XRL System. A modular expandable radiolucent vertebral body replacement system.



Technique Guide



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XRL System. A modular expandable radiolucent vertebral body replacement system.

The XRL System provides surgeons the ability to expand the device in situ to reconstruct the anterior column, restore height, and correct the sagittal curvature of the thoracolumbar spine.

- Modular construction
- 360° implantation
- Tactile feedback during expansion
- Most implant options to accommodate a wide range of anatomies

Material

- PEEK material offers the benefit of radiolucent imaging, so surgeons can better assess fusion progress and/or tumor recurrence.
- Modulus of elasticity of PEEK is approximately between cancellous and cortical bone to aid in stress distribution and load sharing.



Implant Options

Modular—Flexible use

The Modular implant consists of a central body on which two endplates are attached.

- Central body The octagonal shape permits various approach options
- Endplates
 Numerous footprints and angles
 allow the implant to conform to a
 wide range of patient anatomies
- Endplate screws
 Rigidly secures the endplate to the central body

Integrated (no assembly required)

Optimal for procedures where low profile constructs are needed.

Self-locking expansion mechanism

Distracts and locks at 1 mm increments.

Open architecture

The open central body and endplate design allow generous placement of bone graft.

- Medium implant cannulation 8.4 mm diameter
- Large implant cannulation 13.5 mm diameter











In 1958, the AO formulated four basic principles, which have become the guidelines for internal fixation.¹ They are:

- Anatomic reduction
- Stable fixation
- Preservation of blood supply
- Early, active mobilization

The fundamental aims of fracture treatment in the limbs and fusion of the spine are the same. A specific goal in the spine is returning as much function as possible to the injured neural elements.^{2,3}

1 Müller ME, M Allgöwer, R Schneider, H Willenegger. Manual of Internal Fixation. 3rd ed. Berlin Heidelberg New York: Springer. 1991.

2 Ibid.

3 M Aebi, JS Thalgott, and JK Webb. AO ASIF Principles in Spine Surgery. Berlin; Springer-Verlag, 1998.

Indications

The Synthes XRL device is a vertebral body replacement device intended for use in the thoracolumbar spine (T1–L5) to replace a collapsed, damaged or unstable vertebral body due to tumor or trauma (i.e., fracture). The Synthes XRL device is intended to be used with Synthes supplemental internal fixation systems (e.g., USS, including MATRIX, Pangea, and TSLP). The interior of Synthes XRL can be packed with bone (i.e., autograft or allograft).

The Synthes XRL device is designed to provide anterior spinal column support even in the absence of fusion for a prolonged period.

Contraindications

- 1. Use of the Synthes XRL device is contraindicated when there is active systemic infection, infection localized to the site of the proposed implantation, or when the patient has demonstrated allergy or foreign body sensitivity to any of the implant materials.
- 2. Severe osteoporosis may prevent adequate fixation and thus preclude the use of this or any other orthopaedic implant.
- 3. Conditions that may place excessive stresses on bone and implants, such as severe obesity or degenerative diseases, are relative contraindications. The decision whether to use these devices in patients with such conditions must be made by the physician, taking into account the risks versus the benefits to the patient.
- 4. Use of these implants is relatively contraindicated in patients whose activity, mental capacity, mental illness, alcoholism, drug abuse, occupation or lifestyle may interfere with their ability to follow postoperative restrictions, thereby placing undue stresses on the implant during bony healing. This could result in a higher risk of implant failure.

Please refer to package insert for the full list of indications, contraindications, warnings and/or precautions.

1 Access

Various approaches are suitable depending on the affected spinal level involved.

The following surgical technique is described using a lateral approach from the left at L1.

2 Perform corpectomy

Perform a partial or complete corpectomy as required.

Note: Remove the superficial layers of the entire cartilaginous endplates and expose bleeding bone. Excessive removal of subchondral bone may weaken the vertebral endplate. If the entire endplate is removed, subsidence and a loss of segmental stability may occur.

The XRL Vertebral Body Replacement contains a complete line of central body and endplate trial implants that correspond to each central body and endplate implant. Trials are placed into the corpectomy site intraoperatively to determine the appropriate endplate footprint, angle, and central body height.

1

Determine defect

Instruments	
03.661.010	Metal Tape Gauge
324.092*	Measuring Forceps

The metal tape gauge or measuring forceps can be used to measure overall defect.

Note: If final distracted corpectomy height is less than 32 mm, then skip to Step 4 and use the integrated trials.



*Also available

Select endplate footprint size and angle

Instruments

	XRL Medium Endplate Footprint Trials
03.807.364	21 mm round
03.807.365	21 mm x 24 mm
03.807.366	26 mm x 30 mm
	XRL Large Endplate Footprint Trials
03.807.367	27 mm round
03.807.368	28 mm x 33 mm
03.807.369	30 mm x 40 mm

The endplate footprint trial can be adjusted to accommodate the desired approach. Pull the sleeve (1) and turn the endplate trial to the desired position (2). Release the sleeve to lock the position of the trial.

Determine the footprint using the endplate footprint trial. The handles of the endplate footprint trials are color-coded green and blue to match the medium and large sets, respectively. Determine the angle using lateral x-ray imaging.





Determine central body size

The optimal central body height is calculated using endplate trial height which is found on the back of the module lid for reference. The trials do not account for the implant spikes; therefore, 2 mm clearance on each end of the trial is required.

Optimal Central Body Height (CBH) = Overall defect – Cranial trial endplate height – Caudal trial endplate height – Clearance for spikes

Example for 70 mm defect with a 5° cranial endplate and 0° caudal endplate:

CBH = 70 mm - 6.5 mm - 5 mm - 4 mm CBH = 55 mm

Insert the selected trial endplates onto the trial central body. Align the etch lines before pressing the components together. Ensure there is no gap between the endplate and central body trial.

Note: The endplate height is independent of the footprint.

Warning: The trials are not for implantation and must be removed before insertion of the XRL implant. Total construct angle must not exceed 30° lordosis/kyphosis.

Medium Endplate Trial		Large Er	Large Endplate Trial	
Angle	Height (mm)	Angle	Height (mm)	
0°	5	0°	5.5	
5°	6.5	5°	7	
10°	8.5	10°	9.5	
15°	10.5	15°	12	
-5°	6.5	20°	14.5	
-10°	8.5			



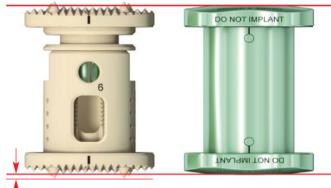
Insert trial

Instruments	
03.807.382	XRL Medium Implant Holder
03.807.384	XRL Large Implant Holder

Using the implant holder, insert the trial into the corpectomy site. Be sure the appropriate endplate is oriented in the cranial/caudal position. The optimal position for the trial is centered on the vertebral bodies with clearance to account for the implant spikes. Trials must always be securely held while in the wound.

Note: Integrated implants do not have tall spikes and therefore the integrated trials are the same height as the corresponding collapsed implant.

Change trial central body and endplates as necessary to achieve the optimal height, angle, and footprint.



2 mm

1 Assemble implant

Select implant based on corresponding trial (see pages 25–26 for cross reference list).

If an integrated assembly is selected, skip to Step 4.

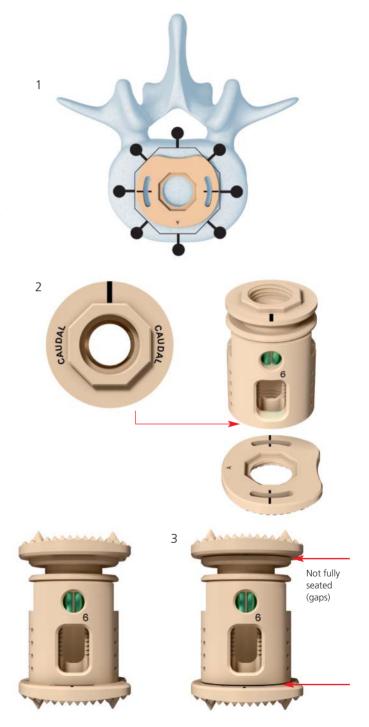
When assembling the implant, orient the caudal endplate with the "A" pointing anterior, and position the central body with the locking ring facing the direction of the desired approach.

Note: The etch line on the ends of the central body, the graft window, and the locking ring may all be used to indicate the direction of approach. Figure 1 shows the orientation of the locking ring with respect to the caudal endplate for each approach option.

Attach the caudal endplate first by pressing the endplate onto the octagon until fully seated (Figure 2). Repeat with the cranial endplate. Ensure that both endplates are in the same direction (lateral left shown).

Warning: When pressing on the endplates, ensure the endplate properly seats on the central body. This can be checked visually (Figure 3). If the endplate is not properly seated, there is a risk that it could detach from the central body.

Warning: The XRL central body should never be implanted without cranial and caudal endplates properly secured with endplate screws.



2 Reposition endplates (optional)

03.807.354	XRL Endplate Removal Tool		

If necessary, the endplates can be repositioned by manually removing them from the central body, except for the round endplates which are removed using the XRL endplate removal tool. Be sure to perform endplate removal over a sterile table.

Warning: Endplates release from central body abruptly. Make sure to have a firm grip on both the central body and the endplate during removal.

To remove round endplates, align the tip of the XRL endplate removal tool with the slot in the endplate. Apply a slight, constant pressure and rotate the tool to release the endplate.



3 Attach endplate screws

Attach endplate screws	
Instruments	
03.807.351	XRL Medium Endplate Screwdriver Tip
03.807.352	XRL Large Endplate Screwdriver Tip
03.807.357	XRL Medium Torque Limiting Handle
03.807.358	XRL Large Torque Limiting Handle



Align the endplate screwdriver tip into the open end of the torque limiting handle.

Press until an audible "click" is heard.

Align the tri-lobal feature of the tip and the etchings on the endplate screw. Lightly press the screw onto the screwdriver tip. The screwdriver tip will retain the screw.

Align the screw with the caudal endplate to prevent cross threading. While gripping the large end of the torque limiting handle, rotate the torque limiting handle clockwise to advance the screw through the caudal endplate and into the central body. Tighten until an audible "click" in the torque handle is heard. Repeat this step to fixate the cranial endplate.

Warning: The torque limiting handles are color-coded green and blue to match the medium and large sets, respectively and may only be used with its corresponding set.

Note: Please follow torque limiting handle calibration instructions to ensure proper functionality.





Prepare implant

Instruments	
instruments	
03.807.371	XRL Medium Graft Packing Tamp
03.807.372	XRL Large Graft Packing Tamp
03.807.374	XRL Medium Graft Packing Preparation Tamp
03.807.375	XRL Large Graft Packing Preparation Tamp

Prior to implanting, use graft packing tamps to facilitate packing of bone graft into the XRL implant. Graft can be packed through the cannulation in the endplate and graft windows.

Warning: Do NOT pack graft into the locking ring. DO NOT use excessive force while packing graft.

Note: Graft packing tamp will not fit inside the window of integrated implant #1.





Assemble spreader instrument

Instruments	
03.807.300	XRL Spreader
03.807.310	XRL Medium Shaft
03.807.311– 03.807.315	XRL Medium Spreader Tops, with 3, 5, 8, 10, or 15 mm distraction
03.807.330	XRL Large Shaft
03.807.331– 03.807.335	XRL Large Spreader Tops, with 3, 5, 8, 10, or 15 mm distraction
03.807.348	XRL Release Tool
03.807.355	XRL Medium Spreader Top, with 5 mm distraction (Integrated)
03.807.356	XRL Large Spreader Top, with 5 mm distraction (Integrated)

Assemble the appropriate size spreader top to the XRL spreader according to the implant central body size selected. The spreader tops are designed to prevent over-distracting the implant.

Note: The underside of the built-in lid of the graphic case shows Spreader Top and its corresponding implant selection.

While holding the spreader with the shaft in the horizontal position, set ratchet lever to the "OFF" position (Figure 1).

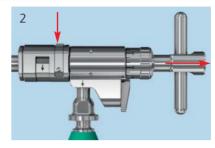
Press T-driver release button and pull back on the T-driver (Figure 2). Release the button to set T-driver in the open position (Figure 3). T-driver should not be fully removed during this operation.

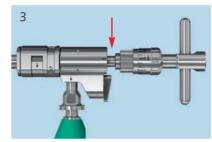
Insert the selected spreader top into the spreader shaft (4) and insert the T-driver (5) by gently pushing and turning the T-driver into the outer shaft assembly.

Check functionality of spreader top by rotating T-driver. If properly assembled, spreader top should translate during T-driver rotation.



4







Secure Implant to Spreader

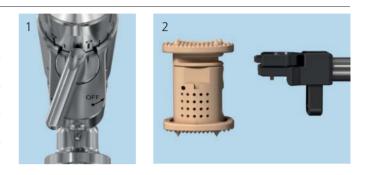
Instruments	
03.807.300	XRL Spreader
03.807.310	XRL Medium Shaft
03.807.311– 03.807.315	XRL Medium Spreader Tops, with 3, 5, 8, 10, or 15 mm distraction
03.807.330	XRL Large Shaft
03.807.331– 03.807.335	XRL Large Spreader Tops, with 3, 5, 8, 10, or 15 mm distraction
03.807.348	XRL Release Tool
03.807.355/ 03.807.356	XRL Spreader Top, Integrated (medium or large)

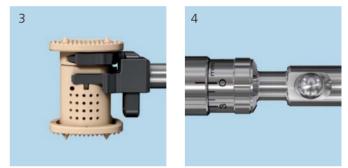
To load the implant, fully collapse the spreader top and set the ratchet lever to the "ON" position (Figure 1).

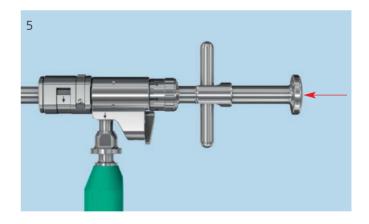
With the locking ring facing the instrument, slide the spreader top into the slots below the cranial endplate (Figure 2). Do not force spreader top onto implant. Slightly turn the T-driver clockwise until the notch on the fork of the spreader shaft engages the implant for a secure hold (Figure 3).

Set the scale to zero (Figure 4).

Completely insert the release tool through the XRL spreader and into the locking ring. An audible "click" will be heard (Figure 5).







7	
Insert implant	
Instrument	
03.807.300	XRL Spreader

The spreader handle can be rotated at 90° increments to aid in visualization. Set ratchet lever to "OFF" position (1). With one hand gripping the spreader shaft, pull back on retaining collar and rotate the spreader handle to the desired position (2,3). Release retaining collar. Verify that the spreader handle is locked into position. Reset scale to zero.

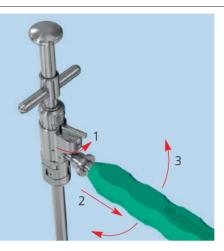
Warning: Do not adjust spreader handle when ratchet lever is set to "ON." This will result in premature distraction of the implant. Do not begin distraction until spreader handle is locked into desired position.

Guide and position the implant with the spreader. Slight distraction of the vertebral bodies may be necessary to ease insertion.

The optimal position for the implant is in the center of the vertebral body endplate. Maintain space around the endplate of the implant to allow peripheral bony fusion.

Warning: Do not impact on spreader. Do not manipulate implant unless both the slot and notch are engaged.

- () Verify the position of the implant using the image intensifier.
 - A titanium locking ring is used to determine orientation of the implant
 - The 1 mm diameter tantalum markers are embedded into the PEEK endplates to provide radiographic markers for intraoperative or postoperative imaging
 - The anterior and medial/lateral markers are located approximately 1 mm from the edges of the implant. The posterior marker is located 1 mm from the edge of the round implant, and 2 mm from the edge of the anatomically shaped endplates. The cranial/caudal locations of the markers are 2 mm from the end of the pyramidal teeth.







8 Distract and check position

Instrument 03.807.300 XRL Spreader

The spreader allows for expansion in both a ratchet mode and continuous mode.

Option A: Ratchet Mode

For ratchet mode, ensure the ratchet lever is set to the "ON" position and the release tool is engaged, then turn the spreader T-driver clockwise (Figure 1) and expand the implant until the desired amount of distraction is achieved.

Option B: Continuous Mode

For continuous mode, ensure the ratchet lever is set to the "OFF" position and the release tool is engaged, then turn the spreader T-driver clockwise and expand the implant until the desired amount of distraction is achieved. With constant clockwise torque on the T-driver, set the ratchet lever to the "ON" position.

Once the implant has been distracted, disengage the release tool, set the release tool to the resting position (Figure 2) and with constant clockwise torque on the T-driver, place the ratchet lever in the "OFF" position.

Close the spreader by turning the T-driver counterclockwise. Remove the spreader from the implant.

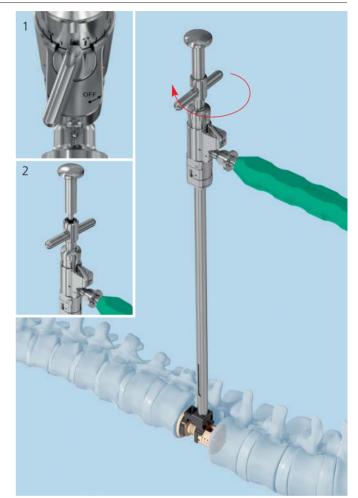
Visually inspect implant/vertebral body interface for gaps to prevent point loading. If a gap is found, repositioning is necessary to ensure full endplate surface contact.

Verify the position of the implant using the image intensifier.

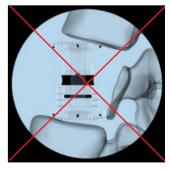
Do not reuse XRL implants once they have been implanted or explanted.

Warning: Do not reposition spreader handle during or after distraction.

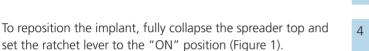
Distraction of the implant is only permitted with the XRL instrument set.







9 Reposition implant (optional)		
Instrument		
03.807.300	XRL Spreader	



Be sure the release tool is disengaged and set to the resting position (Figure 2).

Slide the spreader top into the slots below the cranial endplate. Turn the T-driver clockwise until the notch on the bottom fork engages the implant for a secure hold (Figure 3). Re-engage the release tool until an audible "click" is heard (Figure 4).

Set the ratchet lever to "OFF" position and compress the implant by turning the T-driver counterclockwise. Reposition the implant to the desired location and follow previous step to re-distract implant.

Warning: Do not impact on the XRL endplates when repositioning the implant.

Repositioning of the implant is only permitted with the XRL Instrument Set.





10 Verify lock

When the implant is in its final position, verify the locking ring on the central body is closed. When the slot is approximately 1 mm (Figure 1), the implant is locked and secured. If the slot is larger (Figure 2), re-engage the implant with the spreader, and with the release tool in the resting position, distract the implant slightly to close the locking ring.

Warning: Locking ring must be properly closed to ensure final implant height is maintained.





Apply bone material		
Instruments		
03.807.371	XRL Medium Graft Packing Tamp	
03.807.372	XRL Large Graft Packing Tamp	
03.807.374	XRL Medium Graft Packing Preparation Tamp	
03.807.375	XRL Large Graft Packing Preparation Tamp	

In situ graft packing must not occur until final implant position is achieved, as additional bone graft may obstruct repositioning of the implant.

Before packing additional bone graft in or around the cage, use AP and lateral radiographs to verify the position of the implant in relation to the vertebral bodies using the tantalum beads and locking ring for references.

The graft packing tamp has 2 different ends to fit the corresponding window of the expanded central body. The preparation tamp has an angled end that can be used to gain compression on graft that is not accessible with the graft packing tamp.

Note: Graft packing tamp will not fit inside the window of integrated implant #1, however can still be used to tamp graft material.

Warning: Do not use excessive force while packing graft.



2 Apply supplemental fixation

For spinal stability and to maintain adequate compression on the construct, the XRL system is indicated for use with supplemental fixation.



Modular XRL Implants

XRL Medium

- 21 mm central body diameter
- Endplate footprint options:
 - 21 mm round
 - 21 mm x 24 mm
 - 26 mm x 30 mm
- Construct heights range from 32 mm (fully compressed) to 142 mm (fully expanded)
- Various lordotic/kyphotic angulation options _

XRL Large

- 27 mm body diameter
- Endplate footprint options:
 - 27 mm round
 - 28 mm x 33 mm
 - 30 mm x 40 mm
- Construct heights range from 34 mm (fully compressed) to 145 mm (fully expanded)
- Various lordotic angulation options _

Integrated XRL Implants

XRL Medium

- 21 mm central body diameter
- 21 mm endplate footprint
- Heights range from 22 mm (fully compressed) to 36 mm (fully expanded)
- 0° parallel endplates

XRL Large

- 27 mm body diameter
- 28 mm endplate footprint
- Heights range from 23 mm (fully compressed) to 37 mm (fully expanded)
- 0° parallel endplates











Note: All XRL implants are supplied sterile

XRL Trial Implants

The XRL vertebral body replacement contains a complete line of central body and endplate trials that correspond to each central body and endplate implant. Trials are placed into the corpectomy site intraoperatively to determine the appropriate implant footprint, lordotic angle and central body height.

Use the central body and endplate trials to determine the

Medium Trials (green)

largest implant size (integrated or modular) that will fit the measured corpectomy site. Trials may be secured and lowered into corpectomy defect using the implant holder. Allow 2 mm clearance on each end for the tall spikes on the endplates (modular only).



Description



Part Number

Integrated

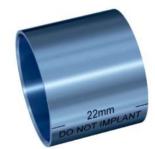
Part Number

Standard





Integrated



Standard

Part Number	Description		Part Number
Trial Implants	Central Bodies (mm)	Size	Corresponding Implants
03.807.501	Integrated 22–25 height, 0°	1	08.807.2015
03.807.502	Integrated 24–29 height, 0°	2	08.807.2025
03.807.503	Integrated 28–36 height, 0°	3	08.807.2035
03.807.504	22–27 height	4	08.807.2045
03.807.505	25–33 height	5	08.807.2055
03.807.506	29–39 height	6	08.807.2065
03.807.507	33–43 height	7	08.807.2075
03.807.508	37–52 height	8	08.807.2085
03.807.509	44–59 height	9	08.807.2095
03.807.510	51–66 height	10	08.807.2105
03.807.511	62–77 height	11	08.807.2115
03.807.512	73–88 height	12	08.807.2125
03.807.513	84–99 height	13	08.807.2135
03.807.514	95–110 height	14	08.807.2145
03.807.515	106–121 height	15	08.807.2155

Part Number	Description		Part Number
Trial Implants	Central Bodies (mm)	Size	Corresponding Implants
03.807.601	Integrated 23–26 height, 0°	1	08.807.3015
03.807.602	Integrated 25–30 height, 0°	2	08.807.3025
03.807.503	Integrated 29–37 height, 0°	3	08.807.3035
03.807.604	22–27 height	4	08.807.304S
03.807.605	25–33 height	5	08.807.3055
03.807.606	29–39 height	6	08.807.3065
03.807.607	33–43 height	7	08.807.3075
03.807.608	37–52 height	8	08.807.3085
03.807.609	44–59 height	9	08.807.3095
03.807.610	51–66 height	10	08.807.3105
03.807.611	62–77 height	11	08.807.3115
03.807.612	73–88 height	12	08.807.3125
03.807.613	84–99 height	13	08.807.3135
03.807.614	95–110 height	14	08.807.3145
03.807.615	106–121 height	15	08.807.3155

Medium Trial Endplates (green)





Large Trial Endplates (blue)





Part Number	Description (mm)	Corresponding Implants
03.807.521	21 Round, 0°	08.807.2215
03.807.522	21 Round, 5°	08.807.2225
03.807.523	21 Round, 10°	08.807.223S
03.807.524	21 Round, 15°	08.807.224S
03.807.531	21 x 24, -10°	08.807.2315
03.807.532	21 x 24, -5°	08.807.232S
03.807.533	21 x 24, 0°	08.807.2335
03.807.534	21 x 24, 5°	08.807.234S
03.807.535	21 x 24, 10°	08.807.2355
03.807.536	21 x 24, 15°	08.807.2365
03.807.541	26 x 30, -10°	08.807.241S
03.807.542	26 x 30, -5°	08.807.2425
03.807.543	26 x 30, 0°	08.807.2435
03.807.544	26 x 30, 5°	08.807.244S
03.807.545	26 x 30, 10°	08.807.2455
03.807.546	26 x 30, 15°	08.807.2465

Part Number	Description (mm)	Corresponding Implants
03.807.621	27 round, 0°	08.807.3215
03.807.622	27 round, 5°	08.807.3225
03.807.623	27 round, 10°	08.807.3235
03.807.624	27 round, 15°	08.807.324S
03.807.625	27 round, 20°	08.807.3255
03.807.631	28 x 33, 0°	08.807.3315
03.807.632	28 x 33, 5°	08.807.3325
03.807.633	28 x 33, 10°	08.807.3335
03.807.634	28 x 33, 15°	08.807.3345
03.807.635	28 x 33, 20°	08.807.3355
03.807.641	30 x 40, 0°	08.807.3415
03.807.642	30 x 40, 5°	08.807.3425
03.807.643	30 x 40, 10°	08.807.3435
03.807.644	30 x 40, 15°	08.807.344S
03.807.645	30 x 40, 20°	08.807.3455

03.807.300

XRL Spreader For implanting, distracting, and compressing (repositioning the implant)

Ratchet Lever

A ratchet lever on the instrument handle allows for the manipulations of the XRL implant.

Note: Release tool must be engaged with locking ring for implant manipulation.



Ratchet Mode "ON" allows expansion of the implant



Continuous Mode "OFF" allows tactile expansion or compression of the . implant

T-driver Release

Allows the T-driver to be disengaged/removed from the spreader

Shaft Release

Allows the spreader shaft to be removed from the from the spreader

Retaining Collar

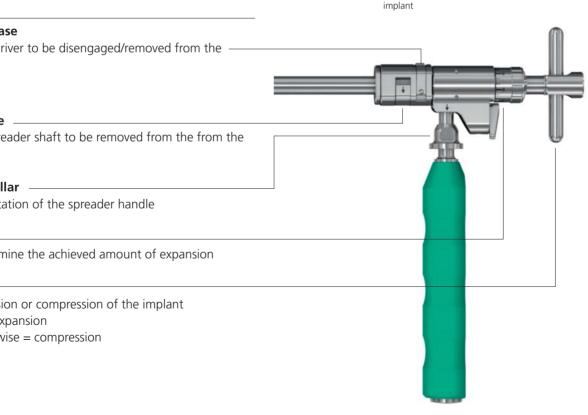
Allows 90° rotation of the spreader handle

Scale -

Used to determine the achieved amount of expansion

T-driver -

Allows expansion or compression of the implant Clockwise = expansion Counterclockwise = compression



03.661.010	Metal Tape Gauge	
03.807.310 03.807.330	XRL Medium Shaft XRL Large Shaft (shown)	
03.807.348	Release Tool Enables implant repositioning	

Spreader tops Medium	
	distraction ranges, dependent on the central
body implant.	
03.807.311	with 3 mm distraction
03.807.312	with 5 mm distraction
03.807.313	with 8 mm distraction
03.807.314	with 10 mm distraction
03.807.315	with 15 mm distraction
03.807.355	with 5 mm distraction (integrated)
Large	
03.807.331	with 3 mm distraction
03.807.332	with 5 mm distraction
03.807.333	with 8 mm distraction

03.807.351 03.807.352	XRL Endplate Screwdriver Tips Medium Large		
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with 10 mm distraction

with 15 mm distraction

with 5 mm distraction (integrated)

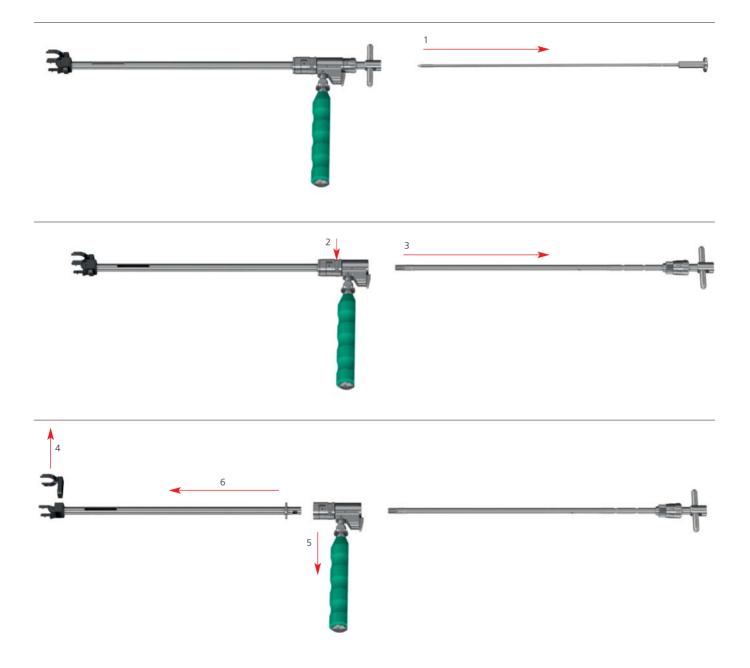
03.807.334 03.807.335

03.807.356

03.807.354	XRL Endplate Removal Tool Allows removal of round endplates from the central body	
03.807.357 03.807.358	XRL Torque Limiting Handles Medium Large	
02.007.204	Endplate Footprint Trials Medium 21 mm round	•
03.807.364 03.807.365	Medium 21 mm round Medium 21 mm X 24 mm	— ———————————————————————————————————
03.807.366 03.807.367	Medium 26 mm X 30 mm Large 27 mm Round	
03.807.368 03.807.369	Large 28 mm X 33 mm Large 30 mm X 40 mm	
	Graft Packing Tamps	
03.807.371 03.807.374	Medium Tamp Medium Preparation Tamp	
03.807.372	Large Tamp	
03.807.375	Large Preparation Tamp	

	Implant Holder
03.807.382	Medium Implant Holder (shown)
03.807.384	Large Implant Holder
	For inserting the implant trials





XRL Medium Implant and Instrument Set (01.807.029)

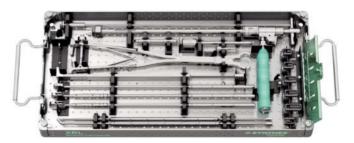
Carry Case and Graphic Case

60.807.026	Carry Case, for XRL Medium Implants
60.807.029	Graphic Case, for XRL Medium Instruments
Instruments	
03.661.010	Metal Tape Gauge
03.807.300	XRL Spreader
03.807.310	XRL Medium Shaft
	XRL Medium Spreader Top
03.807.311	with 3 mm distraction
03.807.312	with 5 mm distraction
03.807.313	with 8 mm distraction
03.807.314	with 10 mm distraction
03.807.314	with 15 mm distraction
03.807.355	with 5 mm distraction (Integrated)
03.807.351	XRL Medium Endplate Screwdriver Tip
03.807.354	XRL Endplate Removal Tool
03.807.364	XRL Medium Endplate Footprint Trial,
	21 mm round
03.807.365	XRL Medium Endplate Footprint Trial,
	21 mm x 24 mm
03.807.366	XRL Medium Endplate Footprint Trial,
	26 mm x 30 mm
03.807.371	XRL Medium Graft Packing Tamp
03.807.374	XRL Medium Graft Packing Preparation
	Tamp
03.807.382	XRL Medium Implant Holder



and Graphic Cases—DJ1305

- Processing Non-sterile Synthes Implants-DJ1304



		Also Available	
(RL Medium In	nplant, Integrated, 0°, sterile	8205	XRL Preoperative Planner, medium
	Height	08.807.2125	XRL Medium Central Body, 73 mm–88 mm
8.807.2015	22 mm–25 mm		height, sterile
8.807.2025	24 mm–29 mm	08.807.2135	XRL Medium Central Body, 84 mm–99 mm
8.807.2035	28 mm–36 mm		height, sterile
		08.807.2145	XRL Medium Central Body,
(RL Medium C	entral Body, sterile		95 mm–110 mm height, sterile
	Height	08.807.2155	XRL Medium Central Body,
8.807.2045	22 mm–27 mm		106 mm–121 mm height, sterile
8.807.2055	25 mm–33 mm		
8.807.2065	29 mm–39 mm		
8.807.2075	33 mm–43 mm		
8.807.2085	37 mm–52 mm		
8.807.2095	44 mm–59 mm		
8.807.2105	51 mm–66 mm		
8.807.2115	62 mm–77 mm		
(RL Medium Er	ndplate, 21 mm round, sterile		
8.807.2215	0°		
8.807.2225	5°		
8.807.2235	10°		
8.807.2245	15°		
	ndplate, 21 mm x 24 mm, sterile		
8.807.2315	-10°		
8.807.2325	-5°		
8.807.2335	0°		
8.807.2345	5°		
8.807.2355	10°		
8.807.2365	15°		
	ndplates, 26 mm x 30 mm, sterile		
8.807.2415	-10°		
8.807.2425	-5°		
8.807.2435	0°		
8.807.2445	5°		
8.807.2455	10°		
8.807.2465	15°		

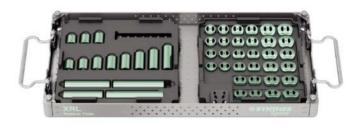
Graphic Case

60.807.032 Graphic Case, for XRL Medium Trials

Trials

XRL Medium Trial	-
	Height
03.807.501	22 mm–25 mm
03.807.502	24 mm–29 mm
03.807.503	28 mm–36 mm
XRL Medium Trial	3
	Height
03.807.504	22 mm–27 mm
03.807.505	25 mm–33 mm
03.807.506	29 mm–39 mm
03.807.507	33 mm–43 mm
03.807.508	37 mm–52 mm
03.807.509	44 mm–59 mm
03.807.510	51 mm–66 mm
03.807.511	62 mm–77 mm
VDL Madium Trial	Endulate 21 mm round
03.807.521	, Endplate, 21 mm round 0°
03.807.522	5°
03.807.523	10°
03.807.524	15°
03.807.524	15
XRL Medium Trial	Endplate, 21 mm x 24 mm
03.807.531	-10°
03.807.532	-5°
03.807.533	0°
03.807.534	5°
03.807.535	10°
03.807.536	15°
	Endplate, 26 mm x 30 mm
03.807.541	-10°
03.807.542	-5°
03.807.543	0°
03.807.544	5°
03.807.545	10°





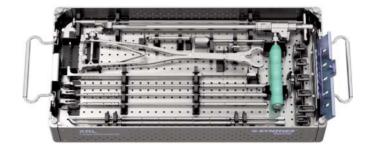
Also Available

03.807.512	XRL Medium Trial, Central Body,
	73 mm–88 mm height
03.807.513	XRL Medium Trial, Central Body,
	84 mm–99 mm height
03.807.514	XRL Medium Trial, Central Body,
	95 mm–110 mm height
03.807.515	XRL Medium Trial, Central Body,
	106 mm–121 mm height

XRL Large Implant and Instrument Set (01.807.030)

Cary Case and Graphic Case

Cary Case and Graphic Case		
60.807.027	Carry Case, for XRL Large Implants	
60.807.030	Graphic Case, for XRL Large Instrument	
Instruments		
03.661.010	Motal Tana Cauga	
	Metal Tape Gauge	
03.807.300	XRL Spreader	
03.807.330	XRL Large Shaft	
	XRL Large Spreader Top	
03.807.331	with 3 mm distraction	
03.807.332	with 5 mm distraction	
03.807.333	with 8 mm distraction	
03.807.334	with 10 mm distraction	
03.807.335	with 15 mm distraction	
03.807.356	with 5 mm distraction (Integrated)	
03.807.348	XRL Release Tool	
03.807.352	XRL Large Endplate Screwdriver Tip	
03.807.354	XRL Endplate Removal Tool	
03.807.367	XRL Large Endplate Footprint Trial,	
	27 mm round	
03.807.368	XRL Large Endplate Footprint Trial,	
	28 mm x 33 mm	
03.807.369	XRL Large Endplate Footprint Trial,	
	30 mm x 40 mm	
03.807.372	XRL Large Graft Packing Tamp	
03.807.375	XRL Large Graft Packing Preparation Tamp	
03.807.384	XRL Large Implant Holder	
08.807.300.025	XRL Large Endplate Screw (PEEK), sterile	



Implants

XRL Large Implant, Integrated, 0°, sterile Height 23 mm-26 mm 08.807.3015 08.807.3025 25 mm-30 mm 08.807.3035 29 mm-37 mm XRL Large Central Body, sterile Height 08.807.3045 22 mm-27 mm 25 mm-33 mm 08.807.3055 08.807.3065 29 mm-39 mm 08.807.3075 33 mm-43 mm 08.807.3085 37 mm-52 mm 08.807.3095 44 mm-59 mm 08.807.310S 51 mm-66 mm 08.807.3115 62 mm-77 mm XRL Large Endplates, 27 mm round, sterile 0° 08.807.3215 08.807.3225 5° 08.807.3235 10° 15° 08.807.324S 08.807.3255 20° XRL Large Endplates, 28 mm x 33 mm, sterile 0° 08.807.3315 5° 10° 15° 20° XRL Large Endplates, 30 mm x 40 mm, sterile 0° 5°

Also Available 0200

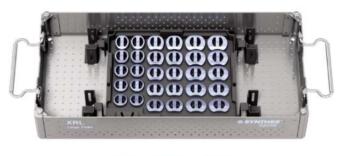
8206	XRL Preoperative Planner, Large
08.807.3125	XRL Large Central Body,
	73 mm–88 mm height, sterile
08.807.3135	XRL Large Central Body,
	84 mm–99 mm height, sterile
08.807.3145	XRL Large Central Body,
	95 mm–110 mm height, sterile
08.807.3155	XRL Large Central Body,
	106 mm–121 mm height, sterile

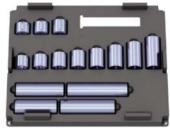
08.807.3325 08.807.3335 08.807.3345 08.807.3355

08.807.2415 08.807.2425 08.807.2435 10° 08.807.244S 15° 20° 08.807.2455

60.807.033 Graphic Case, for XRL Large Trials

Trials	
XRL Large Trial,	Integrated, 0°
	Height
03.807.601	23 mm–26 mm
03.807.602	25 mm–30 mm
03.807.603	29 mm–37 mm
XRL Large Trial,	-
	Height
03.807.604	22 mm–27 mm
03.807.605	
03.807.606	29 mm–39 mm
03.807.607	33 mm–43 mm
03.807.608	37 mm–52 mm
03.807.609	44 mm–59 mm
03.807.610	51 mm–66 mm
03.807.611	62 mm–77 mm
-	Endplate, 27 mm round
03.807.621	0°
03.807.622	5°
	10°
	15°
03.807.625	20°
XRL Large Trial.	Endplate, 28 mm x 33 mm
03.807.631	0°
03.807.632	5°
03.807.633	10°
03.807.634	15°
03.807.635	20°
XRL Large Trial,	Endplate, 30 mm x 40 mm
03.807.641	0°
03.807.642	5°
03.807.643	10°
03.807.644	15°
03.807.645	20°





Also Available

03.807.612	XRL Large Trial, Central Body,
	73 mm–88 mm height
03.807.613	XRL Large Trial, Central Body,
	84 mm–99 mm height
03.807.614	XRL Large Trial, Central Body,
	95 mm–110 mm height
03.807.615	XRL Large Trial, Central Body,
	106 mm–121 mm height



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