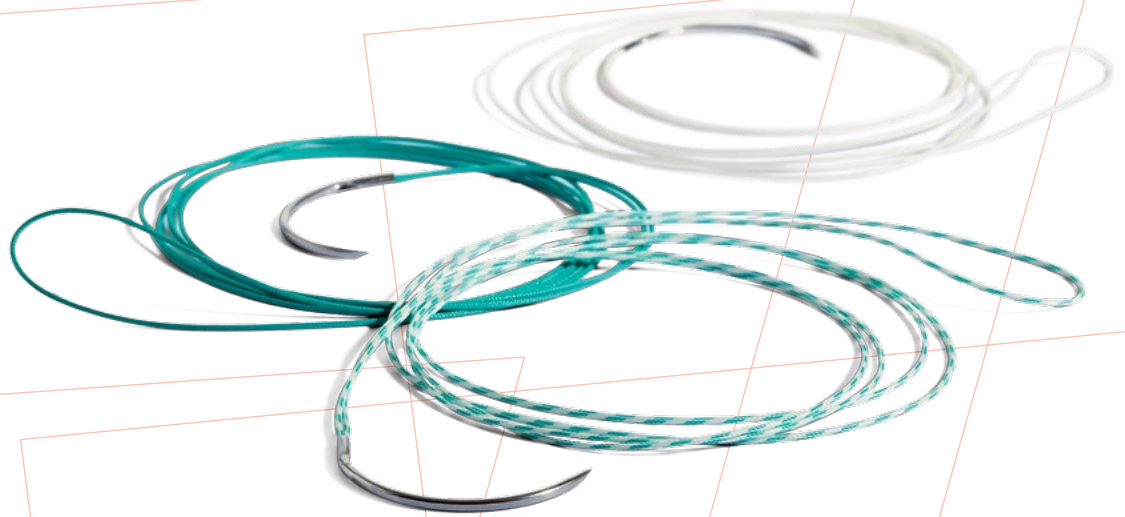


TORNIER
NICELOOP™

Nonabsorbable Braided Suture
for Reliable Bone Fixation



NICELOOP™

The NICELOOP™ is the dedicated solution to perform the Nice Knot, a sliding and self stabilizing knot, invented by Pascal Boileau M.D, PhD, in Nice (France).

For more information about the Nice Knot, Please refer to the WhitePaper "The Doubled Suture Nice Knot" (CAW-1041 rev A).

NICELOOP™ Advantages:

- Strength: the NICELOOP is tied as a doubled-over suture, increasing significantly the knot strength.
- Adjustability: the tightening process of the Nice Knot can be stopped and resumed at any stage, as the knot is self stabilized (preventing from slippage).
- Versatility: the NICELOOP can be used in a variety of indications:
 - Tuberosities fixation during hemi or reversed arthroplasty for fracture
 - Cerclage for humerotomy or femorotomy
 - Fixation of bone fragments (i.e Butterfly)
- Simplicity: by combining a simple initial knot with 3 half hitches, the Nice Knot provides knot security without an excessive amount of complexity and bulkiness.

NICELOOP™ Properties:

- Available in 3 colors: Solid White, Solid Green, White/Green
- 60cm (24 inches) long loop (doubled-over)
- Braided PTFE Impregnated Polyester Fiber Nonabsorbable Surgical Suture
- Needle: KAC-25, 1/2 Circle, K-Point (49mm)

NICELOOP™ Ordering Information

Reference	Description	Size	Needle	Packaging
SMSL50103	Green NonAbsorbable Suture Loop	#5	KAC-25	Box of 12
SMSL50203	White NonAbsorbable Suture Loop	#5	KAC-25	Box of 12
SMSL50303	White/Green NonAbsorbable Suture Loop	#5	KAC-25	Box of 12
SMSL50101	Green NonAbsorbable Suture Loop	#5	KAC-25	Single
SMSL50201	White NonAbsorbable Suture Loop	#5	KAC-25	Single
SMSL50301	White/Green NonAbsorbable Suture Loop	#5	KAC-25	Single

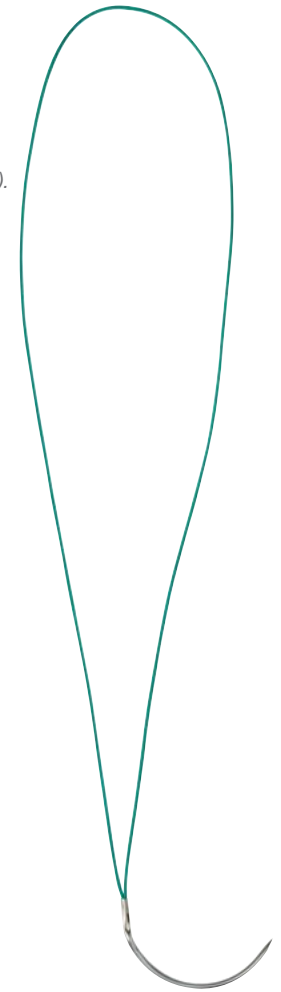
Indications:

The NICELOOP™ Suture is indicated for use in general soft tissue approximation and/or ligation, including use in cardiovascular, ophthalmic, orthopedic and neurological procedures.

Contraindications:

None.

Proper surgical procedures and techniques are the responsibility of the medical professional. This material is furnished for information purposes only. Each surgeon must evaluate the appropriateness of the material based on his or her personal medical training and experience. Prior to use of any Tornier implant system, the surgeon should refer to the product package insert for complete warnings, precautions, indications, contraindications and adverse effects. Package inserts are also available by contacting Wright. Contact information can be found in this document and the package insert.



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