SHARKDREAMS

USER GUIDE – LIVIT 1

Instructions for pharmacy activation and use

Device activation will enable the LIVIT-1 to begin pharmacy mode operation. In this mode the device will be calibrated for empty and full container weights

Keep the container lid on for empty and full weight calibrations

LINK PRESCRIPTION

To link the device to the prescription, scan the prescription label and device serial number with the barcode scanner.

ATTACH DEVICE

To attach the LIVIT device to prescription bottles, remove the adhesive backing and adhere device to the bottom of a bottle.
See “Calibration.”

DEVICE ACTIVATION

To activate the device, pull the battery tab and verify sufficient battery voltage level.

SHARK DREAMS

USER GUIDE – LIVIT 1

Instructions for pharmacy activation and use

Activating the LIVIT-1 device allows it to begin pharmacy mode operation. In this mode, the device is calibrated by weighing the medication bottles empty and full. It is important to weigh the bottles with the lid on for accurate calibration.

CALIBRATION STEPS

1. Scan prescription label with barcode scanner.
2. Scan LIVIT device serial number with barcode scanner.
3. Verify correct prescription ID, device serial number, and pill count.
4. Pull battery tab to activate device.
5. Verify sufficient battery voltage level on the device.
6. Place empty pill bottle and cap onto the scale. Press “Print”.
7. Verify correct prescription information.
8. Fill bottle according to prescription and place full bottle and cap onto the scale. Press “Print”.
9. Press “Finish” to activate the new prescription.

SHIPPING AND END USER MODES

After calibrating the device, enter the courier tracking number into the calibrator system and the prescription can be shipped to the patient.

Upon receiving the prescription, the patient needs only to take their medication as prescribed.

CONTACT US

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WARNING: Changes or modifications to this device not expressly approved by Shark Dreams could void the user's authority to operate the equipment.

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions: (1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

NOTE: This equipment has been tested and found to comply with the limits for a Class B digital device, pursuant to Part 15 of the FCC Rules. These limits are designed to provide reasonable protection against harmful interference in a residential installation. This equipment generates, uses, and can radiate radio frequency energy and, if not installed and used in accordance with the instructions, may cause harmful interference to radio communications. However, there is no guarantee that interference will not occur in a particular installation. If this equipment does cause harmful interference to radio or television reception, which can be determined by turning the equipment off and on, the user is encouraged to try to correct the interference by one or more of the following measures:

- Reorient or relocate the receiving antenna.
- Increase the separation between the equipment and receiver.
- Connect the equipment into an outlet on a circuit different from that to which the receiver is connected.
- Consult the dealer or an experienced radio/TV technician for help.