HEINE QUALITY
MADE IN GERMANY

Hygienic reprocessing

HEINE GAMMA® G5, G7, GP, XXL LF

General warning and safety information

WARNING! This symbol draws attention to a potentially dangerous situation. Non-observance can result in moderate to major injuries.

NOTE! This symbol indicates valuable advice in terms of set up, operation or maintenance, as applicable. Notes are important, but not related to hazardous situations.

	Instructions on hygienic reprocessing must be adhered to, based on national standards, laws and guidelines. They must be implemented in the hospital / practice internal rules and guidelines.
	In the event of suspected contamination, carry out hygienic reprocessing of the instrument.
	The device must not be immersed in liquids.
	The described reprocessing measures do not replace the specific rules applicable for your institution/ department.
	HEINE Optotechnik only approves the agents and procedures listed in this instruction.
	Hygienic reprocessing is to be carried out by persons with adequate hygienic expertise.
	Please consider the instructions of the manufacturer for the applied reprocessing media.
	Avoid liquid entry into the device.
	The cuffs have to be reprocessed after each use.
	The described reprocessing procedures are represented alongside the corresponding material compatibilities. Reprocessing must be carried out in accordance with an approved processing procedure. HEINE Optotechnik GmbH & Co. KG cannot guarantee the sterility and disinfection of these procedures. This has to be validated by the user e. g. Hospital or the manufacturers of the reprocessing equipment.
	For important details regarding the processing procedures, please refer to the FAQs for Hygienic Reprocessing on our Website.

HEINE Optotechnik GmbH & Co. KG • Kientalstr. 7 • 82211 Herrsching • Germany
Tel. +49(0)8152/38-0 • Fax +49(0)8152/38-202 • E-Mail: info@heine.com • **www.heine.com**

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Wipe disinfection

1. Preparation

In the event of suspected contamination, carry out hygienic reprocessing of the instrument.

2. Manual cleaning and disinfection

Pay attention that all surfaces are completely moistened for the complete exposure time specified by the disinfectant manufacturer. If necessary, increase the number of wiping procedures and/or the number of wipes.

Equipment

- Cleaning agent, if necessary: enzymatic (e. g. neodisher® MediClean)
- Disinfectant for HEINE GAMMA® sphygmomanometers:
  quarternary ammonium compounds (e. g. Cleanisept® Wipes , Mikrobac® Tissues or Sani-Cloth® AF3)
- Disinfectant for HEINE GAMMA® blood-pressure cuffs:
  alcoholic (e. g. Incides® N),
  quarternary ammonium compounds (e. g. Cleanisept® Wipes , Mikrobac® Tissues or Sani-Cloth® AF3) or
  alcoholic + quarternary ammonium compounds (e. g. Super Sani-Cloth®)

Implementation

- For heavier soiling, you can first clean with a wipe soaked with cleaning agent before disinfecting with a disinfectant wipe.
- Clean and disinfect the device manually.
- The cuff sleeve can be hand-washed at max. 30 °C after removing the inflatable bladder.
- Pay particular attention to difficult to access areas.
- Thoroughly wipe the touched areas.
- For removing the disinfectant and drying afterwards, follow the instructions provided by the disinfectant manufacturer.

3. Inspection and function testing

- Check the device for any visible contaminants or abrasions. Reprocess again if necessary. Dispose if the contaminants cannot be removed.

4. Storage

Store it in such a way that it is protected from recontamination, dust and moisture.

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