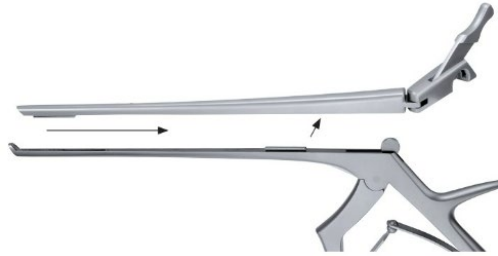


**INSTRUCTIONS FOR USE (EN)  
DISMOUNTABLE INSTRUMENTS: SYSTEMS PROCLEAN AND  
PEACOCK-EYE LOCK**



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**PLEASE READ BEFORE REPROCESSING AND KEEP IN A SAFE PLACE**

**PRODUCT**

These instructions for use apply to the RUDOLF Medical PROCLEAN instruments and to dismountable forceps and scissors.

You are receiving a high-quality product, the proper handling and use of which are described below.

**For professional use only:** The instruments are intended for use by the professional users only (surgeons, operating room nurses, medical device reprocessing technicians).

**Patient population:** There are no restrictions concerning the patient population. It may be left to the discretion and experience of the medical professional to decide whether the benefit outweighs the risk in the given population.



The following instructions refer to the disassembly after the surgical procedure and to the assembly after cleaning and disinfection.

Detailed instructions for reprocessing can be found in the IFU D0462:



RUDOLF Medical instruments are supplied non-sterile and must be cleaned, disinfected and sterilized before the first use and immediately after each use. Protective caps and transport packaging must be removed beforehand.

Before sterilization, we recommend treating the mechanical parts of the instrument with a maintenance oil.

A comprehensive visual and functional inspection of the instruments should be carried out before each use.

**INTENDED PURPOSE**

Punches, conchotomes, skin trephines, bone rongeurs, spoon forceps and septum forceps are intended to cut through skin, mucosa, cartilage or bone tissue and remove it from the surgical site.

Pliers, round forceps, flat forceps, parallel flat forceps and wire forceps are intended to hold and manipulate surgical instruments or wires.

Scissors are intended to cut and manipulate skin, tissue and vessels, as well as to cut suture material and bandages.

Forceps are intended to grip, manipulate and remove organs, bones, tissue and vessels.

**CONTRA-INDICATION**

The medical devices are not intended for the use on the central nervous and circulatory system.

**PRIOR TO EACH USE: VISUAL AND FUNCTIONAL INSPECTION**

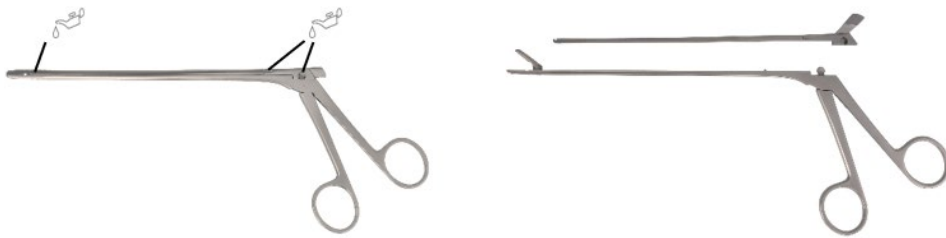
Check for:

- External damage (e.g., deformed shaft, dents, burrs, cracks or sharp edges)
- Correct functioning (e.g., joints, hinges, mobility)
- Cleaning agent or disinfectant residues

## DISMOUNTABLE PROCLEAN INSTRUMENTS

- Dismountable PROCLEAN instruments have hinged and removable sliding shafts.
- There are two PROCLEAN system models:
  - a) Instruments with a removable sliding shaft and a jaw remaining on the lower fixed shaft (forceps)
  - b) Instruments with a completely removable sliding shaft and jaw (punches)

**Figure: Example 1 of PROCLEAN forceps with removable sliding shaft and jaw remaining on the lower fixed shaft**



Disassembled instrument

## DISASSEMBLY

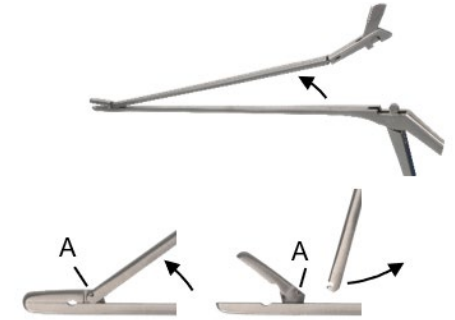
1. Lift the opening lever.



2. Release the opening lever from the peacock-eye lock by pushing it to the right.



3. Open the upper sliding shaft and unhook it from the pin connection (A) of the jaw.



## ASSEMBLY

1. Attach the upper sliding shaft to the pin connection (A) of the jaw.
2. Close the upper sliding shaft parallel to the lower fixed shaft, with the opening lever on the sliding shaft pushed to the right.
3. Push the opening lever to the left into the peacock-eye lock.
4. Push the opening lever down and lock the sliding shaft.
5. Perform a functional check. See the "PRIOR TO EACH USE: VISUAL AND FUNCTIONAL INSPECTION" section in these instructions.

**Figure: Example 2 of PROCLEAN punch with completely removable sliding shaft and jaw**



## DISASSEMBLY

1. Lift the opening lever.



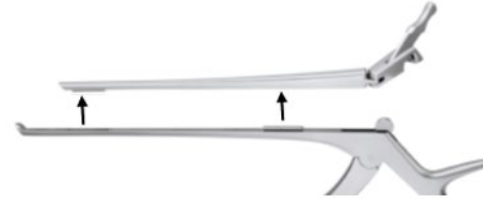
2. Release the opening lever from the peacock-eye lock by pushing it to the right.



3. Slide the upper sliding shaft to the right out of the guide grooves.



4. Remove the upper sliding shaft from the instrument.



## ASSEMBLY

1. The opening lever is flipped to the right. Place the upper sliding shaft on the lower fixed shaft so that the guide grooves engage with each other.
2. Slide the upper sliding shaft to the left until the opening lever can be pressed into the peacock-eye lock.
3. Flip the opening lever down and lock the sliding shaft.
4. Perform a functional check. See the "PRIOR TO EACH USE: VISUAL AND FUNCTIONAL INSPECTION" section in these instructions.

## DISMOUNTABLE INSTRUMENTS WITH PEACOCK-EYE LOCK

- Dismountable instruments with a peacock-eye type lock can be dismantled into three parts, they have a folding-type sliding shaft and, if applicable, a locking latch.

Figure: Example of an instrument with a peacock-eye lock

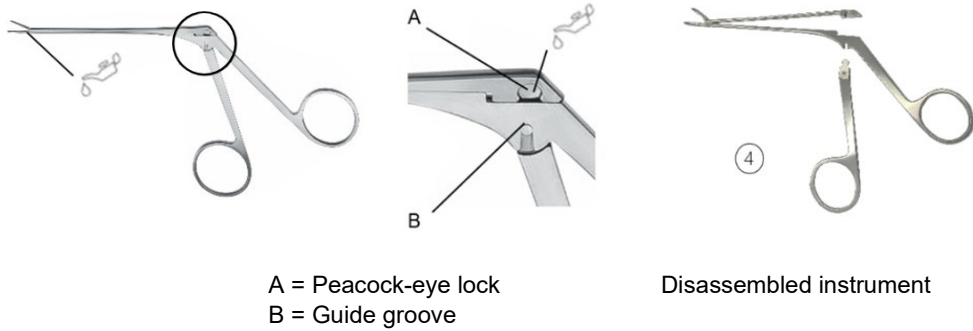
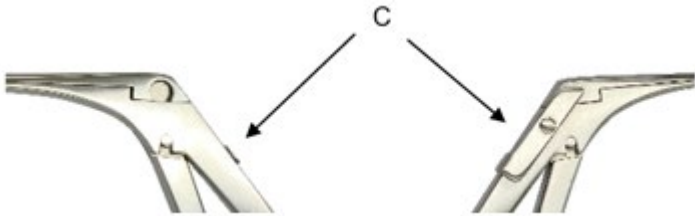
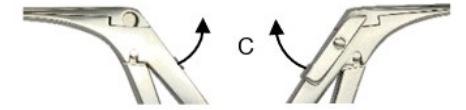


Figure: Example of an instrument with a peacock-eye lock – model with a locking latch (C)

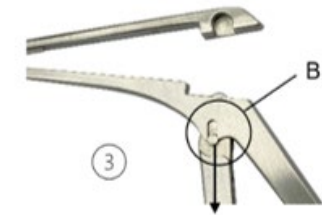


## DISASSEMBLY

1. This step only applies to the model with the locking latch (C): Unlock the locking latch.
2. Press the upper sliding shaft at the outer edge out of the peacock-eye lock and fold it upwards.
3. Release the movable, detachable handle from the guide groove (B) of the fixed instrument part by sliding it downwards.

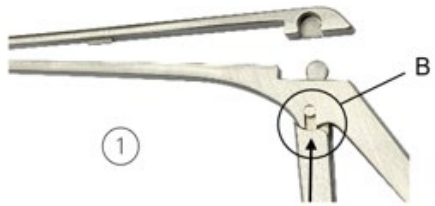


Model with a locking latch



## ASSEMBLY

1. Insert the movable handle completely into the guide groove of the fixed instrument part (B).



2. Push the upper sliding shaft down and then press it lightly against the lower fixed shaft.



3. Squeeze the upper and lower sliding shafts lightly together so that the guide parts of both shafts (D) engage with each other.



4. Connect the peacock-eye lock. To do this, slide the side opening of the upper sliding shaft onto the peacock-eye of the movable handle.









5. This step only applies to the model with a locking latch: Lock it. To do this, press the locking latch down so that it rests against the handle.



6. Perform a functional check. See the "PRIOR TO EACH USE: VISUAL AND FUNCTIONAL INSPECTION" section in these instructions.

## SYMBOLS

	Consult instructions for use.
<b>LOT</b>	Batch code
<b>REF</b>	Article no.
<b>QTY</b>	No. per package
	Non-sterile
	Caution
	Manufacturer
	Date of manufacture
<b>CE</b>	CE mark in accordance with Regulation (EU) 2017/745 for medical devices (MDR) with the identification number of the notified body
	Lubricate with silicon-free, biocompatible white medical oil approved for steam sterilization.
<b>MD</b>	Medical Device