

About This Manual

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Product Model: VLS-55Q/VLS-55T/VLS-51T/VLS-51D

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- The use or application of the product or the use of parts or accessories is approved by SonoScape.
- The electrical installation of the relevant room complies with the applicable national and local requirements.
- The product is used in accordance with the instructions for use.

Documentation

Understand the meanings of the following items clearly before reading this manual.

Item	Meaning
	Indicates a potentially hazardous situation which, if not avoided, could result in death or serious injury.
	Indicates a potentially hazardous situation which, if not avoided, may result in malfunction or damage of the system.
NOTE	Indicates precautions or recommendations that should be used in operating the system.
Boldfaced Word	Indicates controls on the control panel, or on-screen objects such as menu items or keys.
Click	Move the cursor to the controls on the display and press the confirm key.

Item	Meaning
>	Select a menu item or a key following the path.

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1 Safety

This chapter describes important information for operating this device. To ensure the safety of both user and patient, read relevant details in this chapter carefully before use.

The operator should be completely familiar with the precautions provided in this manual. Otherwise, the manufacturer is not responsible for the effects on safety, reliability and performance of the device.

1.1 Intended Use

The device is intended to provide illumination for the endoscope in clinics. It is used with the endoscope, the image processor and other peripheral devices provided or approved by the manufacturer.

The device should be used in the medical institution. The operator of the device should be a physician or a medical staff supervised by a physician, both of whom have received sufficient training in clinical endoscopy technology.

1.2 Device Compatibility

The device is used with the image processor (HD-550 series) and the endoscope (EG-550 series, EC-550 series, EB-5H20 series, EG-UC5T, EG-UR5, EC-P550S series, EC-P560 series, EG-P550N, EC-550Z and EG-550Z) provided by the manufacturer.

1.3 Safety Precautions

Read and understand all precautions in this manual before attempting to use the device. Keep this manual with the device at all times. Periodically review the procedures for operation and safety precautions.

1.3.1 Electrical Safety



- Only the personnel authorized or trained by the manufacturer can maintain the device. Any unauthorized personnel should not assemble or disassemble the device.
- Do not operate the device in an atmosphere containing flammable gases such as anesthetic gases, hydrogen or ethanol, because there is a danger of explosion.
- Connect the protective earth conductor only before powering on the device. Disconnect the ground wire only after powering off the device. Otherwise, electrical shock may occur.
- Do not service or maintain the device while it is in use with a patient.
- Do not position the device to make it difficult to disconnect it.
- Connect the device to other electrical equipment by using the potential equalization lead wire before connecting the power plug of the device to an electrical outlet.
- Do not place the multi-socket outlet on the floor.
- Within the environment that is 1.8 meters (6 feet) around a patient, connect peripherals to the auxiliary power outlet which is capable of isolation protection; or, power the peripherals by the auxiliary output cable or the isolation transformer complied with EN/IEC 60601-1, or the power input of the same safety level.

- Do not pour any fluid onto device surfaces, as fluid seepage into electrical circuitry may cause excessive electrical current leakage or device failure. If any water is spilled onto the device carelessly, stop using the device and contact the local distributor immediately.
- The AC power connector plug for the device is a three-prong grounded plug and should never be adapted to any two-prong outlet or by using an adapter. Connect the AC power plug of the multi-socket outlet to a hospital-grade power outlet.
- Disconnect the power cable from the electric outlet to ensure that the device is powered off.
- Do not connect the device and high frequency surgical instrument to the same power outlet. Place the device away from high frequency surgical instrument.
- Place the device away from the laser instrument which has a high starting voltage.
- A spare light source is recommended to be placed in the examination room in case the device appears abnormal.
- The performance of the device and its accessories may be degraded over time. Perform periodic maintenance as described in this manual to ensure the safety of the device.
- Maintain and store the device as described in this manual after use. Improper maintenance and storage may cause cross infection, damage to the device or performance degradation.
- Do not touch the patient and the ports of the device simultaneously. Otherwise, there is a danger of electric shock.



CAUTION

- Select a qualified multi-socket outlet with protective grounding, and ensure that its maximum output power exceeds the required one of the device.
- The multi-socket outlet can only be used to provide power to the recommended peripherals of this system.
- Do not connect other devices to the multi-socket outlet. Otherwise, it may result in interference and overload.
- If the non-medical electrical equipment used with the system is supplied by a multi- socket outlet with separating transformer, connect the AC power plug of the multi- socket outlet to a hospital-grade power outlet. Consult a professional to ensure that the connection complies with the safety standards.
- Accessory equipment connected to analog and digital interfaces must be certified according to the respective EN/IEC standards (for example, EN/IEC 62368-1 for data processing equipment and EN/IEC 60601-1 for medical equipment). Furthermore, all configurations shall comply with EN/IEC 60601-1.

1.3.2 Accessory Safety



WARNING

Only the accessories provided or approved by the manufacturer can be used. Using other accessories may damage the device and cannot achieve the expected performance described in this manual.

1.4 Safety Symbols

The following table is provided for user's identification of important symbols located in labels on the device.

Symbol	Meaning
	Refer to the user manual
	Caution
	Manufacturer

Symbol	Meaning
	Serial Number
	Alternating current
	Stand-by
	OFF (Power)
	ON (Power)
	This product is provided with a CE marking in accordance with the regulations stated in Regulation (EU) 2017/745.
	Authorized representative in the European community
	Type BF Applied Part
	Hot surface
	Equipotentiality
	Fuse
	Caution! Dangerous voltage.
	This symbol indicates that waste electrical and electronic equipment must not be disposed of as unsorted municipal waste and must be collected separately. Please contact an authorized representative of the manufacturer for information concerning the decommissioning of your equipment.
	Model number
	Country and date of manufacture, CHN is the country code of China.
	Medical device
	Unique device identifier

Symbol	Meaning
	Temperature limit
	Humidity limitation
	Atmospheric pressure limitation
	Stacking limit by number
	Fragile, handle with care
	Keep away from rain
	This way up

2 Overview

This medical light source mainly includes the light source and power cable. To ensure the performance of the device, the operator should be completely familiar with the operations and functions of the device.

2.1 Packing

Make sure that all the following items are in the packaging box of the device.

- Light source
- Water bottle
- Power cable
- Fuse

Others: see the Packing List in the packaging box.

2.2 Component Introduction

2.2.1 Front Panel

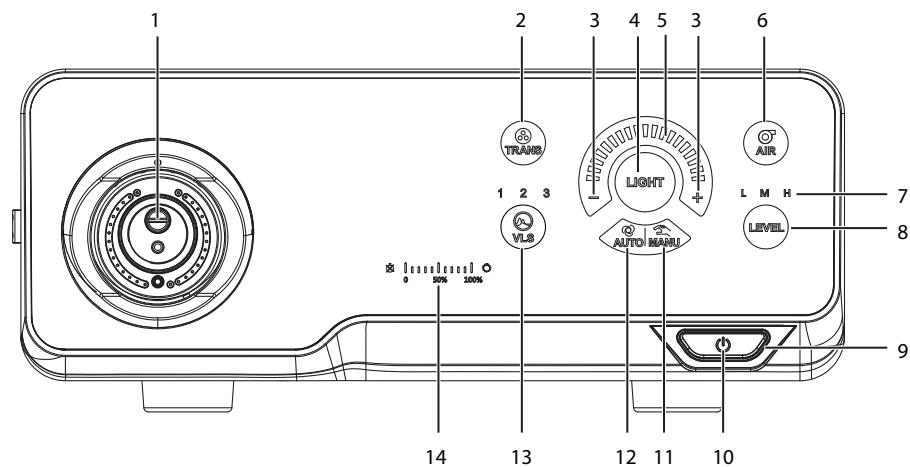


Figure 2-1 Front Panel

No.	Part Name	Description
1	Endoscope Port	Used for connecting the endoscope.
2	TRANS Button	Press it to enable or disable the transillumination feature.
3	⊕ Button	Press them to adjust the brightness.
	⊖ Button	
4	LIGHT Button	Press it to turn on or off the lamps.

No.	Part Name	Description
5	Intensity Indicator	Indicates the brightness level.
6	AIR Button	Press it to enable or disable the air-feeding feature.
7	Air Pump Indicator	Indicates the current air-feeding pressure level (L-Low, M-Medium, H-High).
8	LEVEL Button	Press it to adjust the air pressure.
9	Power Indicator	Indicates the power status of the device
10	 Stand-by Button	Press it to turn on or off the device.
11	MANU Button	Press it to enable manual brightness adjustment.
12	AUTO Button	Press it to enable automatic brightness adjustment.
13	VLS Button	Used to select an illumination mode and the corresponding indicator lights. Illumination modes corresponding to the lighted indicator are determined by the user settings. For detailed setting methods, refer to the image processor user manual.
14	Service Time Indicator	Indicates the accumulated working time of the lamps. This indicator starts blinking when the lamps reach the end of its service time.

2.2.2 Rear Panel

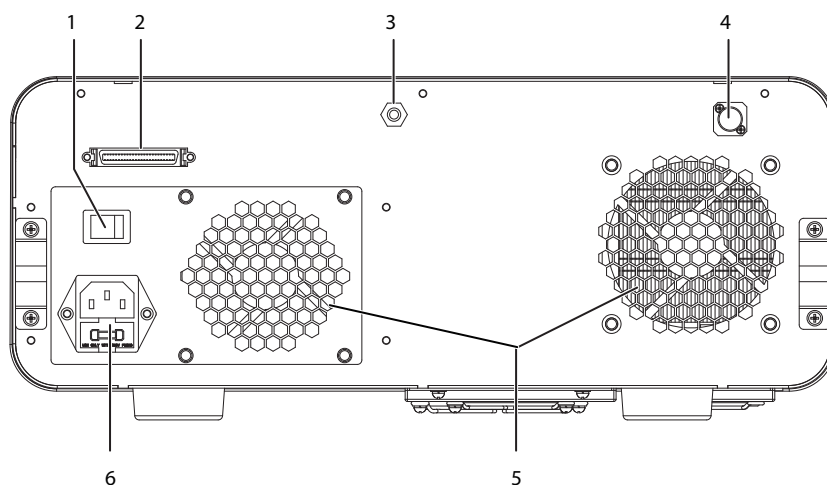


Figure 2-2 Rear Panel

No.	Part Name	Description
1	Main Power Switch	Used for powering on or off the device.
2	LIGHT CONTROL Port	Used for connecting the image processor with the light control cable.
3	Equipotential Terminal	Used for equipotential connection, balancing the protective earth potentials between the device and other electrical equipment.
4	FIBER Port	Used for connecting the image processor.

No.	Part Name	Description
5	Ventilation Holes	Used for ventilation.
6	Power Input Port	Used for connecting the power cable.

2.2.3 Side Panel

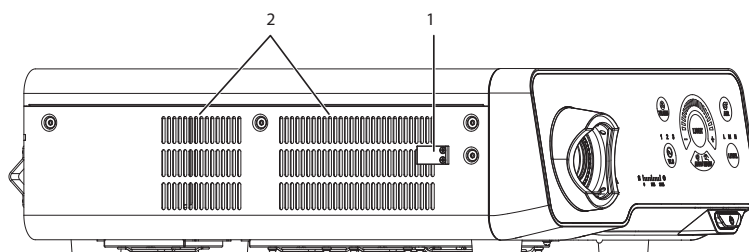


Figure 2-3 Side Panel

No.	Part Name	Description
1	Water Bottle Bracket	Used for holding the water bottle.
2	Ventilation Holes	Used for ventilation.

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3 Preparations

Preparations are necessary before use, which should include device installation, connection, and inspection.

For other devices used with this device, inspect them according to the relevant user manuals. If any malfunction is found, do not use the device.



WARNING

Read through this chapter carefully before using the device to ensure that the installation and connection are correct. Otherwise, damage to the device or personal injury may occur.



CAUTION

- Power off the device and all the peripherals before system connection. Otherwise, damage to the device, malfunction or data loss may occur.
- Only the cables provided by the manufacturer can be used for connection. Otherwise, damage to the device or malfunction may occur.
- Do not block the ventilation holes of the device or place the device in a location without adequate ventilation. Otherwise, the internal heat may cause damage or malfunction.
- Do not place the device where:
 - Water may drip or splash.
 - Flammable and explosive gas exists.
 - It is damp and of high temperature.
 - It is under the sunlight.

3.1 Placing the Device

Place the device on the trolley (sold separately) steadily, and then put the image processor on the device, as shown in Figure 3-1.

NOTE:

- *Ensure that the four feet of the device are positioned within the anti-slip strips.*
- *Refer to the user manual for the installation information of the trolley.*

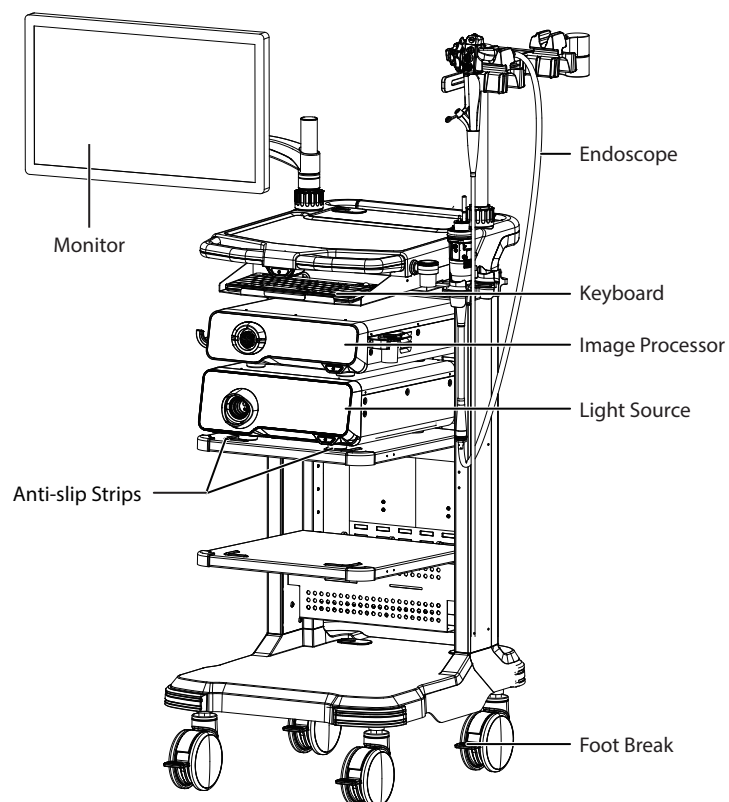


Figure 3-1 Placing the Device

3.2 Connecting the System

After placing the device, connect it to the image processor, the power cable, the endoscope and the water bottle in turn according to Figure 3-2.

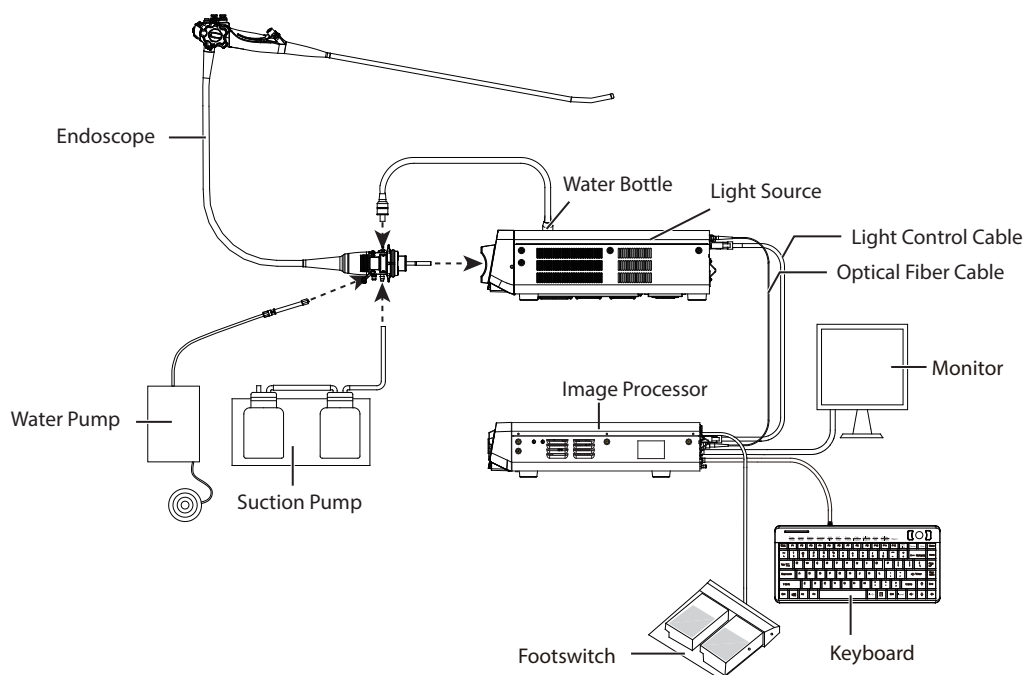


Figure 3-2 Connection the Endoscopy System

3.2.1 Connecting the Image Processor

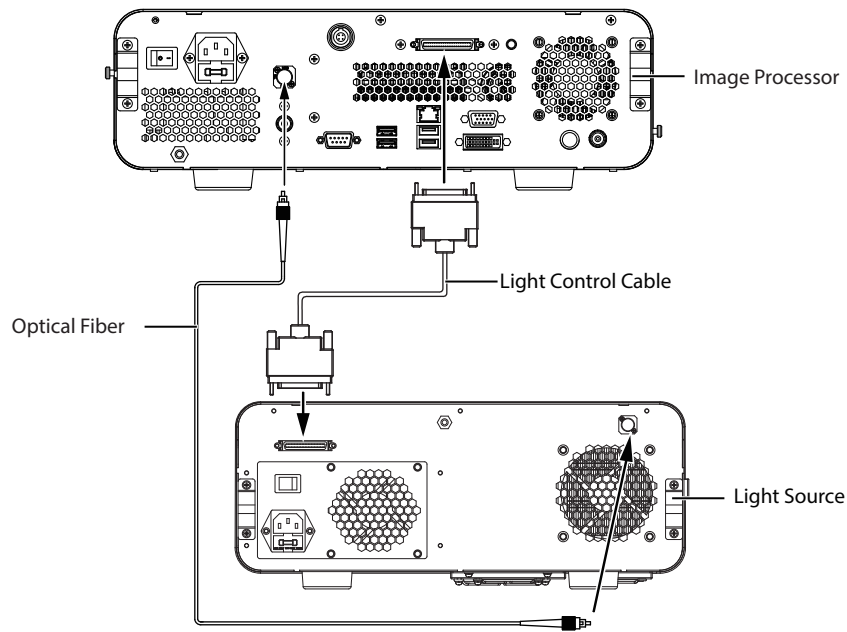


Figure 3-3 Connection the Image Processor

■ To use the light control cable

NOTE:

Only the image processor provided by the manufacturer is allowed to be connected to the light control port.

Connect the light control cable securely to the light control port of the device and the light control port of an image processor.

■ To use the optical fiber cable

Connect the optical fiber cable securely to the optical fiber port of the device and the optical fiber port of the image processor.

3.2.2 Connecting the Power Supply



WARNING Do not bend, drag or twist the power cable excessively. Otherwise, fire or electrical shock may occur.

Perform the following steps.

1. Connect the device to earth or the equipotential terminals of other devices.
2. Connect one end of the power cable to the device and the other end to the AC power outlet.

3.2.3 Connecting the Endoscope

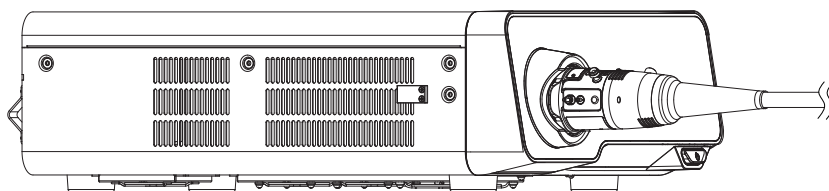


WARNING Wipe the light guide tube of the endoscope carefully and ensure that it is completely dry before connection. Otherwise, electrical shock or damage to the device may occur.

NOTE:

Do not touch the light guide or the endoscope port when the endoscope is just removed from the device. Otherwise, the high temperature may cause skin burns.

Insert the light guide and the air pipe of the endoscope into the endoscope port of the device firmly.

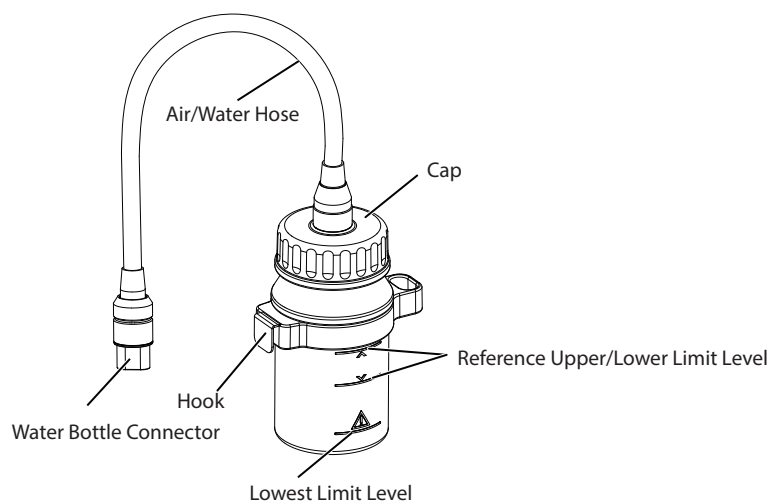


3.2.4 Connecting the Water Bottle



WARNING Do not fill the water bottle with sterile water when it is connected to the device. If water is spilled on or into the device, it may cause electric shock or device damage.

The water bottle is shown as follows.



NOTE:

- Change sterile water in the bottle every day.
- Add sterile water if the water level is under the lowest limit level of the bottle.

Perform the following steps to connect the water bottle.

1. Tighten the bottle cap and fix the water bottle on the bracket, as shown in Figure 3-4.

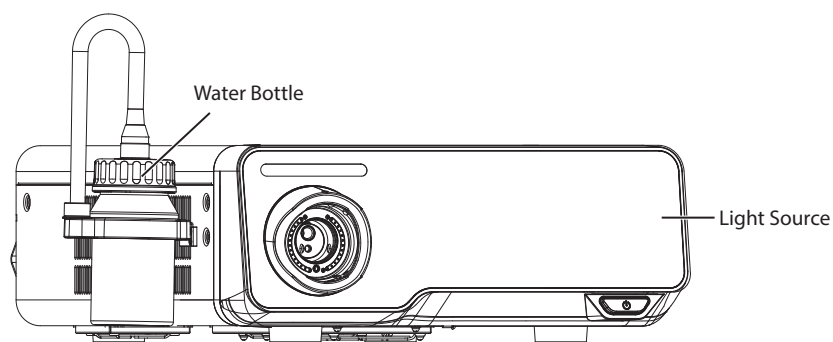


Figure 3-4 Water Bottle Installation

2. Connect the water bottle to the endoscope firmly, as shown in Figure 3-5.

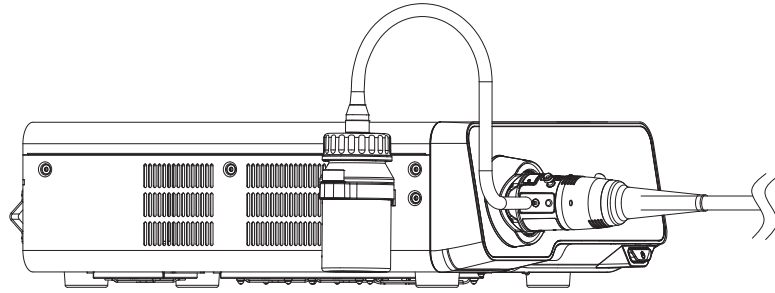


Figure 3-5 Connecting the tube and the connector

3.3 Powering On/Off the Device

■ To power on the device

Perform the following steps.

1. Set the power switch on the rear panel to the **—** position.
2. Press the **⏻** button on the front panel to turn on the device.

■ To power off the device

Perform the following steps.

1. Press the **⏻** button on the front panel to turn off the device. Loosen the button until the indicator lights off.
2. Set the power switch on the rear panel to the **○** position.

3.4 Inspecting the Device

Before each use, strictly follow the descriptions below to inspect the device. Follow the relevant user manuals to inspect the peripherals connected to this device. If any problem exists, refer to Section 5.4 Troubleshooting. If the problem still exists, stop using the device and contact the local distributor.

3.4.1 Inspecting Air-feeding

Perform the following steps.

1. Press the **AIR** button to activate the air-feeding feature. Press the **LEVEL** button repeatedly to set the air pressure to **H**.
2. Immerse the distal end of the endoscope in a container filled with sterile water to a depth of 10 cm.
3. Cover the air/water valve with a finger to feed air. Ensure that bubbles continuously come out from the air/water nozzle, as shown in Figure 3-6. Refer to the relevant endoscope user manual for details.

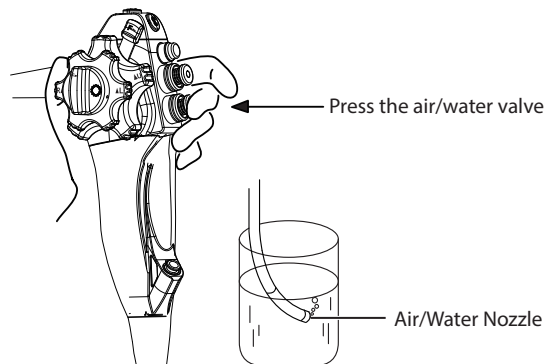


Figure 3-6 Air Feeding Inspection

NOTE:

When the distal end of the endoscope is immersed in the water to a depth less than 10cm, a few bubbles will appear even if the air/water valve is not operated. This is normal.

4. Press the **LEVEL** button repeatedly to adjust the air-feeding pressure level. Ensure that the number of bubbles varies with the air-feeding pressure.
5. Press the **AIR** button to stop air feeding. Ensure that no bubble comes out from the nozzle.
6. Take the distal end out from the sterile water. Check the water feeding feature according to the endoscope user manual.

3.4.2 Inspecting Brightness Adjustment



WARNING Do not stare at the distal end of the endoscope or output light of the device. Otherwise, the strong light may result in injury to eyes.

■ AUTO Mode

Perform the following steps to inspect the brightness adjustment feature in **AUTO** mode.

1. Press the **LIGHT** button on the front panel to turn on the lamps.
2. Press the **VLS** button on the front panel to select the desired illumination mode.
3. Press the **AUTO** button on the front panel to enable the automatic brightness adjustment feature.
4. Press the $\oplus$ or $\ominus$ button to set the brightness to the desired level.
5. Move the distal end of the endoscope up and down. Keep the distance between the distal end and the object (such as a table) within a range of 3 - 100 mm. Ensure that the brightness of the image displayed on the screen does not change obviously.

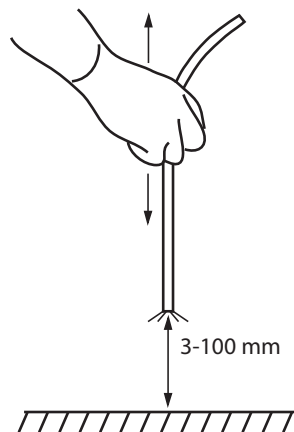


Figure 3-7 Brightness Observation

■ MANU Mode



WARNING To avoid injury to eyes, adjust the brightness to the lowest level necessary for normal clinical observation in **MANU** mode.

Perform the following steps to inspect the brightness adjustment feature in **MANU** mode.

1. Press the **LIGHT** button on the front panel to turn on the lamps.
2. Press the **VLS** button on the front panel to select the desired illumination mode.
3. Press the **MANU** button on the front panel to enable the manual brightness adjustment feature.

4. Press the  or  button to adjust the brightness and ensure that the brightness of the image displayed on the screen changes accordingly.

3.4.3 Inspecting Transillumination

Perform the following steps.

1. Press the **LIGHT** button on the front panel to turn on the lamps.
2. Press the **VLS** button on the front panel to select the desired illumination mode.
3. Press the **TRANS** button on the front panel to enable the transillumination feature.

The light reaches the maximum intensity and starts twinkling. Six to eight seconds later, the light intensity is automatically adjusted to the original intensity.

3.4.4 Inspecting the Illumination Mode Switch Feature

Perform the following steps.

1. Press the **LIGHT** button on the front panel to turn on the lamps.
2. Press the **VLS** button on the front panel to switch the illumination modes and the corresponding indicators lights.

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4 Operations

This manual does not explain or discuss clinical endoscopic procedure. It only describes basic operations and precautions related to the device.

The device will save the current settings automatically when it is powered off.



WARNING Stop the examination immediately if the device works abnormally. Withdraw the endoscope slowly from the patient's body in accordance with the descriptions in the endoscope user manual. Take measures with a reference to Section 5.4 Troubleshooting.

4.1 Turning On the Lamps



WARNING To avoid injury to eyes, set the brightness to the minimum before turning on the lamps.

Press the **LIGHT** button to turn on the lamps.

4.2 Adjusting the Brightness



WARNING To avoid burns, adjust the brightness to the lowest level necessary for normal clinical observation.

For different endoscopy systems and body parts, adjust the brightness of the lamps according to the actual requirement. You can select automatic (**AUTO**) or manual (**MANU**) adjustment mode as required.

■ **AUTO Mode**

Perform the following steps.

1. Press the **AUTO** button on the front panel to enable the automatic brightness adjustment feature.
2. Press the **+** or **-** button to adjust the brightness to the desired level.

Once you set the brightness to the desired level in **AUTO** mode, the light intensity will be automatically adjusted to keep the image brightness roughly the same.

NOTE:

*In **AUTO** mode, the system automatically enters the low brightness mode when the connected endoscope is idle for more than 1 minute.*

■ **MANU Mode**

Perform the following steps.

1. Press the **MANU** button on the front panel to enable the manual brightness adjustment feature.
2. Press the **+** or **-** button to set the brightness to the desired level.

Brightness is fixed once it is set in **MANU** mode. However, the image brightness will be affected by the distance between the distal end and the observed object.

NOTE:

If the endoscope keeps working at highest light intensity for a long time, mist may appear on the image because tissues (such as residual blood and moisture) attached to the lens evaporate due to excessive heat of the light guide. If the examination is interfered under this condition, take out the endoscope, and wipe the distal end with a lint-free cloth dampened with 75% alcohol. Then reinsert the endoscope to continue the examination.

4.3 Selecting the Illumination Mode

These illumination modes can make the details of the gastrointestinal mucosal surfaces more obvious and the images more clear. Different illumination modes can be adopted in different observation distances and their effects differ.

- **White light mode (WL):** This mode features high rendering index and high brightness, which is applicable to the normal examination.
- **Enhanced white light mode (EWL):** This mode can highlight the contour of the mucosal vessels with a color tone similar to the white light mode, which is applicable to the normal examination and lesion identification in the middle and far fields.
- **Spectral focused imaging (SFI):** This mode can highlight the contour of the superficial and middle mucosal vessels, which is applicable to the lesion identification and early cancer screening in the middle and far fields.
- **Versatile intelligent staining technology (VIST):** This mode can highlight the contour of the superficial mucosal vessels and mucosal surface micro-structure, which is applicable to the accurate diagnosis of early cancer in the middle and near fields.

Press the **VLS** button on the front panel to choose the desired illumination mode.

4.4 Using Transillumination



WARNING Use the transillumination feature only when necessary.

You can locate the distal end of the endoscope inside the patient's body by using the transillumination feature.

Press the **TRANS** button to enable the transillumination feature. After this feature is enabled, the light automatically reaches the maximum intensity and starts twinkling. Six to eight seconds later, the light intensity is automatically adjusted to the original intensity.

NOTE:

*You can disable this feature by pressing the **TRANS** button again or other buttons (except the power button) on the front panel.*

4.5 Adjusting Air-feeding Pressure

Perform the following steps.

1. Press the **AIR** button to start air feeding, and press the **LEVEL** button to adjust the air-feeding pressure to the desired level (**L**, **M**, **H**).
2. Perform the air-feeding operation according to the endoscope user manual.
3. Press the **AIR** button again to stop air feeding.

4.6 Turning Off the Lamps

Press and hold the **LIGHT** button for about 2 seconds to turn off the lamps.

5 Maintenance

To ensure the safety and functionality of the device, follow the descriptions in this chapter to clean and maintain the device.

During use, any serious incident that has occurred in relation to the device should be reported to the local distributor and the competent authority of the Member State.



- To avoid electrical shock and damage to the device, power off the device and disconnect it from the AC power outlet before cleaning.
- To ensure the device performance, maintain the device periodically (not more than a year), and ensure that the grounding of the device is correct and complies with the safety requirements.
- Only the local distributor of the manufacturer is allowed to replace the illuminator. Power off the device and unplug the power cable before replacement. Otherwise, electric shock or device damage may occur.

5.1 Cleaning the Device



- Ensure that the device is completely dry before use to avoid electric shock.
- Wear personal protective equipment during the cleaning. Otherwise, the blood, mucous membrane and other potential source of infection adhered to the device may result in cross-contamination.
- Do not wipe the endoscope port or other ports. Otherwise, deformation or bad contact may occur to the pins inside the ports.

NOTE:

If the device is contaminated, perform the cleaning procedure below immediately after use. If cleaning is delayed, the debris may dry up and become difficult to be cleared up.

■ To clean the surface

Perform the following steps.

1. Power off the device and disconnect it from the AC power outlet.
2. In case of potentially infectious substances such as blood or body fluid, use a lint-free soft cloth dampened with neutral detergent solution to clean the surface.
3. Use a lint-free soft cloth dampened with 75% alcohol to clean the surface. Ensure that the device is completely dry.

■ To clean and disinfect the water bottle

Clean and disinfect or sterilize the water bottle according to the water bottle user manual.

NOTE:

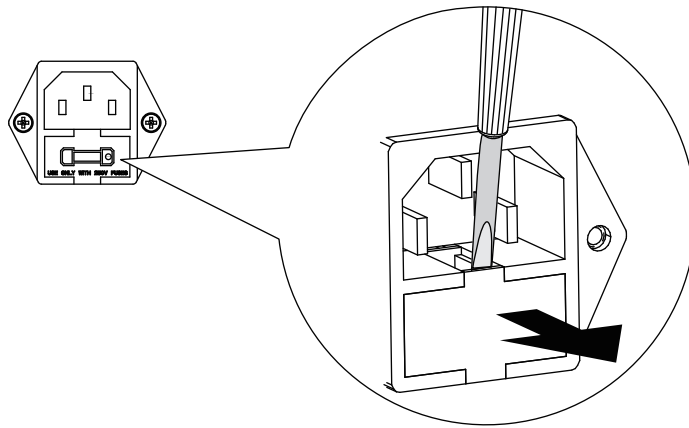
Change the sterile water in the water bottle every day.

5.2 Replacing the Fuse

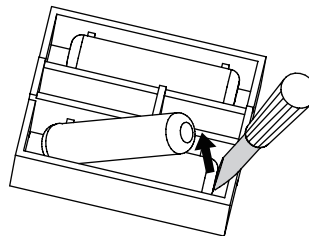
Perform the following step.

1. Power off the device and disconnect the power cable.

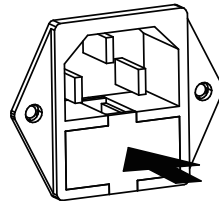
2. Take out the fuse box with a slotted screwdriver.



3. Replace the blown fuse.



4. Push the fuse box back into the place and fasten the screws on the fuse box.



5. Connect the power cable, and press the power button to power on the device. Ensure that the power indicator lights. Contact the local distributor of the manufacturer if the device cannot be powered on.

5.3 Storing the Device

NOTE:

- To avoid malfunction, do not excessively bend, pull, twist or squeeze the power cable during storage.
- Store the device in the environment of good ventilation and avoid direct sunlight.

Perform the following steps.

1. Power off the device and disconnect the power cable.
2. Disconnect all the peripherals from the device.
3. Place the device on a clean and level surface at room temperature.

5.4 Troubleshooting

The troubleshooting should be performed by qualified technical personnel. If a problem persists after the troubleshooting, stop using the device immediately and contact the local distributor of the manufacturer for repair.

Description	Inspection Item	Solution
The prompt that the endoscope is connected is not displayed; or, white screen appears.	Light control cable	Ensure that the light control cable is connected properly.
Blurred screen appears after the endoscope is connected.	Optical fiber cable	Ensure that the optical fiber cable is connected properly.
		Wipe the front end of the optical fiber cable with an alcohol cotton ball, and then insert the optical fiber cable.
The device emits only the white light or the bluish violet light.	Device temperature	Ensure that the rear fan of the case works normally.
		Ensure that ventilation holes are not blocked.
The device cannot be connected to the endoscope.	Endoscope and the device	Ensure that they are compatible with each other.
The device cannot be powered on.	Power cable	Ensure that the power cable is connected properly.
	Fuse	Replace the fuse.
The buttons on the front panel cannot work properly.	Front panel	Ensure that the front panel is clean and free of water drops.
	Power cable	Ensure that the power cable is well connected to the ground.
The indicators cannot work properly.	Front panel	Check if the indicators can respond to the pressing of the buttons on the front panel. Restart the device if the indicators cannot respond.
The intensity of light is low.	Lamps	Check if the lamps have reached to the service time.
No light is emitted from the distal end.	Connection of the endoscope and the device	Ensure that the endoscope is properly connected to the device.
The intensity buttons do not work.	Intensity indicator	Check if the intensity is adjusted to the maximum or the minimum level.
The image is too bright or too dim.	Connection of the endoscope and the device	Ensure that the endoscope is properly connected to the device.
	Transillumination	The device automatically restores to normal state 6 - 8 seconds later.

Description	Inspection Item	Solution
The image color is abnormal.	SFI mode or VIST mode	Check if the device is in the SFI mode or the VIST mode. If yes, press the VLS button to switch to the WL mode or EWL mode.
	White balance	If the current illumination mode is WL mode, perform the white balance feature.
Unable to feed water/air.	Connection of the endoscope and the device	Ensure that the endoscope is properly connected to the device.
	Air pump	Ensure that the air pump is turned on and noise can be heard during working.

5.5 Disposing of the Device

The manufacture date of the device is on the label, and the expected service life of the device is five years with normal use. You should dispose of the device or its accessories in accordance with local laws or regulations. For detailed disposal information, consult the local distributor. The manufacturer is not responsible for any device content or accessories that have been discarded improperly.

The expected service life of the lamps and air pump are shown as follows.

Component	Expected Service Life
Lamps	20,000 hours, continuous operation
Air pump assembly	5,000 hours, continuous operation

5.6 Customer Service

Only the service personnel of or authorized by the manufacturer can service the device. Any feedback or inquiries concerning our product or services should be directed to the manufacturer.

Contact address: 2F, 12th Building, Shenzhen Software Park Phase II, Keji Middle 2nd Road, Nanshan District, Shenzhen, 518057, Guangdong, China

Tel: +86-755-26722890

E-mail: service@sonoscape.net

Appendix A Specifications

Endoscope port		Apply to the video endoscope with a $\Phi 59.5$ mm connector or a $\Phi 32.5$ mm connector.
White light mode (WL)	Infrared rays cut-off performance	In the wavelength range between 300 nm and 1700 nm, the ratio of radiant flux to light flux should not be greater than 6 mW/lm
	Max. output light flux	- VLS-55Q and VLS-55T: ≥ 1400 lm, allowance: - 10% - VLS-51T and VLS-51D: ≥ 800 lm, allowance: - 10%
Lamps	Four LED lamps with different colors.	
Light intensity level		1 - 19
Air-feeding	Air-feeding pressure level	L, M and H
	Pressure range	45 - 65 kPa
	Max. flux	- H Level: 3.7 - 5.5 L/min - M Level: 3.0 - 4.7 L/min - L Level: 2.0 - 3.7 L/min
Power supply	Rated voltage	AC 100-240 V
	Frequency	50 Hz/60 Hz
	Input power	300 VA
	Fuse	T5AH 250 V
Safety Type	Type of protection against electric shock	Class I
	Degree of protection against electric shock	Type-BF applied part
	Degrees of protection against harmful liquid	Non-waterproof enclosed equipment
	According to the degree of safety of application	The equipment is not suitable for use in the presence of a flammable anesthetic mixture with air, oxygen or nitrous oxide.

Environment Conditions	Operation	Environment Temperature	0°C - 40°C
		Relative humidity	30% - 80%
		Atmosphere Pressure	700 hPa - 1060 hPa
	Storage	Environment Temperature	-5°C - + 40°C
		Relative humidity	30% - 80%
		Atmosphere Pressure	700 hPa - 1060 hPa
	Transportation	Environment Temperature	-20°C - +55°C
		Relative humidity	20% - 90%
		Atmosphere Pressure	700 hPa - 1060 hPa

Appendix B EMC Guidance and Manufacturer's Declaration



- The device is suitable for use in professional healthcare facility environment. Do not use it in domestic establishments and those directly connected to the public low voltage power supply network that supplies buildings used for domestic purposes.
- Do not use this device around strong electric field, electromagnetic field (e.g. MRI scan room) and mobile wireless communication devices. Using the device in an improper environment may cause malfunction or damage.
- Only the peripherals (such as endoscope, image processor, etc.) provided or recommended by the manufacturer can be used. Using other devices may increase RF radiation and degrade device the performance of anti-electromagnetic interference.
- Use of this device adjacent to or stacked with other equipment should be avoided because it could result in improper operation. If such use is necessary, this device and the other equipment should be observed to verify that they are operating normally.
- Use of accessories and cables other than those specified or provided by the manufacturer of this device could result in increased electromagnetic emissions or decreased electromagnetic immunity of this device and result in improper operation.
- Portable RF communications equipment (including peripherals such as antenna cables and external antennas) should be used no closer than 30 cm (12 inches) to any part of the device, including cables specified by the manufacturer. Otherwise, degradation of the performance of this device could result.

B.1 Electromagnetic Emissions

Emissions Test	Compliance
Conducted and Radiated RF Emissions CISPR 11	Group 1 Class A
Harmonic Distortion IEC 61000-3-2	Not Applicable
Voltage Fluctuations and Flicker IEC 61000-3-3	Not Applicable

B.2 Electromagnetic Immunity

Immunity Test	Compliance Level
Electrostatic Discharge (ESD) IEC 61000-4-2	±8kV Contact, ±2kV, ±4kV, ±8kV, ±15kV Air
Radiated RF EM fields IEC 61000-4-3	3V/m
Electrical Fast Transient and bursts IEC 61000-4-4	±2kV for power supply lines, ±1kV for SIP/SOP ports

Immunity Test	Compliance Level
Surges IEC 61000-4-5	$\pm 1\text{kV}$ differential mode $\pm 2\text{kV}$ common mode
Conducted Disturbances IEC 61000-4-6	3V 6V at ISM bands
Voltage dips and interruptions IEC 61000-4-11	0% U_T for 0.5 cycle at 0° , 45° , 90° , 135° , 180° , 225° , 270° and 315°
	0% U_T for 1 cycle Single phase: at 0°
	70% U_T for 25 cycle Single phase: at 0°
	0% U_T for 250 cycle
Power frequency Magnetic field IEC 61000-4-8	30A/m