USER MANUAL: IMAGING AND PAIN MANAGEMENT ACCESSORY - FLUORO EXTENDER

PRODUCT USE

The OAKWORKS® Fluoro Extender is radiolucent and offers extra width only where you need it to achieve greater positioning accuracy for optimum imaging.

SAFETY INSTRUCTIONS

DO NOT place undue weight or downward pressure. It is a positioning device for the arms or legs and should not be used as leverage to get on or off the table. Injury can occur.

DIRECTIONS FOR USE

Slide the Fluoro Extender under the table top pad.

Ask the patient to lie down on the table. Position the patient to suit the needs of the procedure. The weight of the patient will hold the Fluoro Extender in place under the table top pad.

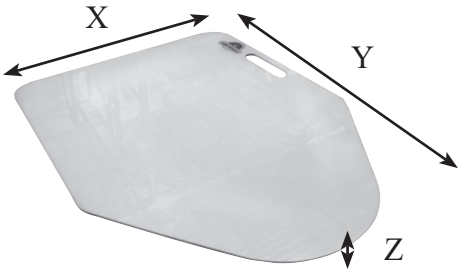
When finished, ask the patient to rise up enough for you to remove the Fluoro Extender from under the table top pad.

CLEANING & DISINFECTION

Reference the Recommended Cleaners and Disinfectant list (MMINML0008-EN) that came with the product or on our website www.oakworksmed.com under product information.

SPECIFICATIONS

Dimensions (X Y Z)	30” x 23-1/2” x 1/4” (76cm x 60cm x .6cm)
Maximum Weight Rating	30 lbs. (13.6 kg)
Radiolucency	Maximum of .85mm AL @ 100 kVp. HVL of 3.6mm
Environmental Conditions	Temperature: 50°F (10°C) to 100°F (38°C) Humidity: 20% to 60% RH Atmospheric Pressure: 98 to 105 kPa



MAINTENANCE & SERVICE

No maintenance required

REPLACEMENT PARTS

- PN 2005: **Fluoro Extender**

OAKWORKS® MEDICAL PRODUCTS

923 East Wellspring Road
New Freedom, PA USA 17349
Phone: 717-235-6807
www.oakworksmed.com

made in the USA
with US & imported parts

USER MANUAL: IMAGING AND PAIN MANAGEMENT ACCESSORY - FLUORO EXTENDER CONT...

SYMBOL IDENTIFICATION

This symbol, when used in this manual and on product labels, represents a caution warning. Be sure to read and comply with all precautions and warnings.

This symbol, when used in this manual and on product labels, indicates the name and address of the manufacturer.

This symbol, when used in this manual or on product labels, indicates the country of manufacture along with date of manufacture of the device next to it.Sitting is prohibited in this area.

UNIQUE DEVICE IDENTIFICATION (UDI) INFORMATION

The UDI Label is located next to the handle cutout

Manufacturer Info

Model Name

EU REP

GTIN

Manufacture Date

Serial Number

Unique Device Information

- GTIN - 14 digit number unique for each variation of a model
- Manufacture Date - Country of manufacture and date the device goes into production in YYYY-MM-DD format
- CE - CE Mark
- SN - Serial Number
- MD - Medical Device Symbol
- QTY - Quantity of the Product

MD CE

EC REP

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