

INTEGRA Intra Uterine Insemination Catheter

Description:

CODE	Description
IG-IUI	INTEGRA Intra Uterine Insemination Catheter

Configuration:

See brochure

Indications for use:

Allwin Intra Uterine Insemination Catheters are intended for the introduction of washed spermatozoa into the uterine cavity.

Contraindications:

- Cervical infection
- Recent pelvic inflammation.
- Sexually transmitted disease
- Pregnancy
- Confirmed or suspected intrauterine device
- Uterine perforation.

Complications:

- Uterine wall perforation
- Bleeding

Caution:

- Federal (USA) law restricts this device to sale by or on the order of a physician.
- To reduce the risk of perforation, if any resistance is felt while inserting the catheter, do not force the catheter against the resistance.

Warning / Precaution:

- Do not use if package is opened or damaged.
- Not intended for IVF, GIFT or other Intra Fallopian tube procedure.
- Always use washed spermatozoa when performing intrauterine artificial insemination. The introduction of unwashed spermatozoa into the uterus may result in severe adverse reactions, including anaphylactic shock. Please refer to published medical literature for methods of preparing spermatozoa for intrauterine artificial insemination before performing the procedure.
- Dispose in accordance with recognized medical practice and under observance of applicable law and regulation.
- For single use only.
- Do not re-sterilize.
- Only experienced surgeon should use this device.
- Storage temperature between 5°C to 30°C.

Instruction for Use:

1. Before to the procedure, check the condition of the cervix prior to the insertion of the catheter, to ascertain the depth of the uterus and any uterine anteflexion or retroflexion present in individual patients.
2. Place the patient in the dorsal lithotomy position with her feet in stirrups.
3. Position a vaginal speculum within the vagina so the external cervical os can be visualized during catheter insertion.
4. Cleanse the cervix with an appropriate cleansing solution.
5. If necessary, gently pre-shape the catheter using the malleable stainless-steel inner cannula already integrated within the device.
6. Adjust the silicone stopper to control insertion depth.
7. Attach a syringe preloaded with washed sperm solution securely to the Luer lock connector.
8. Slowly press the plunger of the syringe to introduce the sperm suspension into the uterine cavity through the side hole near the closed distal tip.
9. Following injection, carefully withdraw the catheter and syringe, and discard all used materials in accordance with recognized medical practice and applicable regulations.

NOTE-1 Human Sperm Survival Assay (HSSA) > 80% of control motility at 24 hours

NOTE-2 Endotoxin Testing: ≤ 20 EU/device

Do not reuse, reprocess & resterilize:

Reuse, Reprocessing & reesterilization may compromise the structural integrity, can also create risk of contamination & may not give desired result or create complications, infections which may result in injury, illness or death.

Limited Express Warranty:

The limited express warranty as set forth herein is exclusive and in lieu of all warranties of merchantability and fitness for use, remedies, obligations and liabilities for consequential damages. The products are being sold only for the purpose described herein and such limited express warranty runs only to the original user. In no event shall ALLWIN be liable for any breach of warranty in any amount exceeding the purchase price of the product. ALLWIN reserves the right to make design changes to products without liability to incorporate said changes in ALLWIN products previously designed or sold.

Catalogue Number
 Batch Code
 Date of Manufacture
 Use By
 Do not re-use
 Do not Re-Sterilize
 Rx only
 Caution: Federal Law restricts this device to sale by or on the order of a physician or a practitioner trained in its use.

Consult Instruction for use
 Do not use if Packing is damaged
 Caution
 Keep out of sunlight
 Keep Dry
 MD Medical Device
 STERILE EO Sterilized using ethylene oxide
 Not made with natural rubber latex
 ^{°C} Temperature Limit

EU REP
 CMC Medical Devices & Drugs S. L.
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Single Sterile Barrier system with protective packaging inside

CE
 0123

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