SPENCER

Manuale d'uso e manutenzione
SED - Estricatore IT

Use and Maintenance Manual
SED - Extrication device EN

Benutzungs- und Wartungshandbuch
SED - Rettungskorsetts DE

Manuel d'utilisation et d'entretien
SED - Attelle d'extraction FR

Manual de uso y mantenimiento
SED – Dispositivo de extricación ES

Manual de uso e manutenção
SED - Dispositivo de extração PT

Εγχειρίδιο χρήσης και συντήρησης
SED - Γιλέκο απεγκλωβισμού EL

Ръководство за употреба и поддръжка
SED - Гърбодържател BG

Návod k použití a údržbě
SED - Vyprošťovací korzet CS

SPENCER ITALIA SRL – Via Provinciale n° 12
43038 Sala Baganza (PR) – Italy

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1. MODELS

The following basic models may be subject to implementation or change without notice.

- SED
- SED XS

2. INTENDED USE

2.1 INTENDED USE AND CLINICAL BENEFITS

The extrication device is a first aid apparatus to be used for the extraction of a traumatised patient from a vehicle.
It must be used after the application of the cervical collar to maintain immobilisation and head-torso alignment.

2.2 TARGET PATIENTS

There are no particular indications related to the patient group.
The product configuration is able to accommodate any subject as long as he/she is within the maximum capacity and within the limits of the size of the device.

2.3 PATIENT SELECTION CRITERIA

The patients expected are typically individuals who require extraction from the place where they are stuck.
The typical situation is car accidents.
Before extraction assisted by the use of the device, the patient must be immobilised with a cervical collar to prevent any aggravation of cervical injuries.

2.4 CONTRAINDICATIONS AND UNWANTED SIDE EFFECTS

No particular contraindications or side effects are known with relation to use of the device, as long as it is used in accordance with the user manual.

2.5 USERS AND INSTALLERS

The intended users are rescue workers with in-depth knowledge related to the immobilisation and handling of people suffering from road traffic injuries, spinal injuries and crush injuries.

- Personnel trained for use of the device must also have training in managing lifting and handling suspended loads with people.

These devices are not intended for lay people.

The extrication device is an apparatus intended for professional use only. Do not allow untrained persons to help while using the product, as they may cause injury to themselves or others.
Despite all efforts, laboratory tests, trials, and instructions for use, standards do not always reproduce practice, so the results obtained under actual conditions of product use in the natural environment may sometimes differ significantly.
The best instructions are the continuous practice of use under the supervision of competent and trained personnel.

Operators using the device should be physically able to use the device and have good muscle coordination, as well as strong back, arms, and legs, should it be necessary to lift and/or support the device and the patient. Operators' ability must be assessed before the definition of roles in use of the stretcher.
Operators must be able to provide the necessary patient care.

2.5.1 USER TRAINING

- Regardless of your level of experience with similar devices in the past, you should carefully read and understand the contents of this manual before installing, operating, or servicing this product. In case of any questions, please contact Spencer Italia S.r.l. for the necessary clarifications.
- Operators must also be trained to perform extrication manoeuvres so as to prevent aggravation of spinal injuries or possibly compromised organs.
- The product must be used only by personnel trained in the use of this product and not on other similar products.
- The suitability of the users for use of this product can be attested by the training registration, in which trained persons, trainers, date and place are specified. This documentation must be kept for at least 10 years after the end of the product's life and must be made available to the competent authorities and/or the Manufacturer when requested. In the absence of such documentation, the relevant bodies will apply any foreseen sanctions.
- The product must be put into use only by personnel trained in the use of this product and not on other similar products.

Note: *Spencer Italia S.r.l. is always available for training courses.*

2.5.2 FORMAZIONE INSTALLATORE

Installation is not required.

3. REFERENCE STANDARDS

As Distributor or End-User of the products manufactured and/or marketed by Spencer Italia S.r.l., users are strictly required to be familiar with the legal provisions in force in the country of destination of the goods, applicable to the devices to be supplied (including regulations relating to technical specifications and/or safety requirements) and, therefore, to understand the requirements necessary to ensure compliance of the products themselves with all legal requirements of the territory.

REFERENCE	DOCUMENT TITLE
EU Regulation 2017/745	EU Regulation on Medical Devices

4. INTRODUCTION

4.1 USING THE MANUAL

The purpose of this manual is to provide healthcare professionals with the information necessary for safe and appropriate use and maintenance of the device.

Note: *The Manual is an integral part of the device and therefore it must be kept for the entire life of the device and must accompany it in any changes of use or ownership. If any instructions for use for products other than the one received are present, please contact the Manufacturer immediately before use.*

Spencer products User Manuals can be downloaded from the site **http://support.spencer.it** or by contacting the Manufacturer. Exceptions are those items whose essentiality and reasonable and predictable use are such that it is not necessary to draw up instructions, in addition to the following warnings and indications on the label.

Regardless of your level of experience with similar devices in the past, it is advisable to carefully read and understand the contents of this manual before installing, operating, or servicing this product.

4.2 DEVICE LABELLING AND TRACEABILITY

Each device is provided with a label, placed on the device itself and/or on the packaging, which contains the Manufacturer's identification data, product, CE marking, serial number (SN) or lot number (LOT). This must never be removed or covered.
In the event of damage or removal, request a duplicate from the Manufacturer, or else the warranty will be void as the device can no longer be traced.
If the assigned Lot/SN cannot be traced, the device must be reconditioned, provided only under the responsibility of the manufacturer.

EU Regulation 2017/754 requires manufacturers and distributors of medical devices to keep track of their location. If the device is in a location other than the address to which it was shipped or sold, or if it was donated, lost, stolen, exported or destroyed, permanently removed from use, or if the device was not delivered directly from Spencer Italia S.r.l., please register the device at **http://service.spencer.it**, or inform Customer Service (see § 4.4).

4.3 SYMBOLS

Symbol	Meaning	Symbol	Meaning
CE	Device in compliance with EU Regulation 2017/745	⚠	Danger – Indicates a hazardous situation that may result in a situation directly related to serious injury or death.
MD	Medical device		See the user manual
	Manufacturer	LOT	Lot number
	Date of manufacture	REF	Product code
UDI	Unique Device Identifier	Rx only	Caution: Federal Law restricts this device to sale by or on the order of a licensed practitioner (only for USA Market)

4.4 WARRANTY AND SERVICE

Spencer Italia S.r.l. guarantees that products are free from defects for a period of **one year from the date of purchase.**
For information on correct interpretation of the use, maintenance, installation or return instructions, please contact the Spencer Customer Assistance at tel. +39 0521 541154, fax +39 0521 541222, e-mail **service@spencer.it**
To facilitate service, always indicate the lot number (LOT) or serial number (SN) on the label attached to the package or device itself.
Warranty and service conditions are available at http://support.spencer.it

***Note**: Record and keep with these instructions: lot (LOT) or serial number (SN), if present, place and date of purchase, date of first use, date of checks, user name and comments.*

5. WARNINGS/DANGERS

⚠ Warnings, dangers, notes, and other important safety information are provided in this section and are clearly visible throughout the manual.

At least every 6 months, it is important to check for updated instructions and any changes involving your product. This information is freely available on the website **www.spencer.it** on the specific product page.

⚠ *Product features*

Use of the product for any purpose other than that described in the User Manual is prohibited.

- Before each use, always check the conditions of the product, as specified in the User Manual. In the event of faults/damage that could compromise its functionality/safety, immediately remove it from service and contact the Manufacturer.
- If the product is found to be malfunctioning, immediately use a similar device to ensure continuity of ongoing operations. Non-compliant devices must be taken out of service.
- The product must not be tampered with or modified without the manufacturer's authorisation (modification, tweaking, additions, repair, use of non-approved accessories), as they may constitute imminent danger of injury to persons and material damage. Should these operations be performed, we decline any responsibility for incorrect operation or any damage caused by the product itself; moreover, the CE marking and the product warranty shall be null and void.
- When using the devices, position and adjust them in such a way that they do not hinder operator works or the use of any other equipment.
- Be sure to take every precaution to avoid hazards from contact with blood or body secretions, if applicable.
- Avoid contact with sharp or abrasive objects.
- The times and methods of such shall be agreed upon between the customer and our Sales Offices.
- Operating temperature: from -5°C to + 50°C. .

Storage

- The product must not be exposed or come into contact with thermal sources of combustion or flammable agents, but must instead be stored in a dry, cool place, away from light and sun.
- Do not store the product under other more or less heavy materials that may damage the structure of the product.
- Store and transport the product with its original packaging, otherwise the warranty shall be invalidated.
- Storage temperature: -10°C to +60°C.

Regulatory requirements:

As Distributor or End-User of the products manufactured and/or marketed by Spencer Italia S.r.l., users are strictly required to be familiar with the legal provisions in force in the country of destination of the goods, applicable to the devices to be supplied (including regulations relating to technical specifications and/or safety requirements) and, therefore, to understand the requirements necessary to ensure compliance of the products themselves with all legal requirements of the territory. .

- Promptly and in detail notify Spencer Italia S.r.l. (already in the quotation request phase) about possible fulfilments by the Manufacturer necessary for the compliance of products with specific legal requirements of the territory (including those deriving from regulations and/or regulatory provisions of another nature).
- Act with due care and diligence to help ensure compliance with the general safety requirements of the devices placed on the market, providing end-users with all the information necessary to carry out periodic revisions on the supplied devices, exactly as indicated in the User Manual.
- **Participate in safety checks on products** placed on the market, transmitting information regarding product risks to the Manufacturer as well as to the Competent Authorities for their respective actions.
- Without prejudice to the above, the Distributor or End-User shall assume wider liability related to non-compliance with non-fulfilment of the above-mentioned obligations, with consequent obligation to indemnify and/or hold Spencer Italia S.r.l. harmless from any possible injurious effect.
- With reference to EU Regulation 2017/745, please note that public or private operators who, when exercising their activity, detect an incident involving a medical product are required to notify the Ministry of Health, within the terms and in the manner established by one or more ministerial decrees, and notify the Manufacturer. Public or private health care professionals are required to notify the Manufacturer of any other incident that may allow the adoption of measures to ensure the protection and health of patients and users.

General warnings for medical devices

⚠ User must carefully read the following in addition to the general warnings.

- The application of the device should not last longer than the time required for first aid operations and subsequent transport to the nearest rescue point.
- Qualified personnel and at least two operators must be present during use of the device.
- Do not use if the device or parts of it are punctured, torn, frayed, or excessively worn.
- Follow the internal procedures and protocols approved by your organisation.
- Do not alter or modify the device arbitrarily, as doing so could result in unpredictable operation and damage to the patient or rescuers and shall void the manufacturer's warranty and release the manufacturer from all liability.
- Disinfection operations must be carried out in accordance with the validated cycle parameters, as stated in the specific technical standards.
- Do not use drying machines to dry the device.

6. SPECIFIC WARNINGS

To use the extrication device, you must also have read, understood and carefully follow all the instructions in the user manual.

- Always comply with the maximum capacity, if any, indicated in the User Manual. Maximum load capacity means the total weight distributed according to human anatomy. When determining the total weight load on the product, the operator should consider the weight of the patient, equipment and accessories. In addition, the operator should assess whether the overall size of the patient reduces the functionality of the product.
- The device must only be used by trained personnel
- The device is not intended for winching manoeuvres.
- Attaching other devices or systems to the device that have not been approved by the manufacturer may cause serious injuries and/or be a source of accidents.
- Practice under simulated conditions to ensure that you are familiar with the manoeuvres.

- If foreseen for the device, make sure that operators are in good physical condition before lifting, as listed in the User Manual.
- **The maximum weight, which weighs on each operator, must comply with local health and safety requirements.**
- Establish a maintenance program and periodic checks, identifying a designated reference person. The person entrusted with routine maintenance of the device must ensure the basic requirements envisaged by the manufacturer within these operating instructions.
- All maintenance activities must be recorded and documented with the relevant technical intervention reports. This documentation must be kept for at least 10 years after the end of the device's life and must be made available to the competent authorities and/or the manufacturer when requested.
- Use only original or Spencer Italia S.r.l. approved components/replacement parts and/or accessories to carry out any operation without causing alterations or modifications to the device. Otherwise, we decline all responsibility regarding incorrect operation or any damage caused by the device to the patient or the operator, invalidating the warranty and invalidating compliance with EU Regulation 2017/745.
- Never leave the patient on the device unsupervised, as they could get injured.
- Avoid contact with sharp objects.
- The extrication device must not be exposed, much less come into contact with thermal sources of combustion or flammable
- Follow approved Emergency Medical Service procedures for patient immobilization and transportation.
- Follow approved Emergency Medical Service procedures for patient positioning and transportation.
- Before lifting, make sure that operators have a secure grip on the supporting structure of the device.
- Do not bend the extrication device excessively or for long periods.
- Do not apply the device unless the cervical spine has been immobilised with a suitable collar.
- Avoid twisting the patient's chest when applying the device and adjusting the belts.
- At least two operators in a suitable physical condition are required to use the device. Consequently, they must have strength, balance, coordination, common sense and be trained in the correct operation and application of the device.
- For patient loading techniques for particularly heavy patients, for operations on steep terrain or in special and unusual circumstances, the presence of more than two operators is recommended in addition to the two minimum operators.
- Before each use, always check the conditions of the device and its accessory components, as specified in the user manual. In case of faults or damage that may compromise the functionality and safety of the device, and therefore the patient and the operator, remove the device from service or replace the components that are not intact.
- Use the device only as described in this user manual.
- Do not arbitrarily alter or modify the device to adapt it to unforeseen conditions of use: doing so could result in unpredictable operation and damage to the patient or rescuers and shall void the manufacturer's warranty and release the manufacturer from all liability.
- To preserve the life of the device, protect it as much as possible from UV rays and adverse weather conditions.
- Operators must be fully aware of the sequence of belt attachment.

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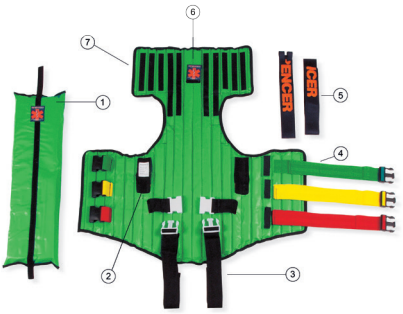
7. RESIDUAL RISK

No residual risks, or rather risks that could arise despite compliance with all warnings in this user manual, have been identified.

8. TECHNICAL DATA AND COMPONENTS

Note: Spencer Italia S.r.l. reserves the right to make changes to specifications without notice.

ELEMENTS/DESCRIPTIONS AND MATERIALS

1	**Nape cushion** Padded and PVC-coated, providing soft support for the back of the patient's neck		5	**Set of chin and forehead bands** Made of polypropylene, they improve the fixation of the patient's head to the device.	
2	**Lifting handles** Made of polypropylene and placed on the back of the device, they allow the device to be gripped when applied to the patient to perform extrication manoeuvres.		6	**Main body** Made of wooden splints, covered with PVC fabric to which the other elements are sewn, it provides horizontal flexibility to be wrapped around the patient and offers vertical rigidity to increase immobilisation.	
3	**Groin belts** Made of polypropylene, they are passed through the groin area, distributing the forces exerted during extrication to the thigh area.				
4	**Abdominal belts** Made of polypropylene, they allow the device to be attached to the patient. The XL accessory set can be applied to these, if appropriate.		7	**Nape area** Part of the main body on which the strap strips are applied for the application of the nape cushion	

	SED	SED XS
Length	830 mm	690 mm
Width	900 mm	640 mm
Maximum thickness (at the hooks)	25 mm	25 mm
Overall dimensions wrapped with bag (approx.)	850x250x120 mm	730x260x100 mm
Belt length	74 ± 2 cm	52 ± 2 cm
Belt length with XL set	-	-
Materials	PVC, Nylon, PP	PVC, Nylon, PP
Weight without bag	2,6 kg	2,10 kg
Weight with bag	2,85 kg	2,40 kg
Maximum load capacity	230 kg	60 kg

9. COMMISSIONING

For first use, check that:

- Packaging is intact and has protected the device during transportation
- Check that all parts included in the packing list are present
- General functionality of the device
- Product cleanliness
- There are no cuts, holes, lacerations or abrasions on the entire structure, including belts and footrests where provided.
- Correct stitching and tightening of belts

See paragraph 11 on how to carry out the above-mentioned checks.
Do not modify the device or its parts for any reason as this could cause damage to the patient and/or rescuers.

⚠ Failure to take the above measures will preclude safe use of the device, resulting in risk of damage to the patient, operators and the device itself.

For subsequent use, perform the operations specified in paragraph 12.

If the above conditions are met, the device may be considered ready for use; otherwise, you must immediately remove the device from service and contact the Manufacturer.

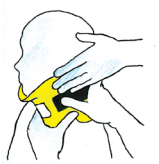
Do not alter or modify the device arbitrarily, as doing so could result in unpredictable operation and damage to the patient or rescuers and will void the warranty and release the Manufacturer from all liability.

10. OPERATING CHARACTERISTICS

See paragraph 11 - Proper use for operating characteristics.

11. PROPER USE

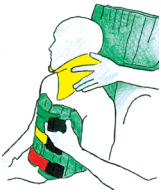
Primary medical evaluations must be carried out before intervening on the patient.



1 **Applying the cervical collar:** Before applying the extrication device, it is necessary to immobilise the cervical section using a collar appropriate to the patient's condition.
Follow local protocols as well as the manufacturer's user manual for the collar in use.

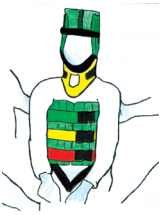
3 Release the groin straps from the back by letting them drop to the sides and wrap the thoracic flaps around the patient's torso, passing them under the arms. Adjust the position of the device by lifting it so that the chest flaps become a support for the armpits. During the application, operators should avoid moving the patient's head to avoid aggravating any injury.
Check that the device has been correctly positioned, i.e. as close as possible to the patient's back and well aligned with the spine.

2 **Positioning the extrication device behind the patient**
Check that there are no items that could obstruct the application of the device (belt, wallet, etc.).
Open the device completely, turn the buckles towards the seat and tilt it 45° to insert it between the patient's back and the seat, aligning it with the patient's spine.



4 Fasten the chest belts in the following sequence. Generally accepted as correct:
1 – Intermediary (Yellow)
2 – Lower (Red)
3 – Upper (Green)
This sequence may not be indicated in all rescue situations. Follow the instructions of your local health authority.
When applying it to pregnant patients, some of the thoracic flaps have to be folded to leave the abdomen uncovered.
A possible closure configuration, which may vary depending on the size of the abdomen and breasts, is as follows:
- Intermediary belt in the respective buckle
- Lower belt in upper buckle
- Upper belt in the lower buckle

5 Close the groin straps by sliding them under the patient's legs using any free space left by the seat.
Check that the belts are straight in relation to the anchorage point and as close as possible to the symmetry axis of the device.
Hook the belts into their buckles and tighten them until they come into contact with the patient.
A further tightening must be carried out before extrication.
The use of crotch belts requires particular care in the case of femur or pelvis fractures. The device handles can help obtain better vertical positioning and alignment of the device.



6 Do not immobilise the head until checking the correct position of the extrication device in relation to the size of the patient. Take care not to cause head and neck movements that could compromise the patient's condition. If there is a gap between the extrication device and the patient's head, collar or shoulders, fill the nape space with the appropriate support cushion, making sure not to hyper extend or flex the patient's neck.
While one operator holds the patient's head in position and positions the flaps, the other operator must secure the headband by tilting the back slightly downwards. This reduces the risk of the headband slipping over the head.
Then apply the chin strap, taking care to support the cervical collar and tilt it towards the patient's ears. This avoids immobilising the jaw and preventing the mouth from opening. Ensure that the straps are correctly secured and symmetrically positioned.

7 When the stretcher or spine board has been brought to the extrication site, the final steps of the application can be carried out, i.e. further tightening of the extrication straps, therefore:
1 – Tighten the upper belt (green)
2 – Check and tighten the lower belt (red)
3 – Check and tighten the groin belts.
After checking the correct application, the two operators can grasp the handles of the device to rotate the patient so that his or her back faces the inside of the vehicle. Rotation must take place without twisting the spinal cord or the cervical spine.
The two operators can then grasp the nearest handle of the extrication device with one hand and the back of the patient's knees with the other. After coordinating the operation correctly, they can lift the patient.
If needed, a third operator could apply a spine board.
The upper belt (green) must be loosened immediately after extrication to allow for proper expansion of the chest.

12. CLEANING AND MAINTENANCE

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Spencer Italia S.r.l. declines all responsibility for any direct or indirect damage which is the consequence of improper use of the product and spare parts and/or in any case of any repair carried out by a person other than the Manufacturer, who uses internal and external technicians authorised to do so; moreover, doing so will invalidate the warranty.

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- The operator must wear suitable personal protective equipment, such as gloves, goggles, etc. during all checking, maintenance and cleaning operations.
- Establish a maintenance schedule, periodic inspections and extend the average life span, if foreseen by the Manufacturer in the User Manual, identifying a reference person who meets the basic requirements set forth in the User Manual.
- **The frequency of inspections is determined by factors such as legal requirements, type of use, frequency of use, and environmental conditions during use and storage.**
- Repairs of products manufactured by Spencer Italia S.r.l. must be carried out by the Manufacturer, who shall make use of specialised internal or external technicians who, using original spare parts, shall provide quality repair service in strict compliance with the technical specifications indicated by the Manufacturer. Spencer Italia S.r.l. declines any responsibility for any direct or indirect damage which is a consequence of improper use of spare parts and/or any repair work carried out by unauthorised parties.

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- Where foreseen, use only original or Spencer Italia S.r.l. approved components/spare parts and/or accessories in order to carry out all operations without causing alterations or modifications to the product.
- All maintenance and overhaul activities must be recorded and documented with the relevant technical operation reports. This documentation must be kept for at least 10 years after the end of the product's life and must be made available to the competent authorities and/or the Manufacturer when requested.

FR

- Cleaning, provided for reusable products, must be carried out in accordance with the Manufacturer's instructions in the User Manual in order to avoid the risk of cross-infection due to the presence of body fluids and/or residues.
- If required, the product and all its components must be washed and left to dry completely before storage.

12.1 CLEANING

ES

Failure to carry out the correct cleaning operations could increase the risk of cross-infection due to presence of body fluids and/or residues.
The operator must wear suitable personal protective equipment, such as gloves, goggles, etc. during all checking and cleaning operations. .

Clean the device and its parts with soap and water using a brush with medium-hard bristles. Rinse thoroughly with lukewarm water, making sure you have removed all traces of soap, which may deteriorate or compromise conditions and durability. **Avoid using high-pressure water.** Let dry completely before storing. Drying after washing or after use in a wet environment must be natural and not forced. Do not use flames or other direct heat sources.

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When **disinfecting**, use products that do not have a solvent or corrosive action on materials constituting the device, in addition to being classified as medical-surgical devices. Be sure to take all precautions to ensure that there is no risk of cross-infection or contamination of patients and operators.
After disinfection, rinse all parts with warm water and allow the device to dry completely before storage, as residual moisture may lead to the formation of mould. .

12.2 ROUTINE MAINTENANCE

EL

The operator must wear suitable personal protective equipment, such as gloves, goggles, etc. during all checking, maintenance and cleaning operations.

The device does not require a routine maintenance program, but checks must be made to verify:

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- All parts are present
- Integrity of the belts
- Integrity of the lifting handles
- Integrity of the groin belts
- Integrity and operation of the buckles
- Integrity of internal slats
- Integrity of the chin and forehead bands
- General functionality of the device
- Nape cushion conditions

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- Cleanliness of the device (remember that the failure of cleaning may cause the risk of cross infections)
- Fulfilment of the requirements of the user manual in section 5 Warnings and 6 Specific Warnings.
- Fulfilment of the requirements of the manual in section 11 Proper use

Use only original or Spencer Italia S.r.l. approved components/replacement parts and/or accessories to carry out any operation without causing alterations or modifications to the device. Otherwise, we decline all responsibility regarding incorrect operation or any damage caused by the device to the patient or the operator, invalidating the warranty and invalidating compliance with EU Regulation 2017/745.

12.3 PERIODIC OVERHAUL

No periodic overhaul is foreseen for the device.

12.4 SPECIAL MAINTENANCE

Special maintenance can only be carried out by the Manufacturer, who uses internal and external technicians specialised and authorised by the Manufacturer itself.

Only maintenance activities carried out by specialised technicians authorised by the Manufacturer are considered valid by Spencer Italia S.r.l..
The end-user can replace only the spare parts indicated in § 15.

12.5 LIFE SPAN

The device, if used as described in the following instructions, has a life span of 5 years from the date of purchase.

Spencer Italia S.r.l. will accept no responsibility for incorrect operation or damage caused by the use of devices that have exceeded the maximum allowable life span.

13. TROUBLESHOOTING TABLE

PROBLEM	CAUSE	REMEDY
Excessive vertical bending	One or more of the internal slats may be broken	Put the device out of service immediately and replace it with a similar one.
The buckles do not close properly	The buckles may be broken	Immediately remove the device from service and contact the manufacturer
The device has mould	The device has been stored in very humid conditions or without waiting for adequate drying following cleaning procedures	Put the device out of service immediately and replace it with a similar one.

If a problem or fault is detected that does not correspond to the above, please contact Spencer Italia Srl customer care service.

14. ACCESSORIES

Code	Description	Compatible
SR00101B	SED XL KIT	SED

15. SPARE PARTS

Code	Description	For model
RISR001	SED ADULT CHIN/NECK STRAP SET	SED
RISR002	REPLACEMENT BAG FOR ADULT SED	SED
RISR003	REPLACEMENT CUSHION FOR ADULT SED	SED
RISR004	BLACK PLASTIC MALE/FEMALE HOOK h 50mm	SED
RISR005	SED PEDIATRIC CHIN/NECK STRAP SET	SED XS
RISR006	REPLACEMENT BAG FOR PEDIATRIC SED	SED XS
RISR007	REPLACEMENT CUSHION FOR PEDIATRIC SED	SED XS
RISR008	WHITE PLASTIC MALE/FEMALE HOOK h 50mm	SED XS

16. DISPOSAL

When devices and their accessories are no longer suitable for use, they can be disposed of as normal municipal solid waste if they have not been contaminated by special agents. Otherwise, follow the regulations in force regarding disposal.

Notice

The information in this manual is subject to change without notice.
The images are included as examples and may vary slightly from the actual device.

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Code	**CCI5288**
Erstausgabe:	**13/04/2021**
Rev. 1	**13/04/2021**
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spencer.it | support.spencer.it - e-mail: info@spencer.it | service: service@spencer.it
Ph./Tel. +39.0521.541111
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