

**IMPORTANT**  
Before using these products, please read the following information thoroughly!

**WARNINGS** 

All instruments must be cleaned, disinfected and sterilized prior to each use.

All reusable Ackermann products are shipped in nonsterile condition. Before the first use transport packaging, coarse dust/pieces of paper/packaging remains need to be removed and each product processed and sterilized according to these instructions. All products have been pre-cleaned to an extent which allows for processing and sterilization by use of the equipment stated here. This only applies to a processing method according to these instructions within a system that has been configured and validated in compliance with ISO 17665 and in which all cleaning/disinfecting devices comply with ISO 15883.


**Thorough cleaning and disinfection is crucial for an effective sterilization!**  
Strong cleaning agents may cause fading of markings.

**LIMITATIONS ON REPROCESSING** 

Instruments have been validated for 50 cycles, based on an average treatment. Products that have been marked as single-use (acc. to EN 980 2008-11/figure 5.2) MUST NOT be reprocessed! With such products materials are being used that are NOT reprocessable under normal conditions or do not withstand more than one sterilization process and, therefore, may break during surgery if reprocessed! (this applies to practically all single-use products featuring plastic components).

**INTENDED USE**

The Ackermann urological sheaths are designed for diagnostic and operative urology surgeries to be used as operation/ irrigation/ aspiration channel, during the obturators are used through the sheaths to access to the urological tract.

**CONTRAINDICATIONS** 

Not intended for use with patients that have allergic reactions to Ni- ; CR- steels or to brass or aluminium.

**INSTRUCTIONS (acc. to ISO 17664)**  
**PLACE OF USE (immediately after use)** 

IMMEDIATELY after each use (within no more than 10 min. or before drying of contaminants) the instruments need to get disassembled and impurities removed under running water, using a soft brush or cloth which are being used solely for this purpose. NEVER use a metal brush, steel wool or other cleaning devices containing metal in order to avoid the imminent risk of corrosion. Rinse under cold, running water until all visible impurities and contaminants have been successfully removed.

**STORAGE AND TRANSPORT** 

Place instruments in a container; make sure that associated parts are being stored together. Keep the inside of the container moist/wet (no contaminants may dry). Reprocess all instruments soonest possible.

**PREPARATION FOR CLEANING** 

Dismantable products are ALWAYS to be disassembled as much as possible (see IFU). Soak instruments in cold water for at least 5 min. and clean them, using a soft brush or cloth which are being used solely for this purpose. NEVER use a metal brush, steel wool or other cleaning devices containing metal in order to avoid the imminent risk of corrosion. Afterwards, wash down the entire surface of the instrument for 10s. by use of a cleaning gun (min. continuous pressure of 4 bar); articulate moveable parts constantly during the preliminary cleaning. Instruments featuring lumina and/or LuerLock flush channels are to be rinsed for an additional 10s. after visibly clear water has emerged from the ports. Place the instruments in an ultrasonic bath for 10min. (35-

40kHz for min. 5min. or longer acc. to specifications). Prior to switching on the ultrasonics make sure that all lumina, sheaths, etc. are filled with cleaning fluid! Note that the preliminary cleaning – even at the use of a disinfectant – is only intended as a preparatory step and DOES NOT replace the actual disinfection!

**MECHANICAL CLEANING** 

Associated parts are to be stored together in order to facilitate a subsequent allocation. Make sure that multiple instruments do not touch; especially different materials such as titanium, brass, aluminum, stainless steel, etc. need to be cleaned separately in order to avoid formation of a rust film. Composite instruments particularly stainless steel combined with ceramics) need to be placed with sufficient distance to other products so they do not break due to the pressure of different thermal expansions. Instruments have been tested with the following devices:

- Washer-Disinfector G 7735 CD (Miele):
- 1. washing cycle: alkaline program (No 105)
- 2. washing cycle: enzymology program (No 105)
- Washer-Disinfector G 7836 CD (Miele)
- 1. two component alkaline/enzymatic program
- 2. OxiVario
- Washer-Disinfector Niagara SI PCF (Medisafe) (RECOMMENDED)
- 1. Cleaning process with pulsed ultrasonic irrigation
- 2. Cleaning process without pulsed ultrasonic irrigation

The water which is to be used needs to be sterile or nearly sterile (<10 microbes/ml) and low in endotoxins (< 0.25 units/ml). The air which is being used for drying needs to be cleaned by means of micro filters which Processing and Sterilization Instructions of Medical Devices (acc. to ISO 17664) are regularly checked and maintained. A maintenance schedule has to be documented.

**DISINFECTION**

Place the instruments into the disinfecting bath (Caution: products need to be fully immersed; at least 1cm below the liquid surface). Multiple instruments may not touch; especially different materials such as titanium, brass, aluminum, stainless steel, etc. need to be disinfected separately in order to avoid formation of a rust film. Composite instruments (particularly stainless steel combined with ceramics) need to be placed with sufficient distance to other products so they do not break due to the pressure of different thermal expansions. Rinse all the lumina of the instrument at least five times using a sterile syringe (min 50ml) and disinfectant.

**AFTER DISINFECTION**

Remove products and rinse for at least 5 min. under running water until all disinfectant is removed from the instruments (the water which is to be used needs to be sterile or nearly sterile with <10 microbes/ml and low in endotoxins with < 0.25 units/ml). Constantly articulate moveable parts. Rinse all the lumina of the instrument with water at least five times using a sterile syringe (min 50ml). Disinfectants that have been successfully tested are:

1. Alkaline, Neodisher FA, pH 12.2, Dr. Weigert
2. Enzymatic, deconex 23 Neutrazym, pH 9.7, Borer
3. 2-Component Alkaline/Enzymatic, deconex TWIN PH, pH 10.9, deconex TWINZYME, pH 7, Borer
4. 2-Component Alkaline, Sekumatic FR, pH 12.1; Secumatic OxiVario. PH 7.8; Neutralizer: Sekumatic FNZ, pH 2.2, Ecolab
5. Enzymatic; M20029 3E-Zyme Scope Plus, pH 6.1, Medisafe
6. Enzymatic + Ultrasound, M20029 3E-Zyme Scope Plus, pH 6.1, Medisafe

**DRYING**

After cleaning and disinfection place the instruments into suitable containers. Make sure that there is NO residue of the disinfectant. When drying as part of the cleaning/disinfection cycle is completed make sure that a temperature of 150°C/300°F is not exceeded. All operations need to take place in a clean, monitored environment!

**MAINTENANCE** 

Apply a small amount of high-grade surgical lubricant on all joints or other moveable parts which are supposed to move smoothly. Sort out all blunt or damaged instruments. Clearly damaged instruments (cracks on the insulation, breakage, strongly bleached polymer handles or coatings) are NOT to be reused but repaired or disposed of.

**TESTING AND INSPECTION**

Jointed instruments are to be tested for ease of movement (avoid too much backlash). The functionality of ratchet mechanisms needs to be checked. All instruments: visually check for damage and wear. Blades should be even and without notches. Long and narrow instruments (especially jointed instruments) should be particularly checked for damages. If instruments are part of a larger set they are to be checked together with all associated components.

**PACKAGING**

Individually: a standardized packaging material may be used. The size of each bag needs to match the individual instrument so that there is no tension applied on the sealing. Sets: sort instruments into designated trays or place on multi-purpose sterilization trays. Blades need to be protected; the weight of each tray may not exceed 8kg (18lbs). For the trays an adequate packaging procedure is to be used.

**STERILIZATION** 

All products have been pre-cleaned to an extent which allows for processing and sterilization by use of the equipment stated here. This only applies to a processing method according to these instructions within a system that has been configured and validated in compliance with ISO 17665 and in which all cleaning/disinfecting devices comply with ISO 15883. With the result of the sterilization process greatly depending on the equipment that is being used a sterilization validation acc. to ISO 17665 MUST be performed at the place of use prior to the first application. All products MAY be used only if the result of this validation is positive.

For the sterilization of medical devices various methods can be applied. Regarding products manufactured by Ackermann gravity steam sterilization with a fractionated process is recommended.

Temperature: 134°C (273°F)  
Pressure: 3 bar  
Duration: ≥ 5 min.

Please comply with all recommendations issued by the manufacturer of your sterilization device with regard to handling and loading. Instruments that are to be sterilized need to be thoroughly exposed to the steam, including inner surfaces. Before using the instruments they need to be cooled down to room temperature.

Other durations and/or temperatures may also be applied. However, when doing so deviations of parameters Processing and Sterilization Instructions of Medical Devices (acc. to ISO 17664) should be validated (Caution: contact the manufacturer of your autoclave to confirm temperatures and/or sterilization durations). Temperature inside the autoclave should not exceed 139°C/182°F. This could cause possible damage to handles, insulation or other non-metallic components. Do not sterilize using hot air or flash autoclave methods. In case only pre-vacuum sterilization can be performed, please adhere to the following parameters:

For Europe: (except Switzerland and France)  
Sterilizer type: Pre-vacuum  
Preconditioning Pulses: 3  
Preconditioning Pressure: 30 psia  
Minimum temperature: 134°C  
Cycle time: 5 min.  
Sample configuration: Individually wrapped

For Switzerland and France:  
Sterilizer type: Pre-vacuum  
Preconditioning Pulses: 3  
Preconditioning Pressure: 30 psia  
Minimum temperature: 134°C  
Cycle time: 18 min.  
Sample configuration: Individually wrapped

Others:  
Sterilizer type: Pre-vacuum  
Preconditioning Pulses: 3  
Preconditioning Pressure: 30 psia  
Minimum temperature: 132°C/270°F  
Cycle time: 4 min.  
Sample configuration: Individually wrapped

**STORAGE**

Store instruments secured against mechanical damage. Use additional wrapping to protect against dust. Do not stack instruments which are packed sterile; especially do not place heavy items on top in order to avoid damage to the sterile packaging of other instruments.

**ADDITIONAL INFORMATION**

Do not exceed maximum loading capacity of the sterilizer when processing multiple instruments in one sterilization cycle.










**APPENDIX**

All product codes covered by these instructions are listed in the following table:

**UROLOGICAL SHEATHS AND OBTURATORS** 

32-4067	32-4205ZV	32-4306
32-4068	32-4221	32-4308
32-4070	32-4221RR	32-4309
32-4071	32-4221ZV	32-4335
32-4075	32-4222	32-4340
32-4076	32-4225	32-4345
32-4080	32-4226	32-4367
32-4081	32-4227	32-4368
32-4085	32-4228	32-4370
32-4086	32-4229	32-4375
32-4090	32-4230	32-4376
32-4091	32-4231	32-4380
32-4095	32-4290	32-4510
32-4100	32-4291	32-4511
32-4105	32-4292	32-4512
32-4106	32-4293	32-4513
32-4110	32-4297	32-4514
32-4115	32-4298	32-4515
32-4205	32-4299	32-4520
32-4205RR	32-4306	32-4521

**SYMBOLS USED ON LABELLING (ACC. DIN EN 980 / DIN EN ISO 1523-1)**

	MANUFACTURER		CONSULT INSTRUCTIONS FOR USE		CAUTION, CONSULT ACCOMPANYING DOCUMENTS
	DATE OF MANUFACTURE		CATALOGUE NUMBER		BSI ASSURANCE UK LIMITED, REGISTERED IN ENGLAND UNDER NUMBER 7605321 AT 389 CHISWICK HIGH ROAD, LONDON W4 4AL, UK
	BATCH CODE		NON-STERILE		PRESCRIPTION USE ONLY