

General instructions for use

Intended use

MHP sterile instruments are intended for single use and are designed for performing [e.g. surgical, diagnostic or therapeutic procedures].
Use only by qualified personnel in a medical environment.

Areas of application

- See areas of application in the technical data sheet
- Suitable for sterile applications
- Not suitable for reuse or for procedures for which it is not intended

Warnings and precautions

- **Disposable product:** Dispose of properly immediately after use.
Do not reuse.
- **Sterility:** Only use the product if the packaging is intact and the sterilisation indicator is correct.
- **Storage:** Store at room temperature, in a dry place away from light. Do not store at temperatures below 0 °C or above 40 °C.
- **Check before use:** Inspect the instrument for visible damage or material defects. **Defective products must not be used.**
- **Do not damage the packaging:** Only open the packaging immediately before use under aseptic conditions.

Preparation:

- Open the sterile packaging in accordance with aseptic rules.
- Check the instrument for integrity and cleanliness.

Application:

- Use the instrument in accordance with specific surgical requirements and in compliance with medical standards.
- Avoid applying improper pressure or excessive strain to the instrument in order to prevent damage or breakage.

After use:

- The instrument must not be cleaned, sterilised or reused after use.
- Dispose of the instrument in accordance with local regulations for medical waste.

Disposal

The product must be disposed of in accordance with regulations for disposable medical waste. Ensure that it is properly separated from other waste to avoid risks to the environment and personnel.

Disclaimer

The manufacturer/importer accepts no liability for damage or injury resulting from improper use or reuse of the instrument.

Note on reporting incidents:

Serious incidents involving this medical device must be reported to the manufacturer and importer, as well as to the competent authority of the Member State in which the user is located.

Contact for reports:**Manufacturer:**

RB Medikal Instruments
S.I.E , Fateh Ghar, Near Umer Town,
51310 Sialkot, PK

Notified body: SGS

Belgium NV Notified
Body n. 1639
Noorderlaan 87
2030 Antwerp, Belgium

**EU authorised representative**

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Importer

**MHP-Handelsgesellschaft Deutschland
mbH**

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