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

MES-CK08-089-00EN-0
PREPARED: (2020-09-24) (Version 3)

Instruction for Use

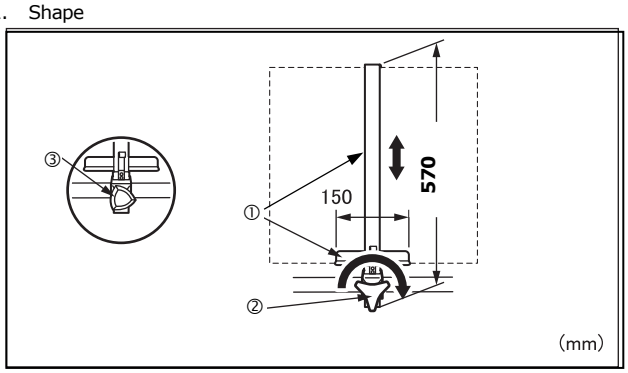
Trade Name: Lateral Cassette Holder

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	Components	Weights (kg)
08-089-03	Lateral cassette holder with clamp	①③ *1	1.3
08-089-03-NC	Lateral cassette holder	① *2	1.1
08-089-05	Lateral cassette holder with clamp Ver.2	①② *3	1.5
08-111-00	Clamp for flat bar	③ *4	0.2
08-111-11	Clamp for flat bar Ver.2	② *5	0.4
08-117-02	Clamp for Flat Bar (R1)	③ *6	0.2
08-117-10	Clamp for flat bar Ver.2 (R1)	② *7	0.4
08-117-12	Clamp for flat bar (R2)	③ *8	0.2
08-117-13	Clamp for flat bar Ver.2 (R2)	② *9	0.4

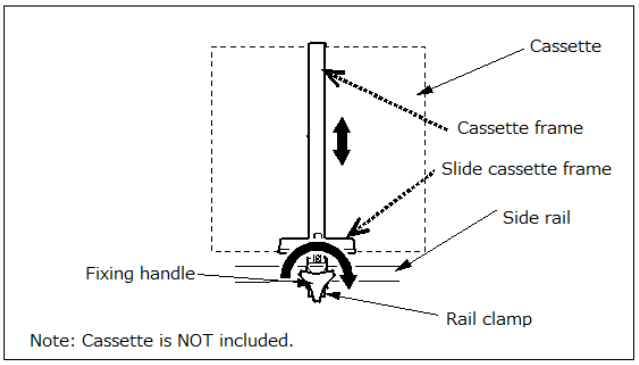
Note: The number corresponds to those used in 1. Shape.
*1: *1 comes with *4.
*2: Please select *6, *7, *8 or *9 to use *2.
*3: *3 comes with *5.
*4 to *9: The size of mountable side rail is different.

2. Material
Lateral cassette holder: Stainless steel rod
Clamp for Flat Bar: Stainless cast steel and plastic molding
Clamp for Flat Bar Ver.2: Stainless cast steel and aluminum

Intended purpose
This is an accessory to an operating table.
It is used for mounting an X-ray cassette to have an X-ray for a lateral part of a patient.

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



The clamp shown above is Clamp for Flat Bar Ver.2

1. Attach the Rail clamp to the side rail of an operating table.
2. Insert the Cassette frame into the Rail clamp and tighten the Fixing handle to temporarily secure the Cassette frame to the side rail.
3. Attach the Cassette to the Cassette frame and Slide cassette frame.
4. Move this product to a position appropriate for the patient's posture.
5. After the adjustment, tighten the Fixing handle to secure the Rail clamp and the Cassette frame.

Warning/Caution

1. Warning
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. Important caution
2-1. When working with an operating table, take care not to allow this product to make contact or interfere with the table top or with other tools and appliances used in combination with this product. Otherwise the product could break.
2-2. 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 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 material moistened with alcohol to disinfect.

3. In case of a malfunction

When this product is broken, clearly label the device as "Broken", "Do not use", "Need repair", etc., and contact your local dealer or Mizuho.

<By agents>

Maintenance and inspection can only be carried out by 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.

Manufacturer

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