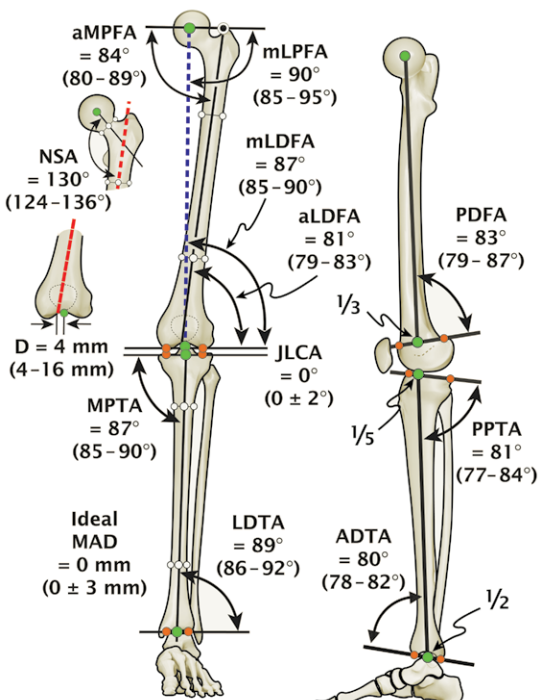
PRECICE®

ADJUSTABLE SOLUTIONS
FOR ORTHOPEDICS

Standard Measurements



PRECICE® Measurement Calculator

NAIL APPROACH	OVERALL LENGTH (mm)	DIAMETER (mm)	MAXIMUM DISTRACTION (mm)
ANTEGRADE TIBIA – 10°	155	8.5, 10.7	50
	160	8.5, 10.7	30
	180	8.5, 10.7	50
	195, 215, 230	8.5, 10.7, 12.5	50
	245, 275, 305, 335	8.5, 10.7, 12.5	80
	365	10.7, 12.5	80
ANTEGRADE FEMUR – TROCHANTERIC 10°	165	8.5	50
	170	8.5, 10.7, 12.5	30
	175	10.7	50
	190, 215, 230	8.5, 10.7, 12.5	50
	245, 275, 305, 335	8.5, 10.7, 12.5	80
	365	10.7, 12.5	80
ANTEGRADE FEMUR – PIRIFORMIS STRAIGHT	215, 230	8.5, 10.7, 12.5	50
	245, 275, 305, 335	8.5, 10.7, 12.5	80
	365	10.7, 12.5	80
RETROGRADE FEMUR – 10°	165	8.5	50
	170	8.5, 10.7, 12.5	30
	175	10.7	50
	190, 215, 230	8.5, 10.7, 12.5	50
	245, 275, 305, 335	8.5, 10.7, 12.5	80
	365	10.7, 12.5	80
RETROGRADE FEMUR – STRAIGHT	215, 230	8.5, 10.7, 12.5	50
	245, 275, 305, 335	8.5, 10.7, 12.5	80
	365	10.7, 12.5	80
UNIVERSAL FEMUR – STRAIGHT	150	8.5	50
	160	10.7	50
	170	8.5, 10.7, 12.5	30
	190	8.5, 10.7, 12.5	50

NuVasive
SPECIALIZED ORTHOPEDICS, INC.

For more information about this product,
please contact your local sales representative.

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CE 0297

The PRECICE Intramedullary Limb Lengthening (IMLL) System is composed of an implantable intramedullary nail, locking screws, reusable instruments, and a hand-held External Remote Controller (ERC). The PRECICE nail is a sterile single use device that is surgically implanted using the instruments and locking screws. The ERC is used daily after implantation to non-invasively lengthen or shorten the implant to a prescribed length. The PRECICE System is intended for limb lengthening of the femur and tibia. Contraindications include infection or pathologic conditions of bone such as osteopenia which would impair the ability to securely fix the device, metal allergies and sensitivities, patients whose distance from the surface of the treated limb to the intramedullary canal is greater than 51 mm for the 10.7 and 12.5 mm diameter implants or greater than 38 mm for the 8.5 mm diameter implant, patients with an irregular bone diameter that would prevent insertion of the PRECICE nail, patients in which the PRECICE nail would cross joint spaces or open epiphyseal growth plates, patients in which there is an obliterated medullary canal or other conditions that tend to retard healing such as blood supply limitations, peripheral vascular disease or evidence of inadequate vascularity, patients unwilling or incapable of following postoperative care instructions, patients weighing in excess of 114 Kg for the 10.7 and 12.5 mm diameter implants for models H, J, K, and U or weighing in excess of 57 Kg for the 8.5 and 10.7 mm diameter implants for models N, M ,P, and Q. The implantable device is only to be used by a trained licensed physician. Please refer to the PRECICE IMLL System instructions for use for complete Important Safety Information.
Caution: Federal law restricts this device to sale by or on the order of a physician.

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