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ORTHOPAEDICS



EXACTA^{RS}

Femoral Stem

**PRODUCT
INFORMATION**

CE
0426

EXACTA^{RS}

12 implant sizes

STANDARD (ccd 132°)

LATERAL (ccd 132°) offset 6mm

Taper

12/14 (top angle 5°42'30") BIOLOX approved

Neck

Reduced section to optimize ROM. Mirror polished finishing to minimize wear on UHMWPE joint liner.

Stability

Horizontal grooves in the proximal portion for improved axial stability.

Bio-active coating

The **HaX-Pore®** double coating of porous Titanium and hydroxyapatite applied with plasma spray technique in the metaphyseal region guarantees an optimal primary fixation and encourages osteoblast colonization thus promoting biological integration.

Distal tip

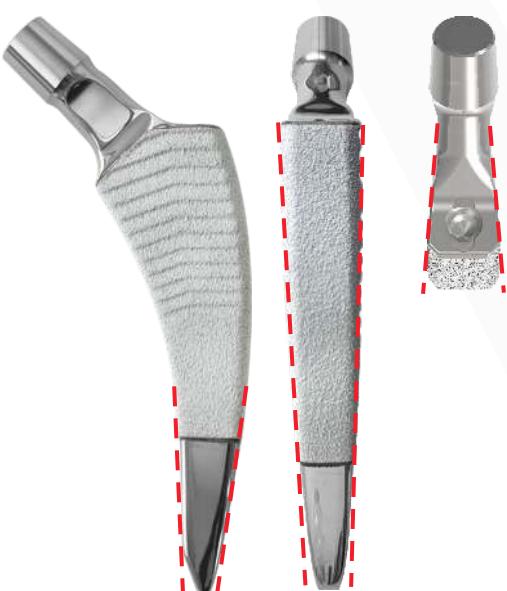
The tapered design and the glossy finishing avoid load concentration on the femoral cortex thus post-operative thigh pain.

Design

The **triple tapered** design (frontal, coronal and sagittal) supports axial and rotational stability favouring load distribution in the metaphyseal region.

The **progressive growth** of the neck proportional to the stem size allows for a better restoration of the natural anatomy.

The **anatomical curvature** and the **optimized dimensions** allow an easy introduction even with minimally invasive techniques or with anterior access, as well as a **limited bone sacrifice**.



EXACTA RS cementless femoral stem

Standard - CCD 132°				Lateral - CCD 132°			
Size	L mm	Offset mm	L2 mm	HaX-Pore®	X-Pore	HaX-Pore®	X-Pore
				reference	reference	reference	reference
1	90	35,0	26,3	13211	13251*	41,0	30,5
2	93	37,0	28,3	13212	13252*	43,0	32,5
3	96	37,5	28,3	13213	13253*	43,5	32,5
4	99	39,5	30,3	13214	13254*	45,5	34,5
5	102	40,0	30,3	13215	13255*	46,0	34,5
6	105	41,6	31,6	13216	13256*	48,6	36,6
7	108	42,2	31,6	13217	13257*	49,2	36,6
8	111	43,9	33,0	13218	13258*	50,9	37,9
9	114	44,5	33,0	13219	13259*	51,5	37,9
10	117	46,2	34,3	13220	13260*	53,2	39,3
11	121	47,9	34,3	13221	13261*	54,0	39,3
12	124	47,8	34,3	13222	13262*	54,7	39,3

Information**INTENDED PURPOSE:**

the EXACTA RS stems are intended for use in total or partial Hip Replacement procedures, combined with a femoral ball head (or a bi-articular head) and an acetabular cup. Indicated for primary hip arthroplasties in cases of serious joint degeneration, mainly due to arthrosis and post-traumatic degenerative factors. Device fixation is obtained by means of primary cementless press-fit stabilization.

MATERIALS:

Titanium Aluminium Niobium forged alloy (Ti6Al7Nb) ISO5832/11.

SURFACE FINISHING:

HaX-Pore®: double coating 300µm pure Titanium+50µm Hydroxyapatite $\text{Ca}_{10}(\text{OH})_2(\text{PO}_4)_6$ plasma sprayed

X-Pore: coating 300µm pure Titanium plasma sprayed

STERILIZATION:

Method: Irradiation (Beta or Gamma rays - nominal dose 25 kGy) or vaporized Hydrogen Peroxide (VH2O2).

Validity: 5 years (Beta) - 10 years (Gamma-VH2O2).

CLASSIFICATION:

Class III as reported in Directive 2005/50/CE (and related D.lgs 26 april 2007 n.65) concerning re-classification of Hip, Knee and Shoulder joint prostheses which modifies classification criteria of Annex IX of Directive 93/42/CEE and next integrations and amendments.



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