d. Slide off the battery compartment cover.
e. Remove the old battery, and install the new battery.
f. Correct polarity is needed for battery proper functionality.
g. Place the battery cover back into the original position.

The SunStim™ or SunStim Plus™ Peripheral Nerve Stimulator should not be used in case of battery leakage, as the acid may impair internal circuits.

11. CLEANING

Only soft, damp, clean cloth is recommended for cleaning of SunStim™ or SunStim Plus™ Peripheral Nerve Stimulator. Liquids or humidity can damage this device, and their usage is strictly prohibited.

12. SERVICE

The SunStim™ or SunStim Plus™ Peripheral Nerve Stimulator should not be serviced by the end user(s). End user(s) may replace batteries only.

In a case of device damage or malfunction, do not attempt any repairs. In such case, contact SunMed Customer Service at 800.433.2797 immediately.

13. STORAGE AND TRANSPORT

The environmental conditions of use including conditions for transport and storage are as listed below:

- Operating Temperature: +10°C to +40°C
- Operating Humidity: 30% RH to 85% RH
- Storage/Transport Temperature Range: -20°C to +70°C
- Storage/Transport Humidity: 10% RH to 95% RH
- Operating and Storage/Transport Atmospheric Pressure: 800hPa to 1,060hPa

14. WARRANTY

The SunStim™ or SunStim Plus™ Peripheral Nerve Stimulator are free from defects in workmanship and materials for one (1) year from purchase, when used for the intended purpose and cared for in accordance with recommended procedure.

No charge repairs will be made during the one year warranty period, only if the SunStim™ or SunStim Plus™ Peripheral Nerve Stimulators have not be abused, and/or subjected to unauthorized repair.

15. SYMBOL MEANINGS

- Caution; consult accompanying documents
- Keep dry
- Consult instructions for use
- Keep away from sunlight
- Type BF applied part
- 800 hPa
- 1060 hPa
- 95%
- -20°C
- 70°C
- Operating and transport atmospheric pressure
- Storage/transport humidity
- Storage/transport temperature range

INSTRUCTIONS FOR USE

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14. WARRANTY .....................................8
15. SYMBOLS MEANING ...........................8
1. INDICATIONS FOR USE
The SunStim™ and SunStim Plus™ Peripheral Nerve Stimulators are 9 Volt DC battery-powered, non-sterile devices, intended for monitoring the magnitude of neuromuscular block in general anesthesia, by delivering an electrical stimulus near a peripheral motor nerve and assessing the response of the muscle innervated by that nerve.

2. EQUIPMENT CLASSIFICATION
Per IEC 60601-1, Medical Device Equipment, General Requirements for Safety, SunStim™ or SunStim Plus™ Peripheral Nerve Stimulators are classified as follows:

Type BF Equipment
Type B of equipment provides a particular degree of protection against electrical shock, predominantly regarding allowable leakage current and reliability of the protective earth connection (grounding).

Type F part applies to extends from the patient into the equipment and is isolated from all other parts of the equipment.

Water Ingress Protection = IPX The SunStim™ or SunStim Plus™ Peripheral Nerve Stimulator do not have protection against ingress of water.

Continuous Operation The SunStim™ or SunStim Plus™ Peripheral Nerve Stimulator is designed for Continuous Operation.

Load Resistance = 510Ω

3. ACCESSORIES
The SunStim™ and SunStim Plus™ Peripheral Nerve Stimulators are provided with:
- Extension Leadwires
- Bipolar Probes

4. EQUIPMENT DISPOSAL
Dispose of SunStim™ or SunStim Plus™ Peripheral Nerve Stimulator should be done in accordance with local regulations.

5. WARNING / CAUTION STATEMENTS
CAUTION: Federal Law (USA) restricts this device to sale by or on order of a physician or appropriate licensed practitioner.

DO NOT: Use SunStim™ or SunStim Plus™ Peripheral Nerve Stimulator for nerve localization for anesthetic regional block.

DO NOT: Use SunStim™ or SunStim Plus™ Peripheral Nerve Stimulator on patients with neuromuscular or skin diseases.

DO NOT: Use SunStim™ or SunStim Plus™ Peripheral Nerve Stimulator in the proximity of equipment which produces electromagnetic fields, short or micro waves.

DO NOT: Modify any components of this equipment.

DO NOT: Use SunStim™ or SunStim Plus™ Peripheral Nerve Stimulator in the presence of a flammable anesthetic mixture with air or in the presence of a flammable anesthetic mixture with oxygen or nitrous oxide.

6. EQUIPMENT DISPOSAL
Dispose of SunStim™ or SunStim Plus™ Peripheral Nerve Stimulator in accordance with local regulations.

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The SunStim™ and SunStim Plus™ Peripheral Nerve Stimulators are provided with:
- Extension Leadwires
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Dispose of SunStim™ or SunStim Plus™ Peripheral Nerve Stimulator in accordance with local regulations.

7. ACCESSORIES
The SunStim™ and SunStim Plus™ Peripheral Nerve Stimulators are provided with:
- Extension Leadwires
- Bipolar Probes

9.4 Electrical stimulus can be performed at the:

9.4.1. Ulnar Nerve - Leads/bipolar probes may be placed:
- 9.4.1.1 Along the medial aspect of the distal forearm (wrist);
- 9.4.1.2 Over the sulcus of the medial epicondyle of the humerus (elbow);
- 9.4.1.3 On hand, by placing the negative electrode on the palm between the base of the thumb, and the second finger, and the positive electrode in the same position on the dorsal side of the hand.

9.4.2 Median Nerve - Leads/bipolar probes may be placed:
- 9.4.2.1 Medial to the wrist;
- 9.4.2.2 At the elbow adjacent to the brachial artery.

9.4.3 Tibial Nerve - Leads/bipolar probes may be placed:
- 9.4.3.1 Along the lateral side of the popliteal fossa.
- 9.4.3.2 Posterior Tibial Nerve - Leads/bipolar probes are placed:
- 9.4.4.1 At the medial maleolus and anterior to the Achilles tendon at the ankle.
- 9.4.5 Peroneal Nerve - Leads/bipolar probes are placed:
- 9.4.5.1 On the lateral aspect of the knee.
- 9.4.6 Facial Nerve - Leads/bipolar probes are placed:
- 9.4.6.1 Negative electrode is placed anterior to the inferior part of the ear lobe, and the positive electrode is placed posterior or inferior to the lobe.
- 9.4.7 Spinal Accessory nerve - Leads/bipolar probes are placed:
- 9.4.7.1 Over the depression between the ramus.

10. MAINTENANCE
10.1 Prior to use, make sure to read and understand provided Instructions for Use. Check condition of the SunStim™ or SunStim Plus™ Peripheral Nerve Stimulator and provided connectors for proper functioning, prior to each use.

10.2 The SunStim™ or SunStim Plus™ Peripheral Nerve Stimulator and all accessories must be visually inspected at regular intervals for any material degradation or battery leakage.

10.3 This device does not require user maintenance/service, except the periodic battery replacement.

10.3.1 Battery Maintenance/Replacement
a. This device may be used with Alkaline 9V battery only.
b. Battery replacement is needed when the Battery LED flashes during the use.
c. Prior to battery replacement, make sure that the device is turned OFF, by rotating the control knob to the “OFF” position.
8. OPERATION INSTRUCTIONS

8.1 Switching the unit on and off

A. The SunStim™ or SunStim Plus™ Peripheral Nerve Stimulator can easily be switched on and off by rotating the rheostat (control knob) clockwise. The rheostat (control knob) is located on the left hand side of the Nerve Stimulator. Once ON, the device will be in the Stand-By mode, and no pulses will be produced.

B. The numbered rheostat (control knob) (numbered from 0-10) intended use it to adjust the stimulation amplitude of the output current, which may range from 0 to 70 mA. If the rheostat (control knob) is set to zero, stimulation amplitude will not be delivered.

8.2 Four Panel Touch Switches

A. Stimulation frequency patterns may be activated by pressing one of four panel touch switches (SunStim™ – STANDBY, TWITCH, 100 Hz, TOF; SunStim Plus™ – DBS, TWITCH, TETANUS, or TOF).

B. Pulse LED will flash each time a pulse is generated.

8.3 Output Connectors

A. Transcutaneously stimulation can be carried out by using surface electrodes.

B. SunStim™ or SunStim Plus™ Peripheral Nerve Stimulator is supplied with metal ball electrodes and lead wires.

C. Two (2) connectors, RED (positive) and BLACK (negative), are located on the top of the SunStim™ or SunStim Plus™ Peripheral Nerve Stimulator case.

D. The output current may reach up to 70 mA, measured with load of 2KOhm load, using a new 9V DC battery.

E. Provided lead wires with BLACK plug should be connected to the black output connector. This connection will create the negative output.

F. Provided lead wires with RED plug, should be connected to the red output connector. This connection will create the positive output.

G. Bipolar Probe Electrodes can be connected to output connectors, by plugging them directly into the two, RED (positive) and BLACK (negative), located on the top of the SunStim™ or SunStim Plus™ Peripheral Nerve Stimulator case, output connectors.

H. It is recommended to create a lead wires loop after they have been attached to electrodes, and a piece of tape placed over them in order to prevent possible electrodes displacement.

I. Nerve Stimulator should be connected to electrodes that are positioned over the selected nerve, prior to anesthesia induction.

9. PERIPHERAL NERVE MONITORING SITES

9.1 The site of stimulations should be away from the surgical field and its location accessible to the anesthesia provider.

9.2 Location of the instrument should approximately five (5) feet above the floor to provide easy viewing of the touch switches and indicators.

9.3 If visual or tactile nerve monitoring is to be used, the site location must be accessible to the anesthesia provider.

a) The SunStim™ or SunStim Plus™ Peripheral Nerve Stimulator may be hazardous to patients with implanted electrical, medical devices.

b) It is important to monitor the correlation of nerve reaction stimulation with the patient’s clinical condition, as there may be a discrepancy between the degree of relaxation and of the monitored muscle at the site of surgery.

c) Tetanus nerve stimulation should be performed only after the anesthetic has been administered.

d) Stimulus current must be increased gradually, until supramaximal stimulus is achieved. Applying currents greater than necessary for supramaximal stimulation may increase the risk of skin burns.

e) Prior to SunStim™ or SunStim Plus™ Peripheral Nerve Stimulator usage, the patient’s skin should be cleaned and completely dried. This area should be free of excessive hair, scar tissue, or any other lesions.

f) This device should not fall into liquids. Liquids should not be spilled over or into the device.

g) The leadwire or bipolar probe electrodes would be attached to the instrument while the numbered control knob is in the OFF position.

h) SunStim™ or SunStim Plus™ Peripheral Nerve Stimulator must be used with 9V alkaline battery only.

i) The operator of this device cannot touch the actual battery and the patient simultaneously.

j) Battery should be removed if the Nerve Stimulator is not likely to be used for some time.

k) In case of battery leakage, the device should not be used.

l) Prior to each use, check the SunStim™ or SunStim Plus™ Peripheral Nerve Stimulator for proper condition and functioning.

m) The SunStim™ or SunStim Plus™ Peripheral Nerve Stimulator should be stored/transport at the room ambient temperature, away from heat. No special ambient temperature range is specified for this device.

n) Caution should be used when switching between DBS (Double Burst) and Train-of-Four stimulation. Up to 92 seconds may be required before the responses are stabilized.
6. TECHNICAL DATA

<table>
<thead>
<tr>
<th>Model</th>
<th>SunStim™</th>
<th>SunStim Plus™</th>
</tr>
</thead>
<tbody>
<tr>
<td>Type of Device</td>
<td>BF</td>
<td>BF</td>
</tr>
<tr>
<td>Membrane Touch Switch</td>
<td>StandBy</td>
<td>Double Burst</td>
</tr>
<tr>
<td></td>
<td>Twitch</td>
<td>Twitch</td>
</tr>
<tr>
<td></td>
<td>100 Hz</td>
<td>Tetanus</td>
</tr>
<tr>
<td></td>
<td>Train-of-Four (TOF)</td>
<td>Train-of-Four (TOF)</td>
</tr>
</tbody>
</table>

**Pulse Characteristics**

| Pulse Width: | 200 Microseconds | 200 Microseconds |
| Pulse Type: | Square Wave Monophasic | Square Wave Monophasic |

**Tetanus**

| 100 Hz | 50 Hz or 100 Hz |

**Double Burst Pulses (DBS)**

| N/A | Two (2) 60 ms bursts of 50 Hz separated by 0.75 seconds |

**Output Current**

| Adjustable 0-70mA | Adjustable 0-70mA |

**Display**

| Pulse LED: | Flash red when pulse is generated |
| Battery LED: | Steady green, when the unit is turned ON |

**Battery Power**

| One 9V alkaline battery | Battery LED flashes when battery voltage is low |

**Power Supply**

| Internally powered ME equipment | Internally powered ME equipment |

**Power Consumption**

| Approximately 15.0 mA | Approximately 13.0 mA |

**Protection Grade**

| IPX0 | IPX0 |

**Operation Mode**

| Continuous Operation | Continuous Operation |

**Size**

| 3.75”H x 2.25”W x 1”D | 3.75”H x 2.25”W x 1”D |

**Weight (Including Battery)**

| 0.25 lb | 0.25 lb |

**Rheostat (control knob)**

| Controls the current | Controls the current |

7. PANEL TOUCH SWITCHES

- **STANDBY**: No stimulus pulses are generated.
- **TWITCH**: Produce twitch stimulation, which is automatically repeated (one pulse per second), until the button is released. This button can be turned off, by pressing the STANDBY button.
- **100 Hz**: Produce tetanic stimulation when pressed and held down. Tetanic stimulation consists of 100 Hz electrical stimuli.
- **Train-of-Four (TOF)**: Generate four (4) equal intensity single pulses in a period of two (2) seconds. This function can be repeated as often as needed.

**SunStim Plus™**

- **DBS (Double Burst Pulses)**: This stimulation produces two short sequences of 50 Hz tetanic stimuli separated by 750 mSec. This stimulation should not be repeated at intervals of less than twelve (12) seconds.
- **TWITCH**: Produce twitch stimulation, which is automatically repeated (one pulse per second), until the button is released. This button can be turned off by pressing the DBS or TOF button.
- **TETANUS**: Produce rapidly repeated stimulation when pressed and held down. Provided electrical stimuli is set for 100Hz. This default can be changed to 50 Hz, by opening the battery cover and removing the battery. 50 Hz/100 Hz slide switch is located within the battery compartment. You may move the slide to the desired frequency (50 Hz or 100 Hz). Once this step is completed, reconnect the battery and close the battery cover, prior to start using the SunStim Plus™ unit again.

The tetanus stimulation should not be repeated more often than every two (2) minutes, as its affect may fade.

- **Train-of-Four (TOF)**: Generate four (4) equal intensity single pulses in a period of two (2) seconds. This function can be repeated as often as needed.