OSTEOREMEDIES®
ADVANCED MEDICAL TECHNOLOGIES

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SURGEON INFORMATION

REMEDY® RADEL® KNEE SPACER TRIAL

The REMEDY® Radel® Knee Spacer Trial is a PPSU device composed of two independent articulating components (REMEDY® Radel® Femoral Trial and the REMEDY® Radel® Tibial Trial – both available in three sizes) that must be combined to fit the anatomy of the patient. A REMEDY® Radel® Tibial Insert Trial is also available and comes in 3 different sizes as well.

Each REMEDY® Radel® Femoral Trial is matchable to each REMEDY® Radel® Tibial Trial, while each REMEDY® Radel® Tibial Insert Trial is matchable only with its corresponding REMEDY® Radel® Tibial Trial.

The correct combination of the REMEDY® Radel® Knee Spacer Trial enables the surgeon to select the appropriately sized REMEDY® Knee Spacer to be implanted.

Step 1: Choose the REMEDY® Radel® Femoral Trial and REMEDY® Radel® Tibial Trial based on the dimension of the removed prosthesis. In case of a large tibial defect, include a REMEDY® Radel® Tibial Insert Trial.

Step 2: Test the device in the patient to determine if the prosthesis is anatomically correct and stable.

Step 3: Once the device has been tested and verified, remove it from the patient, and use it as a reference to prepare the REMEDY® Knee Spacer device which will be implanted.

CLEANING & STERILIZATION

REMEDY® Radel® Knee Spacer Trial components are nonsterile.

Before each use the device must be cleaned and sterilized.

The REMEDY® Radel® Knee Spacer Trials components must be cleaned by the means of most commonly used disinfectants (e.g. bleach solutions, isopropyl alcohol, hydrogen peroxide, phenols) and detergents. It is recommended to avoid the contact of the material with chemicals as esters, aromatic hydrocarbons, chlorinated hydrocarbons and ketones.

RECOMMENDED STERILIZATION CYCLE:

TEMPERATURE	PRESSURE	MINIMUM TIME
121° C	1 atm	15 Min
134° C	2 atm	4 Min

WARNING: The REMEDY® Radel® Knee Spacer Trial must be used only to determine the anatomically correct size of the REMEDY® Knee Spacer to be implanted. The trial device or the single component trial must not be implanted. Do not use the device if it appears damaged (deformation, loss of the information marked on the device, etc.). The device needs to be replaced if it appears damaged after a visual inspection.

CAUTION

Federal law (USA) restricts this device to sale by or on the order of a physician.
Keep out of reach of children.

DESCRIPTION	REF CODE
REMEDY® Radel® Femoral Trial Small	RRTFMSM
REMEDY® Radel® Femoral Trial Medium	RRTFMMD
REMEDY® Radel® Femoral Trial Large	RRTFMLG
REMEDY® Radel® Tibial Trial Small	RRTTBSM
REMEDY® Radel® Tibial Trial Medium	RRTTBMD
REMEDY® Radel® Tibial Trial Large	RRTTBLG
REMEDY® Radel® Tibial Wedge Trial Small	RRTINSM
REMEDY® Radel® Tibial Wedge Trial Medium	RRTINMD
REMEDY® Radel® Tibial Wedge Trial Large	RRTINLG
REMEDY® Radel® Knee Spacer Trial (KIT)	RRKITKN

Symbols: