MIZUHO

MES-CK08-062-00EN-0

PREPARED: (2024-02-02) (Version 6)

Be sure to read this instruction manual before using this product.
Keep this manual available for reference when needed.

CE

Instruction for Use

Trade Name: Anesthesia Screen L Type with Clamp

Contraindication/Prohibition

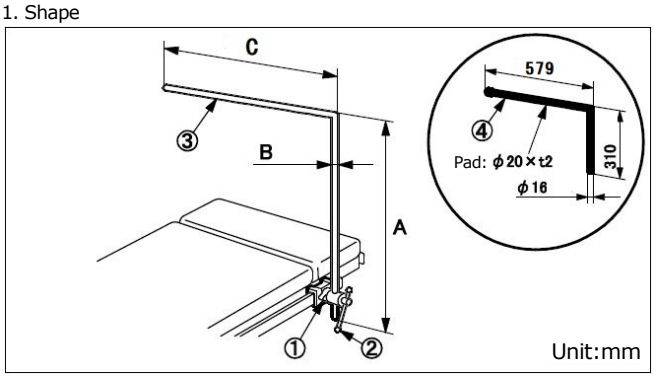
This product is only validated for use with Mizuho operating tables.

Symbol mark for labeling

MD : Medical Device

Specifications

1. Shape



Code No.	Product Description	A (mm)	B (mm)	C (mm)
08-062-00	Anesthesia Screen L Type with Clamp *1	650	16	579
08-062-00-R1	Anesthesia Screen L Type with Clamp (R1) *2	650	16	579
08-062-00-R2	Anesthesia Screen L Type with Clamp (R2) *3	650	16	579

Code No.	Product Description	Reception hole dia. (mm)
08-110-01	Clamp for Round Bar *4	16
08-117-09	Clamp for Round Bar (R1) *5	16
08-117-11	Clamp for Round Bar (R2) *6	16

*1 comes with *4, *2 comes with *5, *3 comes with *6.

*4, *5 and *6: The size of mountable side rail is different.

2. Weight

Main unit (included R1/R2): 1.1 kg

Rail clamp: 0.9 kg

3. Material

① Rail clamp: Stainless steel
② Fixing handle: Stainless steel
③ Screen frame: Stainless steel
④ Pad: Synthetic resin

Note: The number corresponds to those used in 1. Shape.

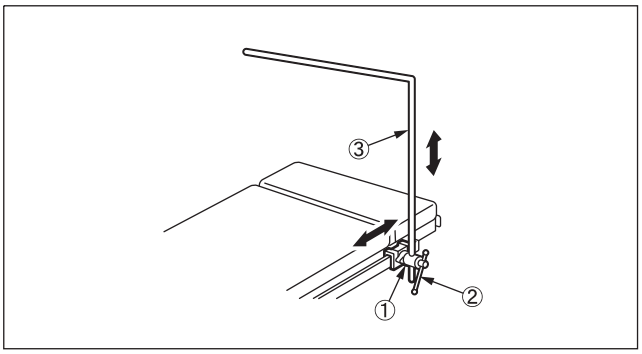
Intended purpose

This is an accessory for the operating table. This is the L-shaped anesthesia screen frame. This frame is to be attached to an angle-adjustable clamp mounted on one of the side rails of the operating table.

Intended user

This product is to be used by health care professionals, including but not limited to surgeons, nurses and biomedical technicians.

Instructions for use



① Rail clamp ② Fixing handle
③ Screen frame

1. Attach the Rail clamp① to the side rail of the operating table.
2. Move the Rail clamp① and the Screen frame③ to a position appropriate for the patient's posture.
3. After the adjustment, tighten the Fixing handle② to fix the Rail clamp① and the Screen frame③.

Warning/Caution

Important caution

1. Ensure all fixing handles are tight. Loose condition can cause the product to come off or move, which may result in a patient injury.
2. When working with the operating table, take care not to allow this product to make contact or interfere with the tabletop or with other tools and appliances used in combination with this product. Otherwise the product could break.
3. For hygiene, be sure to use sterilized drapes on the areas on this product where the patient comes into contact with it.

Storage/Life

1. Do not store the device in high temperatures or in areas with high humidity where the temperature has drastic variations.
2. Service life of this product: 7 years
   (Subject to the specified maintenance and inspection and is stored properly.)

Maintenance/Inspection

<By the user>

1. Check before and after use
   Check that this product is not damaged or broken before and after each use.
2. Cleaning and disinfection
   Wipe off blood, chemicals, contaminants, and other stains with water, and clean the device with gauze or other materials soaked in disinfectants. In addition, use the same disinfectants as the authorized disinfectants on Mizuho's operating table described in the operating table operator's manual.
3. In case of malfunction
   When this product is broken, clearly label the device as "Broken", "Do not use", "Need repair", etc., and contact your local distributor or Mizuho.

<By agents>

Maintenance and inspection can only be carried out certified agents of Mizuho.

Packing

1 piece per pack

Warranty

MIZUHO Corporation will repair defective parts of this product without charge for one year from the date of delivery/installment except for cases of damage caused by a third party's repair, act of nature, improper use or intentional damage. All other warranty terms and conditions are subject to regulations of MIZUHO Corporation.

Disposal

This device must be disposed of in accordance with local regulations. Please contact your local distributor for proper disposal.

Notice

Any serious incident that has occurred in relation to the device should be reported to the manufacturer and the competent authority in which the user and/or patient is established.

MIZUHO Corporation
3-30-13 Hongo, Bunkyo-ku, Tokyo 113-0033, Japan
https://www.mizuho.co.jp

EMERGO EUROPE
Westervoortsedijk 60
6827 AT Arnhem, The Netherlands