

## **DAY - FILM**



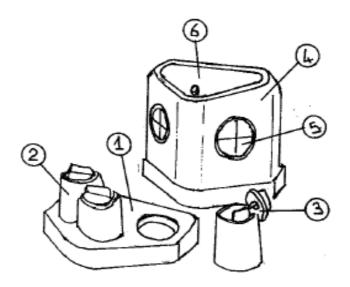
# INSTRUCTION AND INSTALLATION HANDBOOK

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REV.01/2015

#### 1.0 INTRODUCTION

The inactinic container for manual day film light treatment of endoral X- ray film. Made of acidproof material, can reach the max size cm 5 x 7 ,thanks to the appropriate system of introduction of the hands-tight light (5) and the screen control adiactinic (6).



**DIMENSIONS** 

cm 23 x 33 x 31h

#### Legend:

- 1- Base with the slots for the 3 containers.
- 2- Containers with cover for developing, fixing and washing.
- 3- Cover with tweezers.
- 4- Coverage with holes to insert your hands.
- 5- Muffs for kept light.
- 5- Inactinic screen process control.

#### 2.0 USE

Put your hand with the film to be developed in their own right and left muffs (5), then remove the film from container protective and hook it to the tweezers (3).

At this point it is sufficient to dip it in the container (2) before the development and subsequently in that of the fixing.

**ATTENTION!!!** Being the developing process manual, the quality results due to many factors (kind of film, kind of temperature, of the agents etc.) so thay it is hard to indicate the optimal time adlustment. It is advisable to check through the inactinic screen till you get the desired result.

#### 3.0 MAINTENANCE

Occasional care may be needed: the container can be cleaned , if necessary, with a damp cloth. Do not use aggressive agents

#### 4.0 GUARANTEE

The repair and replacement of faulty parts is under guarantee. Guarantee doesn't include indirect damage and every kind of consumer material is excluded.

Guarantee is also not valid in case of failure in observing rules and instructions included in this handbook.



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SCHERMATURE ANTIX - ACCESSORI PER RADIOLOGIA

### CE CONFORMITY DECLARATION

The undersigned "CABLAS S.r.I." office at Via Michelangelo Buonarroti 52/B 20093 Cologno Monzese (MI) - ITALY

DECLARES
Under own responsibility that the product

DAY FILM

Codes

W-DF-01

Which this declaration refers to, meets the Directive 93/42 CEE of the MEDICAL DEVICES, receipt in Italy by D. Lgs.
Nº46 dtd 24/02/97 because considered "
ACCESSORY" (ART. 1 cooma B) and under the classification tethods (enclosure IX) forms part of CLASS 1.

There fore the manufacturer puts CE mark, under the enclosure VII of the same directive

Cologno Monzese, year 2013

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