |
| **2878-04-115** | CONCORDE LIFT™ Instrument Case |

## CAGE DIMENSIONS

| | Convex | | Lordotic | |
|---|---|---|---|---|
| | **9x21 / 11x21** | **9x26 / 11x26** | **9x23 / 11x23** | **9x27 / 11x27** |
| **Fully collapsed (mm)** | 32 | 36 | 32 | 36 |
| **Fully expanded (mm)** | 24 | 28 | 24 | 28 |
| **Endplate length (mm)** | 21 | 26 | 23 | 27 |
| **Expansion range (mm)** | 8-13 | | 10-15 | |

Image below shows 9x21 mm Convex Cage




## INDICATIONS

The CONCORDE LIFT™ Expandable Interbody Device is a lumbar intervertebral body fusion device, and is indicated for use with autogenous bone graft and/or allograft comprised of cancellous and/or corticocancellous bone graft in patients with degenerative disc disease (DDD) at one or two contiguous levels of the lumbar spine, L2 to S1. These DDD patients may also have up to Grade I spondylolisthesis or retrolisthesis at the involved level(s). The CONCORDE LIFT™ Expandable Interbody Device can be implanted via posterior or transforaminal approach.

DDD is defined as discogenic back pain with degeneration of the disc confirmed by history and radiographic studies. Candidates for surgery should be skeletally mature and have had a six-month course of conservative treatment. These patients may have had primary or secondary surgery, but no previous fusion at the involved levels.

The device is not intended to be used as a stand-alone device. It must be used with supplemental internal spinal fixation systems that have been cleared for use in the lumbar spine.

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DePuy Synthes
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