
Instructions for Use Cable Implants

This instruction for use is not intended for distribution in the USA.

Instructions for Use

Cable Implants

Please read these instructions for use, the Synthes "Important Information" and the corresponding Surgical Technique Guide (www.synthes.com/lit) carefully before use. Ensure that you are familiar with the appropriate surgical technique.

Cable Implants consist of an assembly of various devices such as cables, cerclage pins and eyes as well as reattachment devices. The Cable Implants are single packed, and available sterile and/or non-sterile.

Important note for medical professionals and OR staff: These instructions for use do not include all of the information necessary for selection and use of a device. Please see full labeling for all necessary information (corresponding Surgical Technique Guide, Important Information and device-specific label).

Material(s)

Material(s):	Standard(s):
Stainless steel	ISO 5832-1
Stainless steel (CoCr)	ISO 5832-5
TiCP	ISO 5832-2
Titanium alloy	ISO 5832-11

Intended use

Cable Implants are intended for fixation or stabilization of bones in various anatomical regions by using standard cerclage or tension band technique.

Indications

For specific indications for Cable Implants it is mandatory to consult the corresponding Surgical Technique Guide (www.synthes.com/lit) of the product system being used.

Contraindications

For specific contraindications for Cable Implants it is mandatory to consult the corresponding Surgical Technique Guide (www.synthes.com/lit) of the product system being used.

Side effects

As with all major surgical procedures, risks, side effects and adverse events can occur. While many possible reactions may occur, some of the most common include:

Problems resulting from anesthesia and patient positioning (e.g. nausea, vomiting, dental injuries, neurological impairments, etc.), thrombosis, embolism, infection, excessive bleeding, iatrogenic neural and vascular injury, damage to soft tissues incl. swelling, abnormal scar formation, functional impairment of the musculoskeletal system, Sudeck's disease, allergy / hypersensitivity reactions, and side effects associated with hardware prominence, malunion, non-union.


Sterile device

STERILE R Sterilized using irradiation

Store implants in their original protective packaging, and do not remove them from the packaging until immediately before use.

Prior to use, check the product expiration date and verify the integrity of the sterile packaging. Do not use, if the package is damaged.

Single-use device

 Do not re-use

Products intended for single-use must not be re-used.

Re-use or reprocessing (e.g. cleaning and resterilization) may compromise the structural integrity of the device and / or lead to device failure which may result in patient injury, illness or death.

Furthermore, reuse or reprocessing of single-use devices may create a risk of contamination e.g. due to the transmission of infectious material from one patient to another. This could result in injury or death of the patient or user.

Contaminated implants must not be reprocessed. Any Synthes implant that has been contaminated by blood, tissue, and/or bodily fluids/matter should never be used again and should be handled according to hospital protocol. Even though they may appear undamaged, the implants may have small defects and internal stress patterns that may cause material fatigue.

Precautions

For general precautions consult "Important Information".

For application specific warnings related to Cable Implants it is mandatory to consult the corresponding Surgical Technique Guide (www.synthes.com/lit) of the product system being used.

Warnings

For general warnings consult "Important Information".

For application specific warnings related to Cable Implants it is mandatory to consult the corresponding Surgical Technique Guide (www.synthes.com/lit) of the product system being used.

Combination of medical devices

Synthes has not tested compatibility with devices provided by other manufacturers and assumes no liability in such instances.

Magnetic Resonance environment

CAUTION:

Unless stated otherwise, devices have not been evaluated for safety and compatibility within the MR environment. Please note that there are potential hazards which include but are not limited to:

- Heating or migration of the device
- Artifacts on MR images

Treatment before device is used


Synthes products supplied in a non-sterile condition must be cleaned and steam-sterilized prior to surgical use. Prior to cleaning, remove all original packaging. Prior to steam-sterilization, place the product in an approved wrap or container. Follow the cleaning and sterilization instruction given by the Synthes "Important Information".

Processing/reprocessing of the device

Detailed instructions for processing of implants and reprocessing of reusable devices, instrument trays and cases are described in the Synthes brochure "Important Information". Assembly and disassembly instructions of instruments "Dismantling multi-part instruments" can be downloaded from <http://www.synthes.com/reprocessing>

Additional device-specific information


 Reference Number

 Lot or batch number


 Manufacturer


 Manufacturing date

 Expiration date

 Do not use when packaging is damaged

 Notified body

 Caution, see instructions for use

 Consult instructions for use

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Synthes GmbH
Eimattstrasse 3
4436 Oberdorf
Switzerland
www.synthes.com