LUCAS[™]2

Chest Compression System

Instructions for Use



Thank you for choosing the LUCAS™2 Chest Compression System!

With LUCAS™ 2 your cardiac arrest patients will receive 100 chest compressions per minute with a depth of 1.5 to 2 inches as recommended in the ERC (European Resuscitation Council).

If you have any questions about this product or its operation, please contact your local distributor or the manufacturer JOLIFE AB.

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1 Important user information



The information in these Instructions for Use applies to the LUCAS $^{\text{TM}}$ 2 Chest Compression System, also referred to as LUCAS.

All operators must read the complete Instructions for Use before operating the LUCAS Chest Compression System.

The Instructions for Use must always be easily accessible to the operators of LUCAS.

Always follow local and/or international guidelines for cardiopulmonary resuscitation (CPR) when you use LUCAS.

The use of other medical equipment or drugs in conjunction with LUCAS can affect the treatment. Always consult the Instructions for Use of the other equipment and/or drugs to make sure that they are appropriate for use in conjunction with CPR.

LUCAS can only be bought by, or on the order of, a licensed medical practitioner.

TRADEMARKS

LUCAS™ is a trademark of JOLIFE AB.

DECLARATION OF CONFORMITY

LUCAS Chest Compression System complies with the requirements of the European Medical Device 93/42/EEC. It is marked with the CE-symbol:

CE 0434

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2 Introduction

2.1 LUCAS™ Chest Compression System

The LUCAS™ Chest Compression System is a portable tool designed to overcome problems that have been identified with manual chest compression. LUCAS assists rescuers giving 100 chest compressions per minute with a depth of 1.5 to 2 inches as recommended in the ERC (European Resuscitation Council) guidelines¹.

2.2 Intended use

The LUCAS Chest Compression System is to be used for performing external cardiac compressions on adult patients who have acute circulatory arrest defined as absence of spontaneous breathing and pulse, and loss of consciousness.

LUCAS must only be used in cases where chest compressions are likely to help the patient.

2.3 Contraindications

Do NOT use the LUCAS Chest Compression System in these cases:

- If it is not possible to position LUCAS safely or correctly on the patient's chest.
- Too small patient: If you cannot enter the PAUSE mode or ACTIVE mode when the pressure pad touches the patient's chest and LUCAS alarms with 3 fast signals.
- Patient too large: the Upper Part of LUCAS cannot be locked to the Back Plate without compressing the patient's chest.

Always follow local and/or international guidelines for CPR when using LUCAS.

2.4 Side effects

The International Liaison Committee on Resuscitation (ILCOR) states these side effects of CPR²:

"Rib fractures and other injuries are common but acceptable consequences of CPR given the alternative of death from cardiac arrest. After resuscitation, all patients should be reassessed and re-evaluated for resuscitation-related injuries."

Apart from the above, bruising and soreness of the chest are common during the use of the LUCAS Chest Compression System.

2.5 Main parts

The main parts of the LUCAS Chest Compression System include;

- A Back Plate which is positioned underneath the patient as a support during external chest compressions.
- An Upper Part which contains the proprietary and rechargeable LUCAS Battery and the compression mechanism with the disposable Suction Cup.
- A Stabilisation Strap which helps to secure the position of the device relative to the patient.
- A padded Carrying Bag.

European Resuscitation Council Guidelines for Resuscitation 2005, Resuscitation 2005 67S1;S13-

 ²⁰⁰⁵ International Consensus on Cardiopulmonary diopulmonary Resuscitation and Emergency Cardiovascular Care Science with Treatment Recommendations. Resuscitation 2005;67:195

2.6 LUCAS[™] components



- 1. User Control Panel
- 2. Hood
- 3. Patient Strap
- 4. Release ring
- 5. Support leg
- 6. Claw locks
- Back Plate
- 8. DC input
- 9. Bellows
- 10. Suction Cup
- 11. Power Supply
- 12. Power Supply cord
- 13. Battery
- 14. Pressure pad
- 15. Upper Part
- 16. Vent holes
- 17. Car Power Cable
- 18. Carrying Bag

LUCAS Stabilisation Strap

- 19. Cushion strap
- 20. Buckle
- 21. Support leg strap

2.7 User Control Panel





ON/OFF:

LUCAS will power up/power down when you push this key for 1 second. When LUCAS powers up, it automatically does a self-test of the functions and the protection safety system. When the self-test is complete the green LED (Light Emitting Diode) beside the ADJUST key comes on. This procedure takes approximately 3 seconds.



ADJUST:

This mode is used when you want to adjust the position of the Suction Cup. When you push this key, you can move the Suction Cup up or down. To adjust the Start Position of the Suction Cup, manually push the Suction Cup down onto the chest of the patient using two fingers.



PAUSE:

When you push this key, the compression mechanism temporarily stops and is locked in the Start Position. Use this function when you want to stop LUCAS temporarily but still want to maintain the Suction Cup Start Position.



ACTIVE (continuous):

When you push this key, LUCAS performs continuous chest compressions. The green LED signal will blink 8 times each minute to indicate when ventilation is required during compression.



ACTIVE (30:2):

When you push this key, LUCAS performs 30 chest compressions and then temporarily stops for 3 seconds. During the stop time, the operator can perform 2 ventilations. After this stop time the cycle starts again. An intermittent LED combined with an alarm signal will alert the operator before each ventilation pause.



MUTE:

If you push this key when LUCAS is operating, you mute the alarm for 60 seconds. If you push this key when LUCAS is turned OFF, the Battery indicator shows the charge status of the Battery.



Battery indicator:

The three green LEDs show the Battery charge status:



• Three green LEDs: Fully charged



Two green LEDs: 2/3 chargedOne green LED: 1/3 charged



• One intermittent orange LED and alarm during operation: low battery, approximately 10 minutes of operating capacity remaining.



• One intermittent red LED and an alarm signal: the Battery has run down and must be recharged.



• One constant red LED and an alarm signal: the Battery is defective.



Note: When the LED to the far right is orange and not green, the Battery has reached the end of its service life. JOLIFE AB recommends that you replace this Battery with a new one.



Alarm indicator:

A red LED and an alarm signal indicate a malfunction.

Refer to Troubleshooting 8: 8.1 for indicators and alerts during normal operation. 8.3 for malfunction alarms.

3 Safety precautions

To ensure maximum safety, always read this section carefully before operating, carrying out any work on the equipment or making any adjustments.

3.1 Signal words

Throughout the manual, signal words are indicated with, "WARNING" or "CAUTION".

- CAUTION signal word used to indicate a potentially hazardous situation which, if not avoided, could result in minor or moderate injury.
- WARNING signal word used to indicate a potentially hazardous situation which, if not avoided, could result in death or serious injury.

3.2 Personnel

JOLIFE AB recommends that the LUCAS Chest Compression System is only used by persons with medical skills such as: first responders, ambulance personnel, nurses, physicians or medical staff, who have:

- undertaken a CPR course according to resuscitation guidelines, e.g. American Heart Association, European Council of Resuscitation or equivalent,
- AND received training in how to use LUCAS.

3.3 Contraindications

Do NOT use the LUCAS Chest Compression System in these cases:

- If it is not possible to position LUCAS safely or correctly on the patient's chest.
- Too small patient: If you cannot enter the PAUSE mode or ACTIVE mode when the pressure pad touches the patient's chest and LUCAS alarms with 3 fast signals.
- Patient too large: the Upper Part of LU-CAS cannot be locked to the Back Plate without compressing the patient's chest.

Always follow local and/or international guidelines for CPR when using LUCAS.

3.4 Side effects

The International Liaison Committee on Resuscitation (ILCOR) states the following side effects of CPR³:

"Rib fractures and other injuries are common but acceptable consequences of CPR given the alternative of death from cardiac arrest. After resuscitation, all patients should be reassessed and re-evaluated for resuscitation-related injuries."

The above side effects, as well as bruising and soreness of the chest, are common during use of the LUCAS Chest Compression System.

 ²⁰⁰⁵ International Concensus on Cardiopulmonary Resuscitation and Emergency Cardiovascular Care Science with Treatment Recommendations. Resuscitation 2005; 67:195



3.5 Symbols on the device



Symbol	Meaning
	Caution – keep your fingers away Do not put your hands on or below the Suction Cup while the LUCAS is operational. Keep your fingers away from the claw locks when attaching the Upper Part or lifting the patient.
	Caution – do not lift by the Patient Straps Do not use the Patient Straps to lift the patient. The straps are only to attach the patient's arms to LUCAS.
12	Place the lower edge of the Suction Cup immediately above the bottom of the sternum, as indicated in the figure. The Suction Cup should be central on the chest.
	Pull the release rings to release the Upper Part from the Back Plate.
2	Do not reuse - Single use only.
Art no. 300000-00 ⊙-@-⊙ 12-24VDC — +	DC input.

Symbols on type label

Symbol	Meaning
<u> </u>	Caution – see instructions for use All operators must read the complete Instructions for Use before operating the LUCAS Chest Compression System.
\mathbb{M}	Year of manufacture.
X	Battery and/or electronics may not be disposed of in normal waste.
IP 43	Degree of protection provided by enclosure per IEC 60 529.
	DC voltage.
★	Defibrillation protected type BF patient connection.

3.6 General safety precautions

Caution - use only approved accessories

Use only JOLIFE AB-approved accessories with LUCAS. LUCAS may not operate correctly if you use accessories that are not approved. Use only LUCAS Batteries and the LUCAS Power Supply that are designed for LUCAS. If you use other batteries or power supplies you can cause permanent damage to LUCAS. This will also void the warranty.

Caution - liquid

Do not immerse LUCAS in liquid. The device can be damaged if liquid enters the hood.

3.7 Battery

WARNING - LOW BATTERY

When the orange Battery LED shows an intermittent light, either:

- Replace the Battery with one that is charged.
- Connect the external LUCAS Power Supply.

Caution - keep Battery in place

The Battery must always be in place for LUCAS to be able to operate, even when powered by the external Power Supply.

To minimise interruptions, we recommend that there is always a spare, charged, LUCAS Battery in the Carrying Bag.

3.8 Operation

WARNING - UNSATISFACTORY POSITION

Start manual CPR again if it is not possible to position LUCAS safely and correctly on the patient's chest.

WARNING - INCORRECT POSITION ON CHEST

If the pressure pad is not in the correct position in relation to the sternum, there is an increased risk of damage to the rib cage and the internal organs. Also, the patient's blood circulation is compromised.

WARNING - INCORRECT START POSITION

The patient's blood circulation is compromised if the pressure pad presses down too heavily or too lightly on the chest. Push the **ADJUST** key and adjust the height of the Suction Cup immediately.

WARNING - CHANGED POSITION DURING OPERATION

If the position of the Suction Cup changes during operation or during defibrillation, immediately push **ADJUST** and adjust the position. Always use the LUCAS Stabilisation Strap to help secure the correct position.

Caution - defibrillation electrodes

Position the defibrillator electrodes and wires so that they are not under the Suction Cup. If there are already electrodes on the patient, make sure that they are not positioned where the Suction Cup should be placed. If they are, you must apply new electrodes.

Caution - gel on chest

If there is gel on the patient's chest (e.g. from ultrasound examination), the position of the Suction Cup can change during use. Remove all gel before you apply the Suction Cup.

Caution - Stabilisation Strap applicationDelay the application of the LUCAS Stabilisation Strap if this prevents or delays any medical treatment of the patient.

Caution - adjunctive therapies

The use of other medical equipment or drugs in conjunction with LUCAS can affect the treatment. Always consult the Instructions for Use of the other equipment and/or drugs to make sure that they are appropriate for use in conjunction with CPR.

WARNING - ECG interference

Chest compressions interfere with ECG analysis. Push **PAUSE** before you start the ECG analysis. Make the interruption as short as possible. Push **ACTIVE** (continuous) or **ACTIVE** (30:2) to start the compressions again.



WARNING - PATIENT INJURY

Do not let the patient or the device stay unattended when LUCAS is working.

Caution - keep your fingers away

Do not put your hands on or below the Suction Cup when LUCAS is working. Keep your fingers away from the claw locks when attaching the Upper Part or lifting the patient.

Caution - IV access

Make sure that IV access is not obstructed.

Caution - do not block the vent holes Do not obstruct the vent holes under the hood as this can cause the device to overheat.

Caution - device alarms

If there is any malfunction during operation the red Alarm LED will be illuminated and a buzzer signal heard. For troubleshooting, see section 8.3.

WARNING - MALFUNCTION

If there are interruptions, or the compressions are not sufficient, or something unusual occurs during operation: Push **ON/OFF** for 1 second to stop LUCAS and remove the device. Start manual chest compressions.

Caution - do not use the Patient Straps for lifting

Do not use the Patient Straps to lift the patient. The straps are only to attach the patient's arms to LUCAS.

3.9 Service

We recommend yearly servicing of LUCAS to ensure that it operates correctly. Use the original shipping box, when you send LUCAS for servicing. Keep the original shipping box, with padding, for this purpose.

WARNING - DO NOT OPEN

Never open the casing of LUCAS. Do not change or modify external or internal parts of LUCAS.

Unless otherwise specified, all servicing and repairs must be done by service personnel that are approved by JOLIFE AB.

If the above conditions are not followed, this can lead to patient/operator injury or death, and will void the warranty.

Consult your distributor or JOLIFE AB for current information on where to send LUCAS for maintenance.

4 First use preparations

4.1 Delivered items

LUCAS™2 Chest Compression System is supplied in one box with:

- A LUCAS device (Upper Part and Back Plate)
- 3 disposable LUCAS Suction Cups
- · A LUCAS Carrying Bag
- Instructions for Use in the relevant language
- · A rechargeable LUCAS Battery
- · A LUCAS Stabilisation Strap
- · LUCAS Patient Straps

Accessories (optional):

- Disposable LUCAS Suction Cups
- · External LUCAS Battery Charger
- · Extra LUCAS Batteries
- LUCAS Power Supply with Mains cable
- LUCAS 12-24V DC Car Power cable

4.2 The Battery

The proprietary Lithium Polymer (LiPo) Battery is the exclusive power source for LUCAS. You can remove the Battery from LUCAS and recharge it. The Battery is mechanically keyed in LUCAS and in the Battery Charger to make sure it is correctly positioned. The top of the battery has connectors fo both power and data transfer with the Battery charger and with LUCAS.

4.2.1 Charge the Battery

You can charge the LUCAS Battery in two ways:

- In the external LUCAS Battery Charger (optional)
 - put the Battery in the slot on the Battery Charger,
 - plug the Battery Charger power cable in to the mains wall socket.

• Installed in LUCAS:

- put the Battery in the hood slot on LUCAS,
- connect the Power Supply to the DC input on the side of LUCAS,
- plug the Power Supply in to the mains wall socket.



Green LEDs indicate a fully charged Battery.

Caution - keep Battery in position

The Battery must always be in position for LUCAS to be able to operate, even when powered by the external Power Supply.

Caution - use only approved accessories

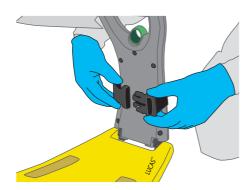
Use only JOLIFE AB-approved accessories with LUCAS. LUCAS does not operate correctly if you use accessories that are not approved. Use only LUCAS Batteries and the LUCAS Power Supply that are designed for LUCAS. If you use other batteries or other Power Supply you can cause permanent damage to LUCAS. This will also void the warranty.



4.3 Prepare the LUCAS™ Stabilisation Strap

Before using LUCAS for the first time, attach the support leg straps, which are part of the Stabilisation Strap, to the LUCAS support legs.

- 1. Fold one support leg strap around each LUCAS support leg.
- 2. Fasten the buckles on the inner side of the support leg.



4.4 Prepare the Carrying Bag

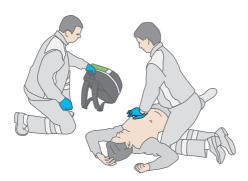


- 1. Insert a fully charged LUCAS Battery in the Battery slot in the hood of LUCAS.
- 2. Make sure that a Suction Cup is mounted correctly.
- 3. Put the Upper Part in the Carrying Bag with the hood towards the open end.
- 4. Put the external Power Supply (optional) in one of the pockets between the LUCAS support legs.
- 5. Put an extra (optional) charged LUCAS Battery in the other pocket.
- 6. Put the cushion strap of the Stabilisation Strap between the support legs.
- 7. Extra Suction Cups can be put in the side pockets close to the hood.
- 8. Position the Back Plate on top of the bag.
- 9. Close the green retaining flap.
- 10. Put the Instructions for Use (IFU) in the transparent IFU pocket in the bag
- 11. Close the bag.

5 Use LUCAS™

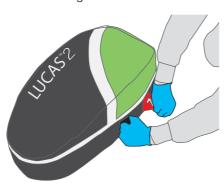
5.1 Arrival at the patient

When you have confirmed a cardiac arrest, immediately start manual cardiopulmonary resuscitation (CPR). Continue with a minimum of interruptions.

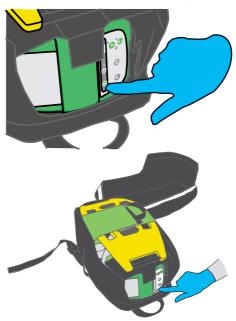


5.2 Unpack LUCAS™

- 1. Position the bag with its top nearest to you.
- 2. Put your left hand on the black strap on the left hand side and pull the red handle so that the bag unfolds.



3. Push **ON/OFF** on the User Control Panel for 1 second to power up LUCAS in the bag and start the self test. The green LED adjacent to the **ADJUST** key comes on when LUCAS is ready for use.



Note: LUCAS powers down automatically after 5 minutes if you leave it in the **ADJUST** mode.

Caution - device alarm

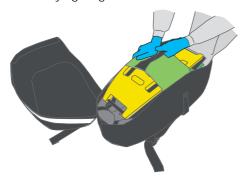
If there is a malfunction, the red Alarm LED comes on and a buzzer signal is heard. For trouble shooting, *refer to section 8.3*.

Caution - keep Battery in position
The Battery must always be in position for
LUCAS to be able to operate, even when
powered by the external Power Supply.

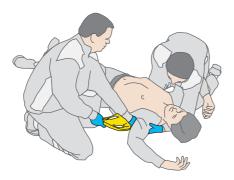


5.3 Assembly

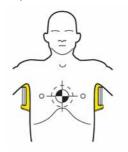
1. Remove the LUCAS Back Plate from the Carrying Bag.



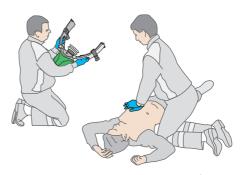
- 2. Stop manual CPR.
- Make sure that you support the patient's head.
- 4. Carefully put the LUCAS Back Plate under the patient, immediately below the arm pits. Use one of these procedures:
 - a. Hold the patient's shoulder and lift the patient's upper body slightly,
 - b. Roll the patient from side to side.

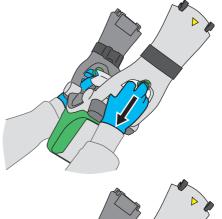


Note: Accurate positioning of the Back Plate makes it easier and faster to position the Suction Cup correctly.



- 5. Start manual CPR again.
- 6. Hold the handles on the support legs to remove the LUCAS Upper Part from the bag. Pull the release rings once to make sure that the claw locks are open.
- 7. Let go of the release rings.



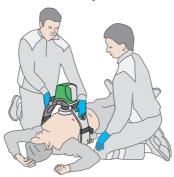




8. Attach the support leg that is nearest to you to the Back Plate.



- 9. Stop manual CPR.
- 10. Attach the other support leg to the Back Plate, so that the two support legs lock against the Back Plate. Listen for a click.
- 11. Pull upwards once to make sure that the parts are correctly attached.



Note: If the LUCAS Upper Part does not attach to the Back Plate, make sure that the claw locks are open and that you have released the release rings.

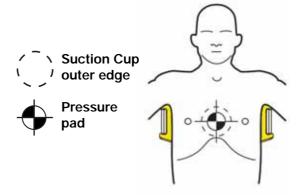
WARNING - PATIENT TOO LARGE

If the patient is too large, the Upper Part of LUCAS cannot lock to the Back Plate without compressing the patient's chest. Continue the manual compressions.

5.4 Adjustment and operation

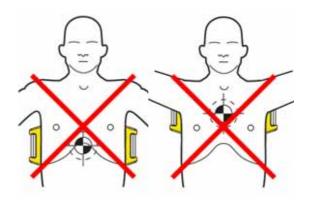
The compression point should be the same place as for manual CPR and as indicated in guidelines.

When the pressure pad in the Suction Cup is in the correct position, the lower edge of the Suction Cup is immediately over the bottom of the sternum.



WARNING - INCORRECT POSITION ON CHEST

If the pressure pad is not in the correct position relative to the sternum, there is an increased risk of damage to the rib cage and internal organs. Also, the patient's blood circulation is compromised.



1. Use your finger to make sure that the lower edge of the Suction Cup is immediately over the end of the sternum



If necessary, move the device by pulling the support legs to adjust the position.

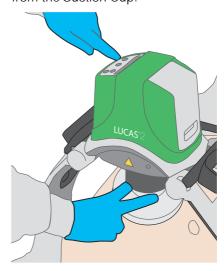


- 2. Adjust the height of the Suction Cup to set the Start Position.
 - a. Make sure that LUCAS is in the **ADJUST** mode.
 - b. Push the Suction Cup down with two fingers until the pressure pad touches the patient's chest without compressing the chest.





c. Push **PAUSE** to lock the Start Position - then remove your fingers from the Suction Cup.



- d. Check for proper position. If not, push ADJUST, pull up the Suction Cup to readjust the central position and/or height for a new Start Position. Push PAUSE.
- e. Push **ACTIVE (continuous)** OR **ACTIVE (30:2)** to start the compressions.

WARNING - UNSATISFACTORY POSITION

Start manual CPR again if it is not possible to position LUCAS safely and correctly on the patient's chest.

WARNING - PATIENT TOO SMALL

Too small patient: If you cannot enter the PAUSE mode or ACTIVE mode when the pressure pad touches the patient's chest and LUCAS alarms with 3 fast signals. Start manual compressions again.

WARNING - INCORRECT START POSITION

The patient's blood circulation is compromised if the pressure pad presses down too heavily or too lightly on the chest. Push the **ADJUST** key and adjust the height of the Suction Cup immediately.



Caution - gel on chest

If there is gel on the patient's chest (e.g. from ultrasound examination), the position of the Suction Cup can change during operation. Remove all gel before you apply the Suction Cup.

Caution - keep your fingers away
Do not put your hands or other body parts on
or under the Suction Cup when LUCAS is
working. Do not touch the claw locks,
especially when you lift the patient.

WARNING - PATIENT INJURYDo not let the patient or the device stay unattended while LUCAS is in operation.

WARNING - CHANGED POSITION DURING OPERATION

If the position of the Suction Cup changes during operation or during defibrillation, immediately push **ADJUST** and adjust the position. Always use the LUCAS Stabilisation Strap to help secure the correct position.

WARNING - MALFUNCTION

If there are interruptions, or the compressions are not sufficient, or something unusual occurs during operation: Push **ON/OFF** for 1 second to stop LUCAS and remove the device. Start manual chest compressions.

WARNING - LOW BATTERY

When the orange Battery LED shows an intermittent light, either:

- Replace the Battery with one that is charged.
- Connect the external LUCAS Power Supply.

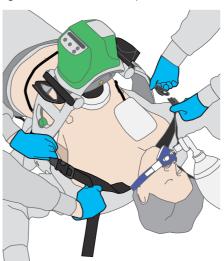
Caution - do not block the vent holes Do not obstruct the vent holes under the hood since this can cause the device to overheat.

5.5 Apply the LUCAS™ Stabilisation Strap

The LUCAS Stabilisation Strap helps secure the correct position during operation. Apply it while LUCAS is working/operational to keep interruptions to a minimum.

Caution - Stabilisation Strap application Delay the application of the LUCAS Stabilisation Strap if this prevents or delays any medical treatment of the patient.

- Remove the cushion strap, which is a part of the Stabilisation Strap, from the Carrying Bag (the support leg straps part of the Stabilisation Strap should already be attached to the support legs).
- 2. Extend the cushion strap fully at the buckles.
- 3. Carefully lift the patient's head and put the cushion behind the patient's neck. Position the cushion as near the patient's shoulders as possible.
- 4. Join the buckles on the support leg straps with the buckles on the cushion strap. Make sure that the straps are not twisted.
- 5. Hold the LUCAS support legs stable and tighten the cushion strap.



6. Make sure that the position of the Suction Cup is correct on the patient's chest.

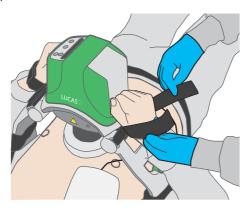
If it is not, adjust the position:

- a. Push ADJUST.
- b. Release the cushion straps from the support leg straps.
- c. Adjust the Suction Cup position (as described in the *section 5.4.2*).
- d. When the Suction Cup is in the correct position, push ACTIVE (continuous) or ACTIVE (30:2) to start the compressions again.
- e. Attach the cushion strap again. *Refer to the steps 2 to 5 above.*

5.6 Move the patient

5.6.1 Secure the patient's arms

When you move the patient, you can secure the patient's arms with the Patient Straps on the LUCAS. This makes it easier to move the patient.



Caution - do not use the Patient Straps for lifting

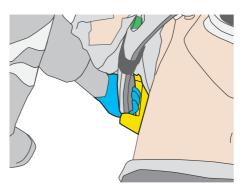
Do not use the Patient Straps to lift the patient. The straps are only to attach the patient's arms to LUCAS.

Caution - IV access

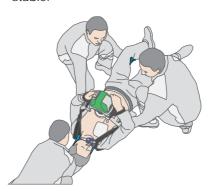
Make sure that IV access is not obstructed.

5.6.2 Prepare to lift the patient

- 1. Make a decision about what equipment you will move and where to put the transportation device.
- 2. Those at the patient's side should:
 - a. put one hand below the claw locks under the support leg



- b. with the other hand, hold the patient's belt, trousers or under the thigh
- 3. Make sure that the patient's head is stable.



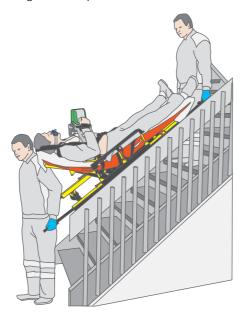
5.6.3 Lift the patient

- Push PAUSE to temporarily stop the compressions.
- 2. Lift and move the patient to a stretcher or other transportation device (backboard, vacuum mattress or similar).
- 3. Make sure that the Suction Cup is in the correct position on the patient's chest.
- Push ACTIVE (continuous) or ACTIVE (30:2) to start the compressions again.

5.6.4 Move the patient

LUCAS can continue working while you move the patient if:

- LUCAS and the patient are safely positioned on the transportation device
- LUCAS stays in the correct position and angle on the patient's chest



If necessary, adjust the position of the Suction Cup.

WARNING - CHANGED POSITION DURING OPERATION

If the position of the Suction Cup changes during operation or during defibrillation, immediately push **ADJUST** and adjust the position. Always use the LUCAS Stabilisation Strap to help secure the correct position.

5.7 Replace the Power Supply during operation

When the Battery is low, LUCAS shows an intermittent orange LED and there is an audible alarm signal.



5.7.1 Change the Battery

Keep interruptions to a minimum while changing the Battery.

Note: To minimise interruptions, we recommend to always have a spare, charged, LUCAS Battery in the Carrying Bag.

- 1. Push **PAUSE** to temporarily stop the compressions.
- 2. Pull the Battery out and then upwards to remove it.



- 3. Install a fully-charged LUCAS Battery. Put it in from above.
- 4. Wait until the green PAUSE mode LED comes on.
- 5. Push ACTIVE (continuous) or ACTIVE (30:2) to start the chest compressions again. The LUCAS Smart Restart feature remembers the settings and Start Position for 60 seconds.

Note: If the Battery change takes more than 60 seconds, LUCAS does a self test and you must adjust the Start Position again.



5.7.2 Connect to the external Power Supply

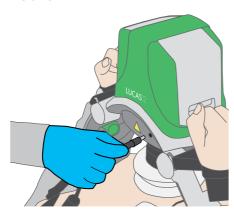
You can connect the LUCAS Power Supply or Car Power Cable when the LUCAS is in all operating modes.

Caution - keep Battery in position

The Battery must always be in position for LUCAS to be able to operate, even when powered by the external Power Supply.

To use the Power Supply cable:

• Connect the Power Supply cable to LUCAS.



 Plug the mains cable in to the wall mains socket (100-240V, 50/60Hz)

To use the Car Power Cable:

- Connect the Car Power Cable to LUCAS
- Plug the Car Power Cable in to the car power socket (12-24V DC)

5.8 Adjunctive therapies

Caution - adjunctive therapies

The use of other medical equipment or drugs in conjunction with LUCAS can affect the treatment. Always consult the instructions for use of the other equipment and/or drugs to make sure that they are appropriate in conjunction with CPR.

5.8.1 Defibrillation

Defibrillation can be performed while LUCAS operates.

- 1. You can apply the defibrillation electrodes before or after LUCAS has been put in position.
- 2. Perform the defibrillation according to the defibrillator manufacurer's instructions.

Caution - defibrillation electrodes

Position the defibrillation electrodes and wires so that they are not under the Suction Cup. If there are already electrodes on the patient, make sure that they are not where the Suction Cup will be positioned. If they are, you must apply new electrodes.

3. After defibrillation, make sure that the position of the Suction Cup is correct. If necessary, adjust the position.



WARNING - CHANGED POSITION DURING OPERATION

If the position of the Suction Cup changes during operation or during defibrillation, immediately push **ADJUST** and adjust the position. Always use the LUCAS Stabilisation Strap to help secure the correct position.

WARNING - ECG INTERFERENCE

Chest compressions interfere with ECG analysis. Push **PAUSE** before you start the ECG analysis. Make the interruption as short as possible. Push **ACTIVE** (continuous) or **ACTIVE** (30:2) to start the compressions again.

5.8.2 Ventilation

Always follow local and/or international quidelines for ventilation.

LUCAS can operate in two different modes:

ACTIVE (continuous)

When you push this key LUCAS performs continuous compressions. The green LED signal will blink 8 times per minute to indicate when to ventilate during ongoing compressions.

ACTIVE (30:2)

When you push this key, LUCAS performs 30 chest compressions and then temporarily stops for 3 seconds. During the stop, the operator can perform 2 ventilations. After the stop the cycle starts again. An intermittent LED in combination with an alarm signal will alert the operator before each ventilation pause.

5.8.3 Use in the catheterisation laboratory

LUCAS can be used in the catheterisation laboratory. Except for the compression mechanism it is mainly radiotranslucent and allows most X-ray procedures to be performed.

5.9 Remove LUCAS™ from the patient

- Push **ON/OFF** for 1 second to turn off the device.
- 2. If a LUCAS Stabilisation Strap is attached to LUCAS, remove the cushion strap, which is part of the Stabilisation Strap, from the support leg straps.
- 3. Pull the release rings to release the Upper Part from the Back Plate.
- 4. If the patient's condition allows it, remove the Back Plate.

6 Care after use and preparation for next use

After each use of the LUCAS Chest Compression System you must:

- 1. Remove the Suction Cup (*refer to section 6.2*).
- 2. If necessary, remove and clean the Patient Straps and the Stabilisation Strap separately (*refer to section 6.1 and 6.3*).
- 3. Clean the device and let it dry (refer to section 6.1).

Preparation for next use:

- Replace the used Battery in the battery slot of the hood with a fully charged Battery.
- 5. Mount a new Suction Cup.
- 6. Attach the Patient Straps again, if they have been removed.
- 7. Attach the support leg straps of the LUCAS Stabilisation Strap again, if they have been removed.
- 8. Pack the device into the Carrying Bag:
 - Put the Upper Part in the Carrying Bag with the hood towards the open end.
 - Put the external Power Supply (optional) in one of the pockets between the LUCAS support legs.
 - Put an extra (optional) charged LUCAS Battery in the other pocket.
 - Put the cushion strap part of the Stabilisation Strap between the support legs.
 - Extra Suction Cups can be put in the side pockets near the hood.
 - Position the Back Plate on top of the bag.
 - Close the green inner lock.
 - Put the Instructions for Use (IFU) in the transparent IFU pocket in the bag.
- 9. Close the bag.

Do routine checks weekly and after each use (refer to the maintenance section).



6.1 Cleaning routines

Clean all surfaces and straps with a soft cloth and warm water containing a mild cleaning agent or disinfectant, e.g.

- 70% isopropyl alcohol solution
- 45% isopropyl alcohol with added detergent
- · Quaternary ammonium compound

Follow the handling instructions from the manufacturer of the disinfectant.

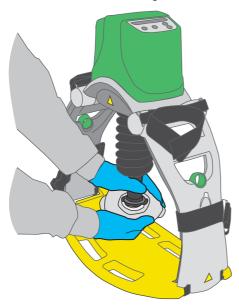
Caution - liquid

Do not immerse LUCAS in liquid. The device can be damaged if liquid enters the hood.

Allow LUCAS to dry before you pack it into the bag.

6.2 Remove and install the Suction Cup

- Pull the Suction Cup off the black mounting tube.
- Discard the Suction Cup as contaminated medical waste.
- Bend a new Suction Cup onto the black mounting tube.
- Make sure the Suction Cup is safely attached to the mounting tube.



6.3 Remove and attach the Patient Straps

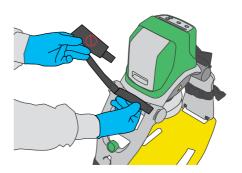
Remove:

1. Open the Patient Straps and pull them out from the metal rings on the LUCAS support legs.

Clean according to 6.1.

Install:

- 1. Thread the Patient Straps through the metal holder on the LUCAS support legs.
- 2. Fold the Patient Strap so that the symbol is visible.
- 3. Press the strap parts firmly together.



6.4 Remove and attach the LUCAS™ Stabilisation Strap

Remove the Support leg straps, which are part of the Stabilisation Strap, by opening the buckles.

Clean the Stabilisation Strap according to 6.1.

Install according to 4.3.

6.5 Remove and recharge the Battery

- Replace the Battery with a fully charged one.
- 2. Recharge the used Battery for future use

You can charge the LUCAS Battery in two ways:

- In the external LUCAS Battery Charger (optional)
 - put the Battery in the slot on the Battery Charger,
 - plug the Battery Charger power cable in to the mains wall socket.
- Installed in LUCAS:
 - put the Battery in the slot in the hood of LUCAS,
 - connect the Power Supply/Car Power Cable to the DC input on the side of LUCAS,
 - plug the Power Supply in to the mains wall socket.



Green LEDs indicate a fully charged Battery.

Caution - keep Battery in position
The Battery must always be in position for
LUCAS to be able to operate, even when
powered by the external Power Supply.

Caution - use only approved accessories

Use only JOLIFE AB-approved accessories with LUCAS. LUCAS does not operate correctly if you use accessories that are not approved. Use only LUCAS Batteries and the LUCAS Power Supply that are designed for LUCAS. If you use other batteries or another Power Supply you can cause permanent damage to LUCAS. This will also void the warranty.

7 Maintenance

7.1 Routine checks

Weekly, and after each use of the LUCAS Chest Compression System, you should:

- 1. Make sure that the device is clean.
- 2. Make sure that a new Suction Cup is fitted.
- 3. Make sure that the Patient Straps are attached.
- Make sure that the two support leg straps of the Stabilisation Strap are attached around the support legs.
- 5. Pull the release rings upwards to make sure that the claw locks are open.
- 6. Make sure that the Battery is fully charged. When LUCAS is in the OFF mode, push **MUTE**. The Battery indicator comes on and shows the Battery charge status (see section 8.1).
- Push ON/OFF to make LUCAS do a self test. Make sure the ADJUST LED illuminates without any alarm or warning LED
- 8. Push **ON/OFF** to turn off LUCAS again.



8 Troubleshooting

8.1 Indicators and warnings during normal operation

Refer to the table below to find the reason for audible and/or LED alarms during normal operation.

Situation	Visual LEI	D indication	Audible signals	User action
LUCAS is in the ON mode and there is more than 90% Battery capacity remaining.	-000	Fully charged Battery: All 3 green Battery indi- cator LEDs show a constant light.	None	None
LUCAS is in the ON mode and there is more than 60% and less than 90% Battery capacity remaining.	•	Battery 2/3 charged: The 2 right hand green Battery indicator LEDs show a constant light.	None	None
LUCAS is in the ON mode and there is more than 30% and less than 60% Battery capacity remaining.	ш	Battery 1/3 charged: The right hand green Battery indicator LED shows a constant light.	None	None
LUCAS is in the ON mode and there is less than 30% Battery capacity remaining (approximately 10 minutes of operating capacity).		Low Battery: The right hand orange Battery indicator LED comes on intermittent- ly.	Intermittent alarm	Replace the Battery or connect to the external power supply.
An external LUCAS Power Supply is connected and charging the Battery.	4	Charging Battery: The 3 green Battery indicator LEDs show a "running" light.	None	None
An external LUCAS Power Supply is connected and the Battery is fully charged.	-000	Fully charged Battery: All 3 green Battery indi- cator LEDs show a constant light.	None	None
The Battery has been used more than 200 times with compressions of more than 10 minutes each or is older than 3 years.		End of Battery service life: The right hand Battery indicator LED shows an orange light instead of green, in all the above situations.	None	Dispose of Battery.
In the ADJUST mode.		The ADJUST LED shows a green light.	None	None
In the PAUSE mode.	0	The PAUSE LED shows a green light.	None	None

Situation	Visual LE	D indication	Audible signals	User action
In the ACTIVE (continuous) mode		The ACTIVE (continuous) Key, LUCAS performs continuous chest com- pressions. The green LED signal will blink 8 rimes per minute	None	This is to indicate when to ventilate during going compressions.
In the ACTIVE (30:2) mode	30:2	The ACTIVE (30:2) LED shows an intermittant green light during compressions number 26, 27, 28, 29 and 30.	Alarm signal alert during compressions number 28 ("ding"), 29 ("ding") and 30 ("dong").	This is to alert the operator to ventilate the patient when LUCAS temporarily stops the compressions at number 30.
Too small a patient. Trying to enter the PAUSE mode or any of the AC- TIVE modes when the Suction Cup is in a lower position than for the mini- mum patient size (sternum height approximately 17 cm)	None		3 fast signals	Continue with manual compressions.

8.2 Battery replacement and Smart Restart feature

If you change the Battery quickly in 60 seconds or less, with LUCAS in the ON mode, the LUCAS Smart Restart feature remembers the settings and Start Position according to the table below. If the Battery change takes more than 60 seconds, LUCAS does a self test and you must adjust the Start Position again.

Mode when you remove the Battery	Mode when the new Battery is in place again
PAUSE	PAUSE (with the same Start Position)
ACTIVE (continuous)	PAUSE (with the same Start Position)
ACTIVE (30:2)	PAUSE (with the same Start Position)
ADJUST	ADJUST
OFF	OFF



8.3 Malfunction alarms

Below is a list of all alarms that can occur on LUCAS. You mute all alarms for 60 seconds if you push **MUTE**.

Start with manual compressions immediately if LUCAS does not operate properly.

Reason	Visual LEI	O indication	Audible alarms	Result
Compression pattern outside limit (too deep, too shallow and timing failure)		Red alarm LED	Alarm	Compressions stop
Rising temperature in LUCAS			Warning alarm	None
Too high temperature in LUCAS		Red alarm LED	Alarm	Compressions stop
Hardware error		Red alarm LED	Alarm	Compressions stop
Too high Battery temperature		Intermittent red Battery warning: The red Battery indicator LED farthest to the right illuminates intermittently.	Intermittent alarm	Compressions stop
Battery charge too low		Intermittent red Battery warning: The red Battery indicator LED farthest to the right illuminates intermittently.	Intermittent alarm	Compressions stop. The Battery must be recharged in the external Battery Charger.
Battery error		Constant red Battery warning: The red Battery indicator LED farthest to the right shows a constant light.	Alarm	Compressions stop. The Battery cannot be used anymore.

If the malfunction described above seems permanent, LUCAS must be examined by approved service personnel. Please consult your local LUCAS representative or JOLIFE AB. Contact information is available at www.JOLIFE.com

9 Technical specifications

All specifications in this chapter apply to the LUCAS $^{\text{TM}}2$ Chest Compression System.

9.1 Patient parameters

Category	Specifications
Patients eligible for treatment:	Adult patients who fit into the device; • sternum height of 6.7 to 11.9 inches / 170 to 303 mm • a maximum chest width of 17.7 inches / 449 mm The use of LUCAS is not restricted by patient weight.

9.2 Compression parameters

Category	Specifications	
Compression depth	1.5 to 2.0 inches / 4 to 5 cm from the Start Position	
Compression frequency	100 ± 5 compressions per minute	
Compression duty cycle	50 ± 5%	
Compression modes (operator selectable)	 30:2 (30 compressions followed by a 3 seconds ventilation pause) Continuous compressions 	

9.3 Device physical specifications

Category	Specifications
Dimensions when assembled (H \times W \times D)	22.4 x 20.5 x 9.4 inches / 57 × 52 × 24 cm
Dimensions Carrying Bag with device inside (H \times W \times D)	25.6 x 13 x 9.8 inches / 65 × 33 × 25 cm
Weight of the device with the Battery	17.2 lbs / 7.8 kg

9.4 Device environmental specifications

Category	Specifications	
Operating temperature	+32°F to +104°F / +0°C to +40°C	
Storage temperature	-4°F to +158°F / -20°C to +70°C	
Relative humidity	5% to 98%, non-condensing	
IP classification (IEC60529)	IP 43	
Operating input voltage	12-24 V DC	



9.5 Battery physical specifications

Category	Specifications
Size (H × W × D)	5.1 x 3.5 x 2.2 inches / 13.0 × 8.8 × 5.7 cm
Weight	1.3 lbs / 0.6 kg
Туре	Rechargeable Lithium Ion Polymer (LiPo)
Capacity	3300 mAh (typical), 86 Wh
Battery voltage (nominal)	25.9 V
Initial Battery runtime (nominal patient)	45 minutes (typical)
Maximum Battery charge time	Less than 4 hours at room temperature (72°F / 22°C)
Required interval for replacement of the Battery	Battery replacement is recommended every 3 years or after 200 uses (of more than 10 minutes use each time)

9.6 Battery environmental specifications

Category	Specifications
Operating temperature	32°F to 104°F / 0°C to +40°C, ambient when installed in the device
Charge temperature	41°F to 95°F / 5°C to +35°C ambient (68°F to 77°F / 20°C to 25°C preferred)
Storage temperature	32°F to 104°F / 0°C to +40°C ambient for less than six months
IP classification (IEC60529)	IP 44

9.7 Electromagnetic environmental declaration

Guidance and manufacturer's declaration - electromagnetic emissions

LUCAS 2 is intended for use in the electromagnetic environment specified below. The customer or the operator of LUCAS 2 must make sure that it is used in the correct environment.

Emissions test	Compliance	Electromagnetic environment - guidance
RF emissions CISPR 11	Group 1	LUCAS 2 uses RF energy only for its internal operation. This makes its RF emissions very low and not likely to cause interference with other electronic equipment near LUCAS 2.
RF emissions CISPR 11	Class B	LUCAS 2 is suitable for use in all
Harmonic emissions IEC 61000-3-2	Class A	buildings including the home and places directly connected to the public
Voltage fluctuations/ flicker emissions IEC 61000-3-3	Complies	low-voltage Power Supply network that supplies buildings used for domestic purposes.

Guidance and manufacturer's declaration - electromagnetic immunity

LUCAS 2 is intended for use in the electromagnetic environment specified below. The customer or the operator of LUCAS 2 must make sure that it is used in the correct environment.

Immunity test	IEC 60601 test level	Compliance level	Electromagnetic environment - guidance
Electrostatic discharge (ESD) IEC 61000-4-2	+/- 6 kV contact +/- 8 kV air	+/- 6 kV contact +/- 8 kV air	Floors must be wood, concrete or ceramic tile. If there is synthetic material on the floor, the relative humidity must be 30% or more.
Electrical fast transient / Burst IEC 61000-4-4	+/- 2 kV for Power Supply lines +/- 1 kV for input/output lines	+/- 2 kV for Power Supply lines n/a. for input/output lines	The mains power quality must be that of a typical commercial or hospital environment.
Surge IEC 61000-4-5	+/- 1 kV differential mode +/- 2 kV common mode	+/- 1 kV differential mode n/a. for common mode	The mains power quality must be that of a typical commercial or hospital environment.
Voltage dips, short in- terruptions and voltage variations on Power Supply input lines IEC 61000-4-11	<5 % UT (>95 % dip in UT) for 0.5 cycle 40 % UT (60 % dip in UT) for 5 cycles 70 % UT (30 % dip in UT) for 25 cycles <5 % UT (>95 % dip in UT)) for 5 sec	<5 % UT (>95 % dip in UT) for 0.5 cycle 40 % UT (60 % dip in UT) for 5 cycles 70 % UT (30 % dip in UT) for 25 cycles <5 % UT (>95 % dip in UT)) for 5 sec	The mains power quality must be that of a typical commercial or hospital environment. If the user of the [Equipment or System] requires continued operation during power mains interruptions, JOLIFE recommends that the [Equipment or System] is energized from a Power Supply or Battery that cannot be interrupted.
Power frequency (50/60 Hz) magnetic field IEC 61000-4-8	30 A/m	30 A/m	The power frequency magnetic fields must be at levels that are characteristic of a typical location in a typical commercial or hospital environment

NOTE UT is the a.c. mains voltage prior to application of the test level.

Guidance and manufacturer's declaration - electromagnetic immunity

LUCAS 2 is intended for use in the electromagnetic environment specified below. The customer or the operator of LUCAS 2 must make sure that it is used in the correct environment.



Immunity	IEC 60601	Compli-	Electromagnetic environment - guidance
test	test level	ance level	
			Portable and mobile RF communications equipment must not be used closer to LUCAS 2 (cables included) than the recommended separation distance calculated with the equation applicable to the frequency of the transmitter.
			Recommended separation distance
Conducted RF IEC 61000-4-6	10 Vrms 150 kHz to 80 MHz	10 Vrms	$d = 1.2\sqrt{P}$
Radiated RF IEC 61000-4-3	10 V/m 80MHz to 2.5GHz	10 V/m	$d = 1.2\sqrt{P}$ $d = 1.3\sqrt{P}$
			where P is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer and d is the recommended separation distance in metres (m).
			Field strengths from fixed RF transmitters, as determined by an electromagnetic site survey, ^a must be less than the compliance level in each frequency range. ^b
			Interference can occur near equipment marked with the following symbol.
			$((\overset{\bullet}{\blacktriangle}))$

NOTE 1 At 80MHz and 800MHz, the higher frequency range applies.

NOTE 2 These guidelines may not apply in some situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.

Recommended separation distances between portable and mobile RF communications equipment and the LUCAS $2\,$

LUCAS 2 is intended for use in an electromagnetic environment in which radiated RF disturbances are controlled. The customer or the operator of LUCAS 2 can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and LUCAS 2 as recommended below, according to the maximum output power of the communications equipment.

^a Field strengths from fixed transmitters, such as base stations for radio (mobile/cordless) telephones and land mobile radios, amateur radio, AM and FM radio and TV broadcasting cannot be predicted theoretically with accuracy. To assess the electromagnetic environment due to fixed RF transmitters, an electromagnetic site survey should be considered. If the measured field strength in the location in which LUCAS 2 is used exceeds the applicable RF compliance level above, LUCAS 2 should be observed to make sure it operates normally. If unusual or incorrect performance is observed, additional measures can be necessary, such as reorientating or relocating LUCAS 2.

^b Over the frequency range 150 kHz to 80 MHz, field strengths should be less than 10 V/m.

Rated maximum output	Separation distance according to frequency of transmitter		
power of transmitter W	nsmitter 150 kHz to 80 MHz 80 MHz to 800 MHz		800 MHz to 2.5 GHz
	$d = 1.2\sqrt{P}$	$d = 1.2\sqrt{P}$	$d = 1.3\sqrt{P}$
0.01	0.12	0.12	0.24
0.1	0.38	0.38	0.73
1	1.2	1.2	2.3
10	3.8	3.8	7.3
100	12	12	23

For transmitters rated at a maximum output power not listed above, the recommended separation distance d in metres (m) can be estimated using the equation applicable to the frequency of the transmitter, where P is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer.

Note 1: At 80 MHz and 800 MHz, the separation distance for the higher frequency range applies.

Note 2: These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.

9.8 Limited warranty

Subject to the limitations and exclusions set forth below, JOLIFE AB ("JOLIFE") guarantees that JOLIFE products which are purchased from authorized JOLIFE representatives or dealers and are used in accordance with their instructions will be free from defects in material and workmanship appearing under normal service and use for the time period listed below. The time limit and the warranty schedule begin on the date of delivery to the first purchaser.

12 Months: LUCAS™2 Chest Compression System (including the LUCAS device (Upper Part and Back Plate), Carrying Bag, Battery, Stabilisation Strap, Patient Straps).

JOLIFE does not guarantee that JOLIFE products will perform error-free or without interruptions. The sole and exclusive remedy under this limited warranty is to repair or replace defective material or workmanship at the option of JOLIFE. To qualify for the repair or replacement, the product must not have been repaired or altered in any way which, in the judgment of JOLIFE, affects its stability and reliability. The product must have been used and maintained in accordance with relevant operating instructions and in the intended environment or setting.

The Limited Warranty does not cover problems with products that have been caused by misuse, abuse, improper maintenance, modifications to the product or accident. JOLIFE or its authorized service provider shall, at its sole discretion, determine whether a reported problem is covered under this Limited Warranty and whether the product is field serviceable. If field serviceable and located within 100 miles of a JOLIFE designated service location, warranty service will be provided by JOLIFE or its authorized service provider at the purchaser's facility during normal business hours. If not field serviceable or if the product is located outside of such areas, all products requiring warranty service should be returned to a location designated by JOLIFE or its authorized service provider, freight prepaid, and must be accompanied by a written, detailed explanation of the claimed failure.

Except for the Limited Warranty provided above, NEITHER JOLIFE NOR ITS AUTHORIZED SERVICE PROVIDER MAKES ANY WARRANTY, EXPRESS OR IMPLIED, INCLUDING, BUT NOT LIMITED TO, ANY IMPLIED WARRANTY OF MERCHANTABILITY OR FITNESS FOR A PARTICULAR PURPOSE, WHETHER ARISING FROM STATUTE, COMMON LAW, CUSTOMER OR OTHERWISE. THIS LIMITED WARRANTY SHALL BE THE EXCLUSIVE REMEDY AVAILABLE TO ANY PERSON OR ENTITY. NEITHER JOLIFE NOR ITS AUTHORIZED SERVICE PROVIDER IS LIABLE FOR DIRECT OR INDIRECT, SPECIAL, INCIDENTAL OR CONSEQUENTIAL DAMAGES (INCLUDING LOSS OF BUSINESS OR PROFITS) WHETHER BASED ON CONTRACT, TORT, OR ANY OTHER LEGAL THEORY.

Any legal action arising from the purchase or use of JOLIFE products shall be commenced within one year from the accrual of the cause of action, or be barred forever. In no event shall JOLIFE liability under this warranty or otherwise exceed the greater of \$50,000 or the purchase price of the product giving rise to the cause of action.

Products are guaranteed in compliance with applicable law. If any part or term of this Limited Warranty is held to be illegal, unenforceable or in conflict with applicable law by any court of competent jurisdiction, the validity of the remaining portions of the Limited Warranty shall not be affected, and all rights and obligations shall be construed and enforced as if this Limited Warranty did not contain the particular part or term held to be invalid. Some countries, and states within United States of America, do not allow the exclusion or limitation of incidental or consequential damages, so the above limitation or exclusion may not apply to you. This Limited Warranty gives the user specific support legal rights. The user may also have other rights which vary from state to state or country to country.

Appendix A; LUCAS™2 parts and accessories

Description	JOLIFE AB Part number
LUCAS Back Plate	150208-00
3 x LUCAS 2 Suction Cup	150205-03
LUCAS 2 Carrying Bag	150200-00
LUCAS 2 Instructions for Use (regional versions)	100666-XX
LUCAS 2 Battery	150201-00
LUCAS Stabilisation Strap	150203-00
LUCAS Patient Straps	150204-00
LUCAS 2 Power Supply (regional versions)	150202-XX
LUCAS 2 Car Power Cable	150206-00
LUCAS 2 Battery Charger	150207-00
LUCAS 2 Back Plate Grip Tape	150209-00









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jolife.com

Chest Compression System

UCAS[®]2 Quick Reference Guide

NOTE: This guide is not a complete Instructions for Use. Refer to the "Instructions for Use" for complete directions for use, indications, contraindications, warnings, precautions, and potential adverse events.













Confirm cardiac arrest and start manual CPR with a minimum of interruptions until LUCAS is applied and ready.



Activate (A)

Push **ON/OFF** for 1 second to start self-test and power up **LUCAS**

5 Push down the **Suction Cup**

- Push the Suction Cup down with two fingers (make sure it is in the **ADJUST** mode)
- Pressure pad inside Suction Cup should touch patient's chest. If the pad does not touch or fit properly, continue manual compressions
- Push **PAUSE** to lock Start Position – then remove your fingers from the Suction Cup







Back Plate (B)

- Pause manual CPR
- Carefully put Back Plate under the patient, below armpits
- Resume manual CPR



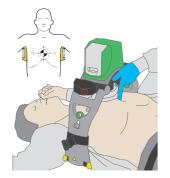
Compressor (C)

- Pull release rings once; claw locks open. Then let go of the release rings
- Attach to Back Plate; listen for "click"
- Pull up once to ensure attachment

Start compressions

- Check for proper position. Adjust if necessary
- Push ACTIVE (continuous) or **ACTIVE (30:2)**
- LUCAS provides compressions with a rate of 100 per minute and 1.5 to 2 inches depth





Position the Suction Cup

- Center the Suction Cup over the chest
- The lower edge of Suction Cup should be immediately above the end of the sternum

LUCAS Stabilization Strap

• Attach the LUCAS Stabilization Strap



Always follow local and/or international guidelines for CPR when you use LUCAS.