

**IFU of Maisway Click™**  
**Three Way Stopcock with click at 45°**  
**Manufactured by Mais India Medical Devices Pvt. Ltd.**

**Intended Use:**

The Three Way Stopcock is intended to control and redirect the flow of fluids or drugs in IV therapy, blood transfusion, infusion pumps, or pressure monitoring systems.

**Materials Used:**

Polycarbonate (PC), High-density Polyethylene (HDPE).

**Indications:**

- For use in conjunction with IV catheters, syringes, or infusion sets.
- To provide multi-directional flow path control in clinical settings.

**Contraindications:**

- Do not use for delivery of high-viscosity fluids unless verified for compatibility.
- Not intended for reuse, implantation, or use with blood products if not explicitly validated.

**Instructions for use:**

- Sanitize your hands using an alcohol-based hand cleanser.
- Ensure I.V. line or connecting device (I.V. Cannula, Infusion sets, Extension lines, Syringes, Pressure transducers) are affixed properly and are in function.
- Visually inspect the product packaging to ensure integrity. Do not use if packaging is damaged.
- Open the packaging using aseptic technique.
- Connect the male luer lock end of the stopcock to the infusion line or any of connecting device.
- Use the rotating tap to select desired flow direction.
- The 45° click provides tactile feedback and ensures accurate positioning.
- Ensure leak-free and bubble-free fluid administration.
- Monitor the patient and flow throughout the procedure.
- After use, disconnect and discard as per hospital biohazard protocols.

**Medical conditions:**

- Read the "Instructions for Use" carefully before using the product.
- The use of the product is restricted to a qualified doctor or a paramedic staff.

**Warnings:**

- Single Use Only: Do not reuse or re-sterilize.
- Sterile Unless Package is Damaged: Inspect packaging before use.
- Ensure secure luer connections before use to prevent leakage.
- Avoid using if cracks, leaks, or discoloration are observed.
- Use only under the supervision of qualified healthcare professionals.

**Cautions:**

- Do not use if the product or its packaging is damaged.
- Do not reinsert any syringe or set forcefully into ports.
- Discard the device if it has been partially or fully disconnected from sterile systems.
- Use within the specified expiry date.

**Target Age Group:** For All age groups

**Duration of Contact of Device:** Up to 72 Hours for best performance.

**Storage conditions:** Store at a temperature between +10°C and +40°C, in a dry place and away from sunlight.

**Device Life:** 05 Years

**Explanation of symbols used:**

Symbols	Title of the Symbol	Symbols	Title of the Symbol
	Consult instructions for use (Indicates the need for the user to consult the instructions for use)		Latex free (Indicates a medical device that is Latex free)
	Keep away from sunlight (Indicates a medical device that needs protection from light sources)		Date of manufacture (Indicates the date when the medical device was manufactured)
	Reference number (Indicates the manufacturer's catalogue number so that the medical device can be identified)		Keep dry (Indicates a medical device that needs to be protected from moisture)
	Sterilized using ethylene oxide (Indicates a medical device that has been sterilized using ethylene oxide)		Use-by date (Indicates the date after which the medical device is not to be used)

Symbols	Title of the Symbol	Symbols	Title of the Symbol
	Single sterile barrier system (Indicates a single sterile barrier system)		Country of manufacture (To identify the country of manufacture of products)
	Do not re-sterilize (Indicates a medical device that is not to be re-sterilized)		Batch Number (Indicates the manufacturer's batch number so that the batch or lot can be identified)
	Do not re-use (Indicates a medical device that is intended for one single use only)		CE Mark (CE Marking with Notified Body Number)
	Non-pyrogenic (Indicates a medical device that is non-pyrogenic)		Medical device (Indicates the item is a medical device)
	Do not use if packaging is damaged and consult instructions for use. (Indicates a medical device that should not be used if its packaging has been damaged or opened, and that the user should consult the instructions for use for additional information)		Authorized representative in the European Union / European Community (Indicates the authorized representative in the European Union / European Community) OBELIS S.A. Boulevard Général Wahis 53, 1030 Brussels, Belgium Contact Number: +32(2)73 25 954 Email ID: regulatory@obelis.net Website: www.obelis.net
	Caution: (To indicate that caution is necessary when operating the device or control close to where the symbol is placed, or to indicate that the current situation needs operator awareness or operator action in order to avoid undesirable consequences.)		Manufacturer (Indicates the medical device manufacturer) Manufacturing Unit: 525P, Sector-37, Pace City II, Gurgaon, Haryana-122001, India. Contact Details: +91 8527589990 Fax No.: 0124 404 7533 Email ID: info@maisindia.com Website: www.maisindia.com
	Unique Device Identifier (Indicates a carrier that contains Unique Device Identifier information)		Temperature limit (Indicates the temperature limits to which the medical device can be safely exposed.)
	This way up		Fragile, handle with care (Indicates a medical device that can be broken or damaged if not handled carefully)

**Manufactured by:**  
**Mais India Medical Devices Pvt. Ltd.**  
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 Gurgaon, Haryana-122001 (INDIA)  
 Mfg. Lic. No.: MFG/MD/2019/000037

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REV BY	Q.A		<b>TITLE : IFU of Maisway Click - Three Way Stopcock</b>		SHEET No. 1/1	APPROVED BY:
	Q.C					
	Pro.					
CHD BY	Mkt.		<b>For Party Code 000</b>		<b>DRW. No.:</b> I02811-IFU-000	
DGN BY	Gopal Singh	DATE: 28.02.2026	<b>COLOR OF PRINT:</b> Black		<b>EDITION:</b> 02, <b>REV.:</b> 01	
SCALE-N.T.S		TOLERANCE : ±2 mm	<b>REVISION DETAILS:</b>		<b>Size:-</b> 225x130mm	
<b>MAIS INDIA MEDICAL DEVICES PVT. LTD.</b> 525-P SECTOR-37, PACE CITY II, GURGAON, HARYANA-122001 (INDIA)			<b>Rev. 01:</b> The humidity limitation symbol has been removed and EC-REP has been changed to EU-REP.			