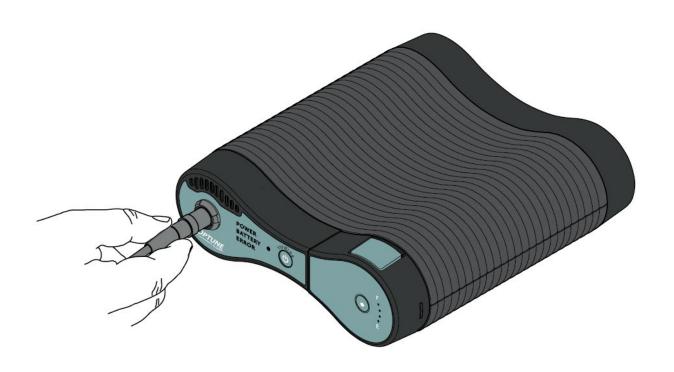




For Non Small Cell Lung Cancer

User Manual



Model Number: TFT9200

Ref Number:TFT9201EU

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	anual is intended for patients receiving TTFields treatment using the Optune Lua ent kit and the ILE Transducer Arrays.

1. ABOUT THE OPTUNE LUA TREATMENT KIT AND THE ILE TRANSDUCER ARRAYS

1.1. DEVICE DESCRIPTION

The Optune Lua treatment kit is a portable device. It produces electric fields, called tumor treating fields ("TTFields"). ILE Transducer Arrays connected to the device deliver TTFields to the chest. The TTFields are intended to destroy lung cancer cells.

The device is intended for home treatment of at least 12 hours a day on a monthly average. The Optune Lua treatment kit refers to the Electric Field Generator (Optune Lua, the device), connection cable, power supply, battery, battery charger and ILE Transducer Arrays.

1.2. INTENDED PURPOSE

The Optune Lua treatment kit is indicated for the treatment of stage IV, non-squamous, non-small cell lung cancer, in combination with Pemetrexed (Alimta), after failure of first line treatments.

The treatment is intended for adult patients, 18 years of age or older, and should be started more than four weeks after the latest surgery, radiation therapy or chemotherapy.

The device is intended for home treatment of at least 12 hours a day and should be used until disease progression in the chest or upper abdomen

1.3. CONTRAINDICATIONS, WARNINGS, PRECAUTIONS AND NOTICES

Contraindications

Do not use the Optune Lua treatment kit if you have an active implanted medical device. Examples of active electronic devices include deep brain stimulators, spinal cord stimulators, vagus nerve stimulators, pacemakers and defibrillators. Use of the Optune Lua treatment kit together with implanted electronic devices have has not been tested and may lead to malfunctioning of the implanted devices.

Do not use the Optune Lua treatment kit if you are known to be sensitive to conductive hydrogels like the gel used on electrocardiogram (ECG) stickers or transcutaneous electrical nerve stimulation (TENS) electrodes. In this case, skin contact with the gel used

with the Optune Lua treatment kit may commonly cause increased redness and itching, and rarely may even lead to severe allergic reactions such as shock and respiratory failure.

Do not use the Optune Lua treatment kit if you have clinically significant hepatic or renal disease.

Do not use the Optune Lua treatment kit if you have significant additional neurological disease (primary seizure disorder, dementia, progressive degenerative neurological disorder, meningitis or encephalitis, hydrocephalus associated with increased intracranial pressure)

Do not use the Optune Lua treatment kit if you have one of the following conditions: Congestive heart failure, angina pectoris that is not medically controlled, history of myocardial infarction 1 year from start of device treatment, uncontrolled hypertension or arrhythmias, active infection requiring i.v. antibiotics, unstable diabetes mellitus or other contraindication for corticosteroid therapy and active ulcer.

Warnings

Warning - Use the Optune Lua treatment kit only after receiving training from qualified personnel, such as your doctor, a nurse, or other medical personnel who have completed a training course given by the device manufacturer (Novocure GmbH Switzerland). Ask to see a certificate signed by Novocure that says they completed a training course. Your training will include a detailed review of this manual and practice in the use of the treatment kit. In addition, you will be trained in what to do if there are problems with treatment. Use of the Optune Lua treatment kit without receiving this training can result in breaks in treatment and may rarely cause increased skin rash, open sores on your body, allergic reactions or even an electric shock.

Warning - In case of skin irritation, which appears as redness under the transducer arrays (a mild rash), talk to your physician before starting any treatment for skin irritation. Your physician may recommend using over-the-counter topical steroids when replacing the transducer arrays. This will help relieve your skin irritation. If you do not use this cream, the skin irritation can become more serious and may even lead to skin break down, infections, pain and blisters. If this happens, stop using the topical steroid cream and contact your doctor. Your doctor will supply you with an antibiotic cream to use when replacing the transducer arrays. If you do not use this cream, your symptoms may continue and your doctor may ask you to take a break from treatment until your skin heals. Taking a break from treatment may lower your chance to respond to treatment.

Warning - All servicing procedures must be performed by qualified and trained personnel. If you attempt to open and service the treatment kit alone you may cause damage to the

treatment kit. You could also get an electric shock by touching the inner parts of the device.

Warning - No modification of this equipment is allowed. – Do not use the Optune Lua device with applied parts or accessories other than those described in this manual.

Precautions

Caution – Do not use any parts that do not come with the Optune Lua treatment kit, or that were not sent to you by the device manufacturer or given to you by your doctor. Use of other parts, manufactured by other companies or for use with other devices, can damage the device. This may lead to a break in treatment.

Caution - Do not use the Optune Lua treatment kit if any parts look damaged (torn wires, loose connectors, loose sockets, cracks or breaks in the plastic case). Use of damaged components can damage the device and cause a break in treatment.

Caution - Do not wet the Optune Lua device or transducer arrays or use in the shower or in heavy rain. Getting the device wet may damage it, preventing you from receiving treatment for the right amount of time. Getting the transducer arrays very wet is likely to cause the transducer arrays to come loose from your skin. If this happens, the device will turn off and you will need to change the transducer arrays.

Caution - Before connecting or disconnecting the transducer arrays, make sure that the Optune Lua power switch is in the OFF position.

Caution - Do not use the Optune Lua treatment kit if you are pregnant, think you might be pregnant, or are trying to get pregnant. If you are a woman who is able to get pregnant, you must use birth control when using the device. The Optune Lua treatment kit was not tested in pregnant women. It is unknown what side effects the device may cause if you are pregnant or if it will be effective.

Caution -Connection cable may pose a hazard of strangulation. Avoid wearing the connection cable around your neck.

Caution – do not place the power supply to make it difficult to disconnect the wall plugin from the wall socket.

Caution – Covering the power supply may result in the power supply overheating.

Notices

Notice! The Optune Lua treatment kit and the transducer arrays will activate metal detectors.

Notice! You should use the Optune Lua treatment kit for at least 12 hours a day to get the best response to treatment. Using the Optune Lua treatment kit for less than 12 hours a day lowers the chances that you will respond to treatment.

Notice! Do not stop using the Optune Lua treatment kit even if you have used it less than the recommended 12 hours per day. You should stop using the device only if your doctor tells you to. Stopping treatment could lower the chances that you will respond to treatment.

Notice! If you plan to be away from home for more than 1 hour, carry an extra battery and/or the power supply with you in case the battery you are using runs out. If you do not take an extra battery and/or the power supply, you may have a break in your treatment. Breaks in treatment may lower your chance to respond to the treatment.

Notice - Make sure you have at least 12 extra transducer arrays at all times. This will last you until the next transducer array shipment arrives. Remember to order more transducer arrays when there are at least 12 extra transducer arrays left. If you do not order transducer arrays in time, you may have a break in your treatment.

Notice! Batteries may weaken over time and need to be replaced. You will know this has happened when the amount of time the device can run on a fully charged battery begins to shorten. For example, if the low battery indicator light flashes within only 1 hour from the start of treatment, replace the battery. If you do not have replacement batteries when your batteries run out, you may have a break in your treatment.

Notice - You should carry the Troubleshooting Guide (Section 12 of the patient user manual) at all times. This guide is necessary to ensure the Optune Lua treatment kit works properly. If you do not operate the treatment kit correctly, you may have a break in your treatment.

Notice! Do not block the device vents located on the front and back of the Optune Lua device. Blocking the vents may cause the device to overheat and turn off, leading to a

break in treatment. If this happens, unblock the vents, wait 5 minutes and restart the device. In case the vents are blocked with pet hair or dust, return the device for service.

Notice! Do not block the battery charger vents located on the left and right sides of the battery charger. Blocking the vents may cause the charger to overheat. This could prevent your batteries from charging. If the vents become blocked with pet hair or dust, return the charger for service.

Notice - The transducer arrays are for single use and should not be taken off your body and put back on again. If you put a used transducer array back on your chest again, it may not stick well to your skin and the device could turn off.

Notice - Keep the Optune Lua treatment kit out of the reach of children and pets.

Notice - The device has a cord that may cause tripping when connected to an electric socket.

2. CLINICAL BENEFIT AND CLINICAL EVIDENCE

Expected clinical benefit for the patient

Patients using Optune Lua together with cancer drugs lived longer compared to patients who used cancer drugs alone.

Median progression free survival of advanced (stage IV) NSCLC patients treated with the Optune Lua device together with Pemetrexed, following at least one line of prior chemotherapy, was more than double that of the expected median with Pemetrexed alone based on comparison with historical control data. Also, a statistically significant increase in time to disease progression was seen (median overall survival of 13.8 months compared to 8.3 months in historical control patients). The one-year survival rate was 57% compared to the historical control of 30% reported to Pemetrexed alone.

Clinical experience — a multi-center clinical trial has shown that Optune Lua (formerly NovoTTF-100L) treatment together with standard chemotherapy (Pemetrexed) was well tolerated with no device related serious adverse events seen in any of the 42 patients treated with an average follow up of 6 months. No cardiac or other electric field based serious adverse events were seen in any of the patients. No increase in chemotherapy related toxicity was seen. Compliance with treatment was very high with 85% of patients receiving treatment on average 12 hours a day.

Mild to moderate contact dermatitis appeared beneath the transducer array gel in all patients during treatment, which was manifested by a red rash in most cases. In rare cases, blisters, itching, or pain were seen beneath the transducer arrays. The skin reaction improved with use of topical corticosteroids. In persistent cases, the condition resolved with the use of low dose oral corticosteroids. Regular relocation of the transducer arrays was necessary in order to allow for continuous treatment.

3. WHAT ARE THE RISKS OF USING OPTUNE LUA TREATMENT KIT AND ILE TRANSDUCER ARRAYS?

Skin irritation is often seen under the ILE Transducer Arrays when using the Optune Lua treatment kit. This will look like a red rash, small sores or blisters on your body. In general, this will not cause skin damage that cannot be fixed. The irritation can be treated with steroid cream or by moving the ILE Transducer Arrays. If you do not use steroid cream, the skin irritation could become more serious. This may lead to open sores, infections, pain and blisters. If this happens, stop using the steroid cream and contact your doctor.

4. MECHANISM OF ACTION AND PERFORMANCE

Your doctor has prescribed the Optune Lua treatment kit for use at home because you are a good candidate for treatment with the device.

The Optune Lua treatment kit is a portable device. It produces electric fields, called tumor treating fields ("TTFields"). Transducer arrays connected to the device deliver TTFields to your chest. The TTFields are intended to destroy lung cancer cells.

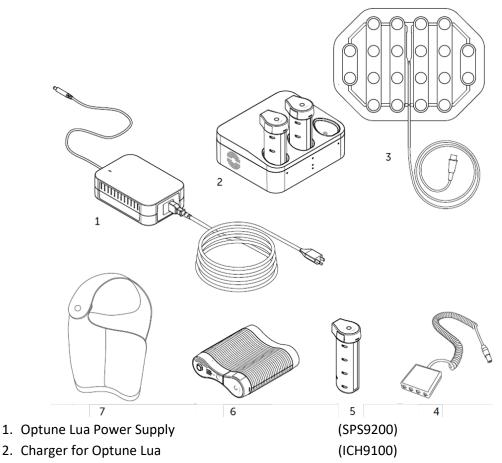
The device and battery are carried in a shoulder bag. You should use them as much as you can.

In this manual, the term "Optune Lua treatment kit" refers to the Electric Field Generator (also called "the device"), connection cable, power supply, battery, battery charger and ILE Transducer Arrays.

Optune Lua is a portable medical device that delivers electric fields called "TTFields" to the chest using transducer arrays. TTFields are intended to kill cancer cells.

The Science - The TTFields produce electric forces which disrupt cancer cell growth in cell cultures and in animals by inhibiting microtubule polymerization and by causing disruption of cell integrity during cytokinesis/division (see Kirson et al., Cancer Research 2004, Kirson et al., PNAS 2007, Salzberg et al., Onkologie 2008 and Kirson et al., BMC Medical Physics 2009).

5. OVERVIEW OF THE OPTUNE LUA TREATMENT KIT AND ILE TRANSDUCER ARRAYS



1. Optune Lua Power Supply

3. ILE Transducer Array

4. Optune Lua Connection Cable

5. Optune Lua Battery

6. Optune Lua electric field generator – the device

7. Carrying Bag

(Small: ILE1010, ILE1010W)

(Large: ILE1030, ILE1030W))

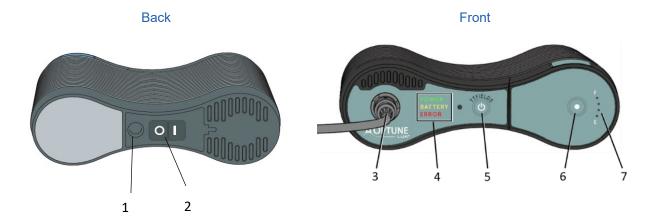
(CAD9100)

(IBH9200)

(TFT9200)

The Optune Lua device is an automatic system. The TTFields treatment should be kept on as continuously as possible (at a minimum 12 hours a day, 7 days a week). Breaks from treatment should be kept as short as possible.

You will need to learn how to place it in a carrying bag, connect a battery and operate the treatment kit. The following controls will allow you to do this:



- 1 Power Supply Port
- 2 Optune Lua Power Switch
- 3 Connection Cable (CAD) Socket
- 4 POWER / BATTERY / ERROR Indicators
- 5 TTFields ON / OFF Button
- 6 Battery Test Button
- 7 Battery Gauge

6. GLOSSARY OF SYMBOLS

	Follow instructions for use
MD	Medical Device
	Manufacturer information: Novocure GmbH, Business Village D4, Park 6/Platz 10, 6039 Root, Switzerland
#	Model number
REF	Part Number
SN	Serial number
LOT	Lot Number
UDI	Unique Device Identifier Indicates a device carries Unique Device Identifying information.
_W]	Manufacture date

YYY-MM	Use-by date/Expiry date
<u> </u>	Caution Consult the instructions for use for important cautionary information such as warnings and precautions
Z	Waste Electrical and Electronic Equipment recycling "WEEEE disposal" Contact technical support to arrange for proper disposal of transducer arrays that are used up or no longer in use.
Li-ion	Batteries are Lithium Ion. Contact technical support to arrange for proper disposal of batteries that are used up or no longer in use
	Do not re-use: The ILE Transducer Arrays are for single use and should not be re-used.
STERILE R	Indicates that the packaged products are sterile, the products have been sterilized by irradiation and the packaging is a single sterile barrier system
STERILE R	Sterile/sterilization method The ILE Transducer Arrays are sterilized by Gamma irradiation
STERMUZE	Do not re-sterilize
	Do not use if package is damaged Do not use the ILE Transducer Arrays if their packaging is breached.

	Protect from heat and radioactive sources
	The Optune Lua device, additional parts and ILE Transducer Arrays should be kept away from extreme heat and sources of radiation
IPxx	IP code: A coding system to indicate the degrees of protection provided by an enclosure against access to hazardous parts or water.
	IP21: The power supply protects persons against access to hazardous parts with fingers. Protects the equipment inside the enclosure against ingress of solid foreign objects of 12.5 mm in diameter or greater and against ingress of vertical falling water drops.
	IP22: The device protects persons against access to hazardous parts with fingers. Protects the equipment inside the enclosure against ingress of solid foreign objects of 12.5 mm in diameter or greater and against ingress of vertical falling water drops when enclosure is tilted up to 15°.
j	Keep dry. Do not enter rooms with high humidity or danger of direct exposure to water while wearing the device. Do not use the device if not within its carrying bag.
	Do not expose the device to direct rain.
	For indoor use only
	Class II equipment per IEC 60601-1
	BF type applied part
⚠	Symbolizes the part which comes in contact with the patient

	Storage Temperature range Do not expose to temperatures below -5°C or above 40°C
%	Storage humidity range. Do not expose to humidity below 15% or above 93%
Ţ	Fragile, handle with care
P1 P2 N1 N2	P1 P2 N1 N2 black and white coding on the connection box
C€ 0197	CE Mark with Notified Body Number
EC REP	European authorized representative MDSS GmbH Schiffgraben 41 30175 Hannover, Germany
	Importer details: Novocure Netherlands B.V., Prins Hendriklaan 26, 1075 BD, Amsterdam, The Netherlands
-0	Power ON / OFF switch for the device and battery charger: When the switch is in the I position the device is ON and will light up green. When the switch is in the O position the device is OFF

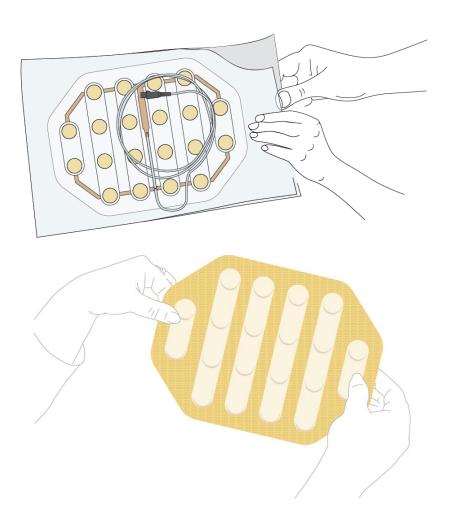
7. BEFORE YOU BEGIN

- You will need four (4) ILE Transducer Arrays (sterile) every 3-4 days in order to maintain treatment with the Optune Lua treatment kit.
- You will need to make sure you have the right sized flank transducer arrays for your chest size.
- Make sure you have ample supply of ILE Transducer Arrays to keep you going until your next visit to your physician.
- Before using an ILE Transducer Array make sure the package is sealed by gently rubbing the package between thumb and pointer finger on all four sides. The package should be closed on all sides. There should be no openings in the package seal. If the package is not sealed, the transducer array may be damaged. A damaged transducer array will not work properly and may cause the device to turn off. Do not use an ILE Transducer Array which has been opened previously.
- The ILE Transducer Arrays are for single use and should not be re-used.
- Maintenance and cleaning The ILE Transducer Arrays are supplied sterile for single use and thus do not require maintenance, cleaning or disinfection.
- The ILE Transducer Arrays are provided sterile for single use.

8. DIRECTIONS FOR USE

8.1. REMOVING THE TRANSDUCER ARRAY FROM ITS PACKAGE

- The ILE Transducer Arrays are supplied sterile and are to be used with the Optune Lua treatment kit only.
- ILE Transducer Arrays come in two sizes small and large. You should use two large transducer arrays on the back and front of your chest. You should use either two large or two small transducer arrays on both your flanks (on your sides beneath your arm pits), depending on your chest size.
- Open the see-through envelopes of each of the four (4) ILE Transducer Arrays, by gently pulling apart the opposing edges of the envelope. Hold the transducer array as shown in the illustration



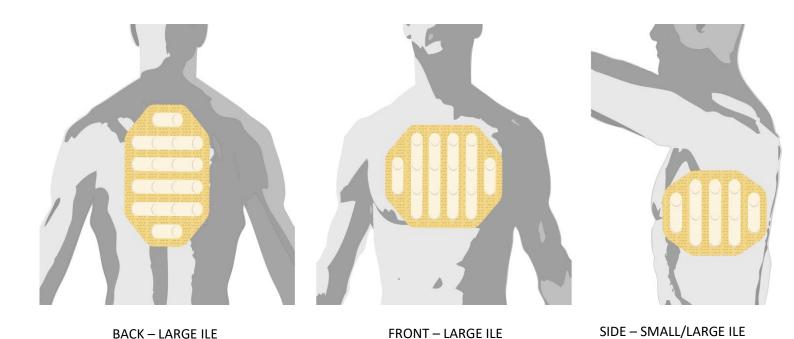
8.2. PREPARING YOUR SKIN FOR TRANSDUCER ARRAY PLACEMENT

- Wash your skin on the chest and flanks using a gentle soap.
- Remove any remnant adhesive from your skin from prior transducer arrays by wiping with baby oil.
- If you have significant chest hair, shave your entire torso using an electric shaver. Make sure no stubble is left.
- Wipe your skin with 70% Alcohol (medical grade any manufacturer).
- If the skin is red, apply the steroid cream prescribed to you by your physician.
- If you have any sores on your skin treat them as instructed by your treating physician.
- Wait at least 15 minutes and gently wipe your skin again with 70%
 Alcohol to facilitate adhesion of the transducer arrays to your skin.

8.3. PLACING THE TRANSDUCER ARRAYS

Once every 3-4 days perform the following steps to replace your transducer arrays:

- Remove the transducer arrays already on your chest and flanks by peeling the medical tape away from your skin.
 Note the black and white color of the transducer array connectors - each pair of the same color will be positioned opposite to each other on your body.
- 2) Remove the transducer array liner from the first transducer array.
- 3) Place the transducer array on your chest in the same location as before, but shifting the transducer array 2 cm to avoid areas of redness
- 4) Place the other three ILE Transducer Arrays in the same fashion.
- 5) You will need to ask for assistance from a friend or family member to place the back transducer array.
- 6) Press the entire rim of the transducer array tape to your skin.



8.4. CONNECTING THE ILE TRANSDUCER ARRAYS TO THE OPTUNE LUA DEVICE

- Connect the four black and white transducer array connectors to the corresponding black and white coded sockets on the Optune Lua connection cable.
- Make sure the transducer arrays connect the following way:
 - o Front transducer array (large) connects to P1 (black)
 - Back transducer array (large) connects to N1 (black)
 - o Right transducer array (either large or small) connects to P2 (white)
 - o Left transducer array (either large or small) connects to N2 (white)
- Press firmly to verify the connectors are inserted all the way.
- Collect the transducer array wires together and bind with a small piece of tape where convenient.
- You may clip the connection cable clip to your belt.



8.5. THE CONNECTION CABLE

The connection cable is the coiled, stretchy cord that runs from the connection box to the device. The four transducer array connectors (two blacks and two whites) are plugged into the connection box. The black and white coding matches with the transducer array position on the body.

Follow the instructions to connect to the device:

- Verify that the arrow on the connection cable is facing up and is aligned with the arrow on the connection cable socket of the device and plug in the connection cable.
- Push in the connection cable until you hear a snap. It indicates that the connection cable is in its place.



8.6. STARTING & STOPPING THE DEVICE

To start treatment:

The ILE Transducer Arrays should be attached to your body

- 1. Plug the ILE Transducer Arrays into the connection cable box (see Sections 8.4 and 8.5)
- 2. Plug the connection cable into the device, aligning the connector arrow with the socket arrow (see Section 8.5).
- 3. Connect a power source either a charged battery (Section 8.7) or a power supply (Section 8.9) to the device.
- 4. Turn ON the device by using the power switch.



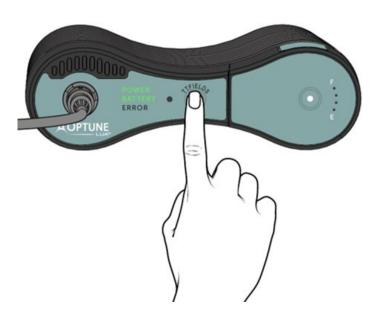
Wait about 10 seconds for the self-check to be completed, until the green "POWER" indicator illuminates



NOTE: If a charged battery is installed (and no power supply is connected), the green "BATTERY" indicator illuminates. If the device is connected to the power supply, it will be operated from the power supply and the "BATTERY" indicator will turn off.



Activate TTFields by pressing the TTFields ON/OFF button



The "TTFIELDS" indicator, above the TTFields ON/OFF button, should illuminate in blue and stay on while the treatment is ON.

NOTE: If the blue indicator doesn't illuminate, then the treatment is OFF and you should check the setup and restart the procedure. If, after this, the indicator lights do not light up, refer to the Troubleshooting Guide (Section 12). If you still have problems, contact Novocure technical support (Section 13).

The green, blue and yellow indicators automatically dim in a dark room. The red "ERROR" indicator illumination level is permanent.

If the TTFields button isn't pressed within about 10 minutes after the device is turned ON, a notification signal sounds along with a flashing blue "TTFIELDS" indicator, indicating that the treatment is OFF. This is a reminder to start the therapy. The

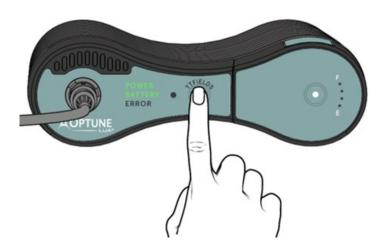
TTFields button should be pressed once to silence the alarm and again to start the therapy. The blue "TTFIELDS" indicator will then illuminate.

TO STOP TREATMENT:

Stopping treatment may be performed in each of the following situations:

- A. When the device is running properly and you need to take a break:
- 1. Stop treatment by pressing the TTFields ON/OFF button. TTFields therapy stops, indicated by the blue "TTFIELDS" indicator turn OFF.

NOTE: Device power is still ON.



2. Turn off the device by using the power switch



B. When an Error condition occurs:

If an error occurs, the device stops the treatment and sounds a loud beeping alarm. The red "ERROR" indicator illuminates (as shown below).

- 1. Press the TTFields button to stop the alarm. The red "ERROR" indicator will turn OFF. If the alarm sound persists, proceed to the next step to silence the alarm.
- 2. Turn OFF the device by using the power switch.



C. When the low battery indicator lights up:

When your battery runs out (after about one hour), an alarm will sound, the TTFields output will shut down (device stops the treatment) and an alarm will sound.

NOTE: The alarm sound is identical to the alarm that the device sounds when an error occurs. However, in this case, both the yellow "BATTERY" and red "ERROR" indicators light up.

- 1 Press the TTFields button to stop the alarm. The red "ERROR" indicator turns OFF.
- 2 Turn OFF the device by using the power switch.
- 3 Replace the battery (see Section 8.7).



8.7. CONNECTING AND DISCONNECTING THE BATTERY

The Optune Lua treatment kit is provided with four rechargeable batteries. Optune Lua operation requires one battery at a time. The other three batteries should stay in the battery charger.

If you plan to be away from home for more than one hour, carry extra batteries.

- 1 Slide the battery into the device.
- 2 Gently push the battery down until a click is heard, indicating it is fully latched.

NOTE: Take care not to drop the battery in place or force it into the battery slot.

3 Replace the battery each time it runs out (when the green "BATTERY" indicator turns yellow)





Gently press down to lock the battery in place.

To remove the battery from the slot, press both blue buttons on the sides of the battery and lift up.

Recharge the batteries in the charger (Section 8.8) for two to four hours. The batteries will keep most of their charge after being removed from the charger for several days but eventually will lose their charge. It will not hurt the batteries to keep them in the charger after they are fully charged so you can leave them there if they are not needed.

You can charge and use the batteries many times for about six to nine months. Over time, the length of time that the batteries can run the device (before the yellow low "BATTERY" indicator illuminates and the alarm beeps) will get shorter. If the time from treatment start with a full battery to low battery alarm, audible alarm sounds and the red "ERROR" indicator illuminates falls below 50 minutes contact technical support (Section 13) to get replacement batteries.

The "BATTERY" indicator will turn from green to yellow when the battery charge falls below a threshold. This is an indication that the battery should be changed soon. The treatment will continue to run while the yellow low "BATTERY" indicator is illuminated until the audible alarm sounds and the red "ERROR" indicator illuminates. Once this happens the treatment will stop and the device must be turned off and the battery replaced.

When the "BATTERY" indicator turns yellow, there are two ways to continue your treatment:

A. Option one:

If you are near the direct wall power supply, connect the power supply to the wall outlet to provide continuous therapy. This can be used before the battery is completely depleted, and before the device has alarmed. Follow the instructions:

- 1 Plug in the wall power supply to the back of the Optune Lua device (Section 8.9). Treatment continues while the device indicator indicates that it is no longer operated by battery power.
- 2 Press the two blue buttons on both battery sides and remove the battery by lifting it out of the device.
- 3 Charge the removed battery (Section 8.8).
- 4 Continue the treatment using the wall power supply.

B. Option Two:

If you are not near a wall power supply, follow the instructions to replace the battery:

NOTE: If the battery is totally depleted, start from step 2

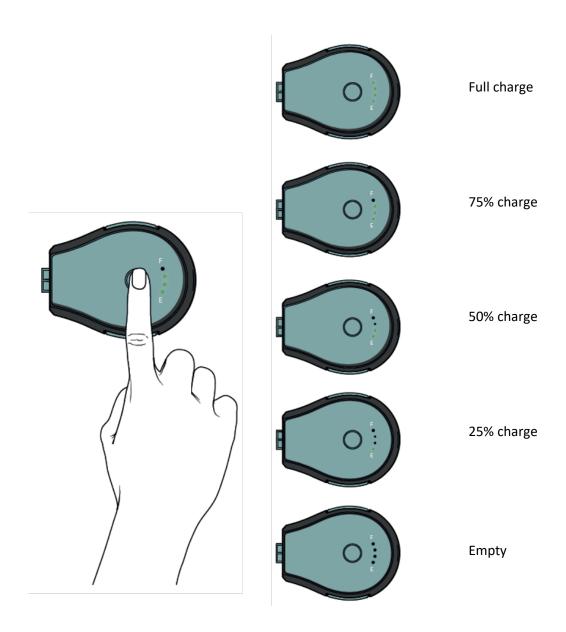
- 1. Press the TTFields button to stop the treatment.
- 2. Turn OFF the device by using the power switch (on the back of the device).
- 3. Press the two blue buttons on both battery sides and remove the battery by lifting it out of the device.
- 4. Select another fully charged battery.
- 5. Slide the fully charged battery into the device.
- 6. Gently push the battery down until a click is heard, indicating it is fully latched.
- 7. See Section 8.8 to check the battery gauge.
- 8. Turn ON the device by using the power switch and wait about 10 seconds until the device completes with the self-check.
- 9. Start treatment by pressing the TTFields button (Section 8.6).
- 10. Insert the used battery into the battery charger for recharging (Section 8.8).

8.8. CHARGING THE BATTERY

Checking the Battery Gauge

While you are using the Optune Lua treatment kit, you may want to check how much energy is left in your battery. Checking the battery will not interfere with, or stop, your treatment.

To check the battery capacity, press once on the button on the top of the battery. The battery capacity will be indicated by the lighted gauge to the right of the button. The gauge reads from Full (F) to Empty (E) like a gas gauge in a car.



The battery charger recharges used batteries. The battery charger uses power from a standard wall outlet. Each battery sits in a slot that connects it directly to the charger.

Before charging the batteries, plug the charger power cord into a standard wall outlet and turn ON the power switch at the charger rear side. The front lights of the charger will come on during a self-check then the small light in the center of the front panel will light up green indicating power is applied.

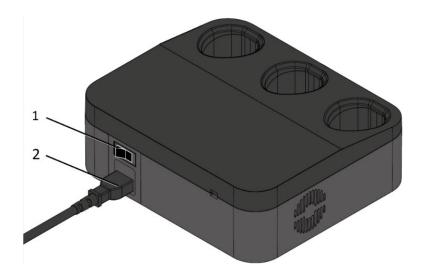
To recharge a used battery:

- 1. Place the used battery in one of the three openings in the top of the charger. Slide the battery in until it is fully in place.
- 2. The light directly in front of the opening where the battery is plugged in will illuminate flashing green. This indicates the battery is charging. The green light will flash faster once the battery has been charged to 95% of its capacity. You can also check the battery gauge while charging to get information regarding the amount of charge in the battery.
- 3. When the battery is fully charged (about 2 to 4 hours), the charge light will turn from flashing green to solid green. The solid green light will disappear upon removal of the battery or the disconnection of the charger from the standard wall outlet.

If a light on the front panel turns red, this indicates that there is a fault with the battery or charger and you should contact technical support for assistance. Do not use a battery if it creates a red light on the charger.

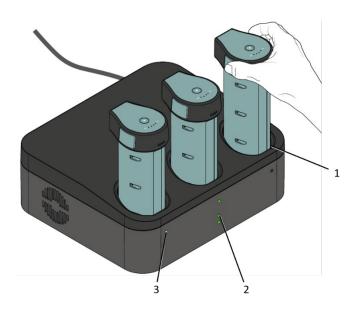
Keep the batteries in the charger even after they are fully charged. This will not harm the batteries

- 1. Power Switch
- 2. Power Cord



Battery Charger Rear View Showing the Power Switch and Where the Power Cord Connects

- 1. Battery Charging Slot
- 2. Charger Power Indicator
- 3. Battery Charge Indicator



Front view of the battery charger showing how the batteries are inserted into the charger

NOTE: The charger is not intended for use in the presence of flammable mixtures.

8.9. USING THE PLUG IN POWER SUPPLY

When you plan to stay in one place for a while, like when you are sleeping, you may use the plug-in power supply instead of the batteries. Unlike the batteries, there is no limit to how long the device can work when you use the plug-in power supply. The plug-in power supply will work with either US (120 VAC) or European (230 VAC) outlets.

NOTE: It is normal for the power supply to become warm when in use. If the power supply becomes too hot to touch, unplug it and contact technical support (Section 13).

When the device has a battery in, and is also connected to the wall power supply, it will utilize the wall power supply as the preferred power source. When the wall power cord is plugged in while the device is operated from the battery, the device will automatically switch from battery power to wall supply power.

Connecting the Plug-In Power Supply

1. Plug in the power supply cord into a standard wall outlet.

NOTE: You do not need to remove the battery from the device to use the wall power supply.

Please note that a battery in the device will not charge while the device is plugged into the wall power supply.

If the TTFields are activated, you do not need to turn them OFF.

- 2. Plug the power supply connector into the power supply port, located on the back side of the device (next to the power switch).
- 3. If the TTFields are already activated, the device will automatically switch to wall power supply without interruption of the treatment.
- 4. If the device is OFF, turn ON the power switch and wait about 10 seconds until the device completes with the self-check. Then, push the TTFields button to start the treatment (as described in Section 14).

Disconnect the Plug-In Power Supply and Go Back to Battery Power

Ensure that a charged battery is properly inserted in the device before removing the wall power supply. If the TTFields are activated, you need to turn them OFF before removing the wall power supply. The device will shut down and restart using battery power once the power supply is removed. In that case you will be required to push the TTFields button to start the treatment (as described in Section 8.6), after the self-check is completed.

- 1. Remove the power supply connector from the back side of the device. After about eight seconds, the "BATTERY" indicator on the front panel illuminates.
- 2. Store the plug-in power supply for future use.

8.10. DISCONNECTING FROM THE DEVICE

There are two ways to unplug the device in order to take a break from treatment:

- To unplug the connection cable from the device.
- To unplug the four transducer arrays from the connection cable.

To Unplug the Connection Cable from the Device

- 1. Stop treatment by pressing the TTFields button.
- 2. Turn OFF the device by using the power switch.
- 3. Hold the connector latch-sleeve and pull out the connection cable from the socket.

CAUTION! Do not pull on the cord!

You may now move around without the device, but you will still be connected to the connection cable and box.

To start treatment again after your break:

- 1. Plug the connection cable into the port with the arrows pointing up.
- 2. Turn ON the device by using the power switch. Wait about 10 seconds until the device completes with the self-check.
- 3. Activate TTFields by pressing the TTFields button.

To Unplug the Transducer Arrays from the Connection Cable

To take a break from treatment and completely disconnect from the device, unplug the ILE Transducer Arrays from the connection cable box. The four transducer arrays are plugged into the connection cable box (as described in Section 8.5). The connection cable remains plugged into the device socket.

- 1 Stop treatment by pressing the TTFields button.
- 2 Turn OFF the Optune Lua device by using the power switch.
- 3 Unplug the four transducer arrays from the connection box by pulling their connectors.

NOTE: You may have to wiggle the transducer array connectors gently to remove them. Do not pull on the cord.



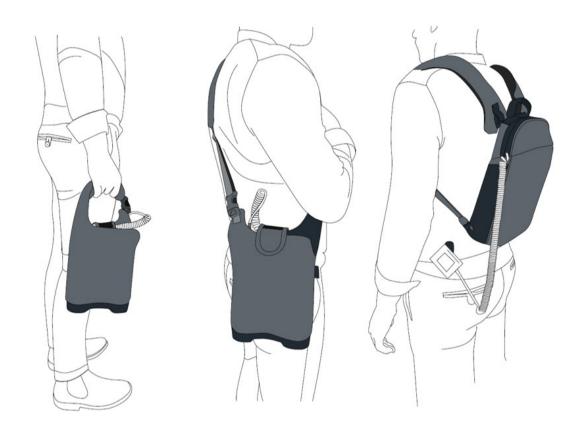
To restart treatment:

- 1 Plug the four transducer arrays into its matching color (black or white) in the connection box.
- 2 Turn ON the device by using the power switch and wait about 10 seconds until the device completes with the self-check.
- 3 Activate TTFields by pressing the TTFields button.

8.11. CARRYING THE DEVICE

Both the electric field generator and the battery fit in a carrying bag. The bag can be carried in one of three ways: by the handle on top or over the shoulder/ cross-body with a carrying strap attached, or as a backpack as shown below.

NOTE: Do not place the device in a different bag. The Optune Lua device has a fan on the inside that needs air flow. The bag that comes with the device is designed to allow for proper air flow. If you put the device in a bag without proper air flow, it could overheat and stop the treatment. If this happens, you will hear an alarm.



9. ENVIRONMENTAL CONDITIONS FOR OPERATION, STORAGE AND TRANSPORTATION

Conditions for operation

All treatment kit components shall be normally used under conditions specified below:

- Mainly for home use
- For indoor use only (chargers, power supply)
- Not for use in shower, bath tub or sink, or in heavy rain
- Not for use in presence of flammable mixtures
- Can be dropped on the floor with no safety hazard, but is no longer expected to function.

Conditions of visibility: any

Cleaning: all treatment kit components can be periodically cleaned with damp cloth, to remove dust and regular soil.

Physical operation conditions for all treatment kit components:

Temperature range: -5°C to +40°C (23°F to 104°F)

Relative Humidity range: 15-93%

Ambient pressure range: 700-1060hPa

Conditions for storage

- Temperature range: -5°C to +40°C (23°F to 104°F) for the device and additional parts
- Temperature range: 5°C to +27°C (41°F to 80°F) for the transducer arrays

Conditions for transport

Transportation of the device, ILE Transducer Arrays and additional parts shall be possible using air/ground transportation in weather protected conditions as specified below:

- Temperature range: -5°C to +40°C (23°F to 104°F)
- Maximal relative humidity 15-93%
- No direct exposure to water

10. EXPECTED LIFE

The expected service life for the Optune Lua device and all components of the treatment kit is 5 years.

The expected service life of the ILE Transducer Arrays is 9 months. The ILE Transducer Arrays have an expiration date. Please do not use the transducer arrays after the expiration date.

11. DISPOSAL

Please contact Novocure to arrange for proper disposal of used transducer arrays.

Do not throw them in the trash.

Novocure contacts local authorities for the determination of proper disposal method for potentially biohazardous parts.

12. TROUBLESHOOTING

Problem	Possible causes	Actions to be taken
Device POWER indicator does not light up after turning ON the device	 Device not connected to power source Battery depleted Battery malfunction If power supply – not properly plugged into the wall Device malfunction Power supply malfunction 	 If on battery – check battery gauge to verify it is not depleted. If it is – replace with a charged battery or to power supply Verify both the device and the power source are properly connected and retry Evaluate the integrity of all connectors. Nothing should appear to be damaged or broken in any way If device cannot be powered on by either the battery or the wall power supply or if anything appears to be damaged do not use the device Call technical support
Any cable detached from transducer array/ connection cable/ device	 Too much physical force to cables Device malfunction 	 Silence the notification signal by pressing the TTFields button Evaluate the connectors. If intact – reconnect and re-start therapy If anything appears damaged or cannot be properly connected do not try to use the device Call technical support
Device dropped or wet	Incorrect use	 Press TTFields button to stop therapy Turn OFF power switch Disconnect from power Call technical support
Device alarm on, and low BATTERY indicator is yellow	Low battery Device is turned ON, but the therapy has not been activated	 Replace battery as described above in Section 8.7 Turn ON treatment Press the TTFields button to stop the alarm Wait a few seconds then press the TTFields button again If the blue lights around the TTFields button light up – the therapy has now been activated If the notification signal recurs within a few minutes: Silence the notification signal and

Problem	Possible causes	Actions to be taken
		power the device down completely 2. Disconnect all equipment and make sure that nothing appears to be damaged or broken. If something is — replace the damaged item before trying to power the device back 3. Re-connect all equipment in proper order and power the device back up. Verify the self-check is completed and press the TTFields button 4. Check vents on device to make sure they are not blocked 5. If lying down, get up and move your body 6. Make sure transducer arrays are well stuck to the body, add tape if needed 7. Restart treatment 8. If alarm keeps going, turn OFF the device and call technical support
Device alarm is flashing, the "TTFIELDS" indicator above the TTFields button will flash blue and audio sound 3 very short beeps, stops for 2.5 seconds and beeps 3 times again	Therapy Timeout	The notification alarm on the device will sound if it is powered on for about 10 minutes, but therapy is not initiated. This is a reminder to start therapy and does not indicate a malfunction. 1. Silence the notification alarm by pressing the TTFields button then wait a few seconds and press the TTFields button again to initiate treatment. The blue indicator around the TTFields button will illuminate to indicate therapy is now on 2. If you encounter further alarms please review the following troubleshooting descriptions in this section.
Low BATTERY indicator remains on after battery replaced	 Charger malfunction Battery malfunction Device malfunction 	1. Replace battery with an additional charged battery 2. If problem is not fixed — call technical support
When powering on the device a continuous notification alarm sounds and all lights remain on	Device is too hot Device malfunction Power Source Malfunction	 Power the device off completely using the power switch Verify the device is not hot to the touch Connect the device to a different power source and try powering on again If device cannot be powered on by

Problem	Possible causes	Actions to be taken		
indefinitely. Device does not complete the self-check.		either the battery or the wall power supply or if anything appears to be damaged, please contact technical support		
Redness of the skin beneath the transducer arrays	Common side effect	 Use steroid cream prescribed by your doctor when replacing transducer arrays. Place transducer arrays in a location shifted by 3/4 of an inch (2 cm) from the last location (so the adhesive gel is between the red marks). If the redness gets worse: See your treating doctor 		
Blisters beneath the transducer arrays	Rare side effect	See your treating doctor		
Itching beneath the transducer arrays	Rare side effect	 Use steroid cream prescribed by your doctor when replacing transducer arrays. Place transducer arrays in a location shifted by 3/4 of an inch (2 cm) from the last location (so the adhesive gel is between the red marks). If the itching gets worse: See your treating doctor 		
Pain beneath the transducer arrays	Rare side effect	Stop treatment See your treating doctor		

13. ASSISTANCE AND INFORMATION

Technical support

For technical support, contact your Device Support Specialist. His/her contact information will be supplied to you separately.

If you are unable to get a hold of your Device Support Specialist, you can contact the EMEA Novocure technical support email: SupportEMEA@novocure.com or patientinfoEMEA@novocure.com.

Please describe the problem and provide the following information when you contact:

NAME (First/Last)

EMAIL

TELEPHONE (optional)

COUNTRY:

QUESTION:

Please also have the device serial number on hand when you contact the DSS or Technical Support. The serial number can be found on the bottom of the device (TTFields Generator)

Clinical support

If you feel any change in your health or any side effects from the treatment call your doctor right away.

Reporting

If you experience a serious incident that occurs while using the Optune Lua treatment kit and ILE Transducer Arrays you should report it to the manufacturer (Novocure) and the competent authority of the Member State in which you reside.

Traveling with Optune Lua

The treatment kit's batteries contain lithium ion material and are restricted from being checked as luggage for passenger aircraft travel. They can be carried in the passenger cabin. Please contact your DSS if you have questions related to travel restrictions.

Note: The Optune Lua device and transducer arrays will activate metal detectors.

14. GLOSSARY

Cancer – abnormal cell division that spreads without control

Chemotherapy – medication used to destroy cancer cells

Clinical trial – a research study that involves people

Contraindications – situations when a treatment should not be used

EKG – Electrocardiogram

EN 60601-1 - Harmonized standards series for safety of medical devices

Electric Field Generator (the device) – a portable device for delivering TTFields to the lungs of patients with MPM

ILE – Insulated Lung Transducer Arrays

Local – in one part of the body

NSCLC – Non small cell lung cancer

Optune Lua – NovoCure's Tumor Treatment Fields Device for the treatment of advanced NSCLC

Optune Lua treatment kit – treatment kit containing the Optune Lua device (TFT9200); Connection cable (CAD9100); power supply (SPS9200); battery (IBH9100); charger (ICH9100); ILE Transducer Arrays.

Progression – when cancer comes back after being treated

Radiation – a treatment involving x-rays used to kill tumor cells

Steroids – When used on the skin, a medication that can reduce inflammation

Systemic – throughout the body

Topical – on the surface of the skin

Transducer array – adhesive bandages that hold insulated ceramic discs that deliver TTFields to the chest

TTFields – Tumor Treating Fields: Alternating electric fields, delivered using transducer arrays to the part of the body with a solid tumor. The fields have been shown to destroy tumor cells

Tumor – an abnormal growth of tissue

15. APPLICABLE STANDARDS

The Optune Lua treatment kit electronic components and the sterile transducer arrays comply with the latest editions of the following safety standards:

- EN 60601-1Medical electrical equipment Part 1: General requirements for safety
- EN 60601-1-2 Medical electrical equipment-Part 1-2: General requirements for safety-Collateral standard: Electromagnetic compatibility-Requirements and tests
- EN 60601-1-11- Medical electrical equipment -- Part 1-11: General requirements for basic safety and essential performance -- Collateral standard: Requirements for medical electrical equipment and medical electrical systems used in the home healthcare environment
- EN 60601-1-6 Part 1-6: General requirements for basic safety and essential performance - Collateral Standard: Usability
- EN 62366-1 Application of usability engineering to medical devices
- EN 62304 Medical device software. Software life-cycle processes

16. INPUT OUTPUT SPECIFICATIONS

The Optune Lua treatment kit is considered class II equipment according to EN 60601-1.

Mode of operation – continuous. The device is portable when battery operated and stationary equipment when connected to the power supply.

The applied part is classified as BF.

The Optune Lua treatment kit is not intended for use in the presence of flammable mixtures.

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17. EMITTED RADIATION & ELECTROMAGNETIC COMPATIBILITY

The Optune Lua device and the accompanying battery charger (ICH9100) and power supply (SPS9200) need special precautions regarding EMC and need to be installed and put into service according to the EMC information provided below.

Portable and mobile RF communications equipment can affect the Optune Lua treatment kit and the accompanying battery charger.

The Optune Lua device (TFT9200) should be used with the following cables and additional parts only:

- 1 connection cable (CAD9100)
- 2 ILE Transducer Arrays (ILE1010; ILE1030; ILE1010W; ILE1030W)
- 3 battery (IBH9200)
- 4 power supply (SPS9200)
- 5 Battery charger (ICH9100)
- 6 Unshielded AC mains cables for indoor use only with a maximal length of 1.5m

The use of accessories, parts and cables other than those specified may result in increased EMISSIONS or decreased IMMUNITY of the Optune Lua treatment kit.

Table 1 – Guidance and MANUFACTURER'S declaration – ELECTROMAGNETIC EMISSIONS – for all ME EQUIPMENT and ME SYSTEMS

Guidance and manufacturer's declaration – electromagnetic emissions

The Optune Lua treatment kit is intended for use in the electromagnetic environment specified below.

The customer or the user of the Optune Lua should assure that it is used in such an environment.

Emissions test	Compliance	Electromagnetic environment – guidance		
RF emissions CISPR 11	Group 1			
RF emissions CISPR 11	Class B	The Optune Lua treatment kit is suitable for use in all establishments, including domestic		
Harmonic emissions IEC 61000- 3-2	Class A	establishments and those directly connected to the public low-voltage power supply network that		
Voltage fluctuations/ flicker emissions IEC 61000-3-3	Complies	supplies buildings used for domestic purposes.		

Guidance and manufacturer's declaration – electromagnetic emissions

The ICH9100 charger and the SPS9200 power supply are intended for use in the electromagnetic environment specified below. The customer or the user of the ICH9100 charger and the SPS9200 power supply should assure that it is used in such an environment.

Emissions test	Compliance	Electromagnetic environment – guidance
RF emissions CISPR 11	Group 1	The ICH9100 charger and the SPS9200 power supply use RF energy only for their internal function. Therefore, their RF emissions are very low and are not likely to cause any interference in nearby electronic equipment.
RF emissions CISPR 11	Class B	The ICH9100 charger and the SPS9200 power supply are suitable for use in all establishments,
Harmonic emissions IEC 61000-3-2	Class A	including domestic establishments and those
Voltage fluctuations/ flicker emissions IEC 61000-3-3	Complies	directly connected to the public low-voltage power supply network that supplies buildings used for domestic purposes.

Warning: The Optune Lua treatment kit, the ICH9100 charger and the SPS9200 power supply should not be used adjacent to or stacked with other equipment.

Table 2 – Guidance and MANUFACTURER'S declaration – ELECTROMAGNETIC IMMUNITY – for all ME EQUIPMENT and ME SYSTEMS

Guidance and manufacturer's declaration – electromagnetic immunity

The Optune Lua treatment kit is intended for use in the electromagnetic environment specified below.

The customer or the user of the Optune Lua treatment kit should assure that it is used in such an environment.

Emissions test	IEC 60601 Test level	Compliance level	Electromagnetic environment – guidance	
Electrostatic discharge (ESD) IEC 61000-4-2 ±8 kV contact, ± 2 kV, ± 4 kV, ±8 kV, ± 15 kV air		±8 kV contact, ± 2 kV, ± 4 kV, ±8 kV ± 15 kV air	The relative humidity should be at least 5%.	
Electrical fast transient/burst lines ±2 kV for power supply lines ±1 kV for input/output lines		±2 kV for power supply lines ±1 kV for input/output lines 100 kHz repetition frequency	Mains power quality should be that of a typical commercial or hospital environment.	
Surge IEC 61000-4-5	± 0,5 kV, ±1 kV line to line ± 0,5 kV, ± 1 kV, ±2 kV line to ground	± 0,5 kV, ±1 kV line to line ± 0,5 kV, ± 1 kV, ±2 kV line to ground	Mains power quality should be that of a typical commercial or hospital environment.	
Voltage dips, short interruptions and voltage variations on power supply input lines IEC 61000-4-11 O % UT; 0,5 cycle At 0°, 45°, 90°, 135°, 180°, 225°, 270° and 315° interruptions and voltage W UT; 1 cycle and 70 % UT; 25/30 cycles Single phase: at 0° 0 % UT; 250/300 cycle		0 % UT; 0,5 cycle At 0°, 45°, 90°, 135°, 180°, 225°, 270° and 315° 0 % UT; 1 cycle and 70 % UT; 25/30 cycles Single phase: at 0° 0 % UT; 250/300 cycle	Mains power quality should be that of a typical commercial or hospital environment.	
Power frequency (50/60 Hz) magnetic field IEC 61000-4-8		30 A/m	Power frequency magnetic fields should be at levels 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

The ICH9100 charger and the SPS9200 power supply are intended for use in the electromagnetic environment specified below. The customer or the user of the ICH9100 charger and the SPS9200 power supply should assure that they are used in such an environment.

Emissions test	IEC 60601 Test level	Compliance level	Electromagnetic environment – guidance	
Electrostatic discharge (ESD) IEC 61000-4-2	± 8 kV contact ± 2 kV, ± 4 kV, ± 8 kV, ± 15 kV air	± 8 kV contact ± 2 kV, ± 4 kV, ± 8 kV, ± 15 kV air	The relative humidity should be at least 5%.	
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 ±1 kV for input/output lines 100 kHz repetition frequency	Mains power quality should be that of a typical commercial or hospital environment.	
Surge IEC 61000-4-5	\pm 0,5 kV, \pm 1 kV line to line \pm 0,5 kV, \pm 1 kV, \pm 2 kV line to ground	\pm 0,5 kV, \pm 1 kV line to line \pm 0,5 kV, \pm 1 kV, \pm 2 kV line to ground	Mains power quality should be that of a typical commercial or hospital environment.	
Voltage dips, short interruptions and voltage variations on power supply input lines IEC 61000-4-11	0 % UT; 0,5 cycle At 0°, 45°, 90°, 135°, 180°, 225°, 270° and 315° % UT; 1 cycle and 70 % UT; 25/30 cycles Single phase: at 0° 0 % UT; 250/300 cycle	0 % UT; 0,5 cycle At 0°, 45°, 90°, 135°, 180°, 225°, 270° and 315° 0 % UT; 1 cycle and 70 % UT; 25/30 cycles h) Single phase: at 0° 0 % UT; 250/300 cycle	Mains power quality should be that of a typical commercial or hospital environment.	
Power frequency (50/60 Hz) 30 A/m magnetic field IEC 61000-4-8		30 A/m	Power frequency magnetic fields should be at levels 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 = 120V and 230V

Table 3 – Guidance and MANUFACTURER'S declaration – ELECTROMAGNETIC IMMUNITY – for ME EQUIPMENT and ME SYSTEMS that are not LIFE-SUPPORTING

Guidance and manufacturer's declaration – electromagnetic immunity

The Optune Lua treatment kit is intended for use in the electromagnetic environment specified below. The customer or the user of the Optune Lua treatment kit should assure that it is used in such an environment.

Immunity test	IEC 60601 test level	Compliance level	Electromagnetic environment – guidance
Conducted RF IEC 61000-4-6	3 V 0,15 MHz – 80 MHz 6 V in ISM bands between 0,15 MHz and 80 MHz	3 V 0,15 MHz – 80 MHz 6 V in ISM bands between 0,15 MHz and 80 MHz	Portable and mobile RF communications equipment should be used no closer to any part of the Optune Lua treatment kit, including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter. Recommended separation distance
Radiated RF IEC 61000-4-3	80 % AM at 1 kHz (table 8.5.1)	80 % AM at 1 kHz 10 V/m 80 MHz to 2.7 GHz 80 % AM at 1 kHz	$d=\frac{6}{E}\;\sqrt{P}$ Where P is the maximum power in W, d is the minimum separation distance in m, and E is the IMMUNITY TEST LEVEL in V/m. Field strengths from fixed RF transmitters, as determined by an electromagnetic site survey ^a , should be less than the compliance level in each frequency range.
Radiated fields in close proximity Standard IEC 61000-4- 39	8A/m 30kHz CW 65A/m 134.2kHz pulse modulated 2.1kHz 7.5A/m 13.56MHz pulse modulated 50kHz	5cm distance	

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

a. Field strengths from fixed transmitters, such as base stations for radio (cellular/cordless) telephones and land mobile radios, amateur radio, AM and FM radio broadcast and TV broadcast 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 the Optune Lua treatment kit is used exceeds the applicable RF compliance level above, the Optune Lua treatment kit should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as reorienting or relocating the Optune Lua treatment kit.

Guidance and manufacturer's declaration – electromagnetic immunity

The ICH9100 charger and the SPS9200 power supply are intended for use in the electromagnetic environment specified below. The customer or the user of the ICH9100 charger and the SPS9200 power supply should assure that they are used in such an environment.

Immunity test	IEC 60601 test level	Compliance level	Electromagnetic environment – guidance			
			Portable and mobile RF communications equipment should be used no closer to any part of the ICH9100 charger and the SPS9200 power supply, including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter.			
	3 V	3 V				
	0,15 MHz – 80 MHz	0,15 MHz – 80 MHz	Recommended separation distance			
	6 V in ISM bands	6 V in ISM bands				
Conducted RF IEC	between 0,15 MHz and	between 0,15 MHz and 80 MHz	$d = \frac{6}{F} \sqrt{P}$			
61000-4-6	80 MHz	and so winz	L			
	80 % AM at 1 kHz	80 % AM at 1 kHz	Where P is the maximum power in W, d is the minimum separation distance in m, and E is the IMMUNITY TEST LEVEL in V/m.			
Radiated RF IEC	(table 8.5.1)					
61000-4-3		10 V/m	Field strengths from fixed RF transmitters, as deter-mined by an electromagnetic site survey ^a , should be less than the			
	10 V/m	80 MHz to 2.7 GHz	compliance level in each frequency range.			
		80 % AM at 1 kHz				
			Interference may occur in the vicinity of equipment marked with the following symbol:			

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

a. Field strengths from fixed transmitters, such as base stations for radio (cellular/cordless) telephones and land mobile radios, amateur radio, AM and FM radio broadcast and TV broadcast 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 the ICH9100 charger and the SPS9200 power supply are used exceeds the applicable RF compliance level above, the ICH9100 charger and the SPS9200 power supply are should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as re-orienting or relocating the ICH9100 charger and the SPS9200 power supply.

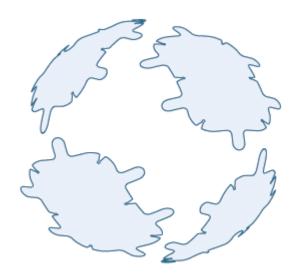
Normal operation: The Optune Lua treatment kit is working properly when the blue LED surrounding the TTFields button are lit and no notification signal sounds. The ICH9100 charger is working properly when all the LEDs are lit. The SPS9200 power supply is working properly when the blue LEDs surrounding the TTFields button on the Optune Lua device are lit and no notification signal sounds.

Table 4 – Recommended separation distances between portable and mobile RF communications equipment and the ME EQUIPMENT or ME SYSTEM – for ME EQUIPMENT and ME SYSTEMS that are not LIFE-SUPPORTING

	Separation distance according to frequency of transmitter m						
Rated maximum output power of transmitter W	380 – 390MHz	430 – 470MHz	704 – 787MHz	800 – 960MHz	1700 – 1990MHz	2400 – 2570MHz	5100 – 5800MHz
The Optune Lua is intended for use in an electromagnetic environment in which radiated RF disturbances are controlled. The customer or the user of the Optune Lua can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and Optune Lua as recommended below, according to the maximum output power of the communications equipment.							
0.2	0.3	0.3	0.3	0.3	0.3	0.3	0.3
1.8	0.3	0.3	0.3	0.3	0.3	0.3	0.3
2	0.3	0.3	0.3	0.3	0.3	0.3	0.3

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

For transmitters rated at a maximum output power not listed above, the recommended separation distance d in meters (m) can be determined 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.



novœure®



Manufacturer information:

Novocure GmbH, Business Village D4,Park 6/Platz 10, 6039 Root, Switzerland



Importer details:

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