HyBase V8 Classic Operating table

Versatility beyond Expectation





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With increasingly complex surgical procedures, today's surgical environment requires a table that instantly adapts to all the versatility to provide the highest level of care.

- An extreme load capacity of up to 460 kg.
- All positions are permitted up to overall load of 250 kg.
- The longitudinal shift up to 350mm for free to access to C-arm.
- 600mm lowest table position for neurosurgery, while 1,050 mm highest table position for orthopedics.
- 12 electric movements, such as table top up/down, slope and tilt, adapting to various surgical needs.
- Bluetooth hand control for unobstructed control in operating rooms.
- Independent manual override system for dual safety.
- Triple-layer decompression pad to redistribute the patient's weight.



Flexibility for multiple disciplines

Meeting the needs of different surgical positions

Modular Design

The modular design of the tabletop is tailored for various surgical discipline needs, which can also lower the cost for hospitals as they only need to buy the specific module, instead of a new table, for an additional surgical type.

Surgical Disciplines





















Flexibility for multiple disciplines

Cater to increasing endoscopic surgeries

Higher Imaging Quality

With the development of minimally invasive surgery (MIS), requirement for better images during procedures arises. HyBase V8 Classic allows clear and high-quality images that are crucial for today's MIS, cardiothoracic, orthopedic, and neurosurgical procedures.

Huge Imaging Access





(Optional) Carbon fiber operating plate provides extreme imaging space.

The carbon fiber tabletop ensures good radiolucent imaging. With a wide imaging access and better radiolucency, HyBase V8 Classic is an optimal solution for cardiovascular surgery.

Bidirectional longitudinal shift



Head-end shift:160 mm Foot-end shift:190 mm

Built-in body elevator



Build-in body elevator(120 mm) for kidney surgery.

Carbon fiber tabletop



Radiolucent carbon fiber tabletop provides better imaging.

Optimal Stability

With a load capacity of up to 460 kg, the table is designed to accommodate the highest safety and stability. All positions are permitted up to overall load of 250 kg.



Better Exposure to Operating Field

With gas spring assisted, HyStirrup allows simultaneous adjustment for abduction and lithotomy while reducing pressure under the popliteal fossa.



Lithotomy position with HyStirrup-3 leg support.



Electric brake provides higher stability.





Smart backlit keypad makes operating table status and function keys visible in the dark.



IPX5 protection against water jets from any direction.

Safety for various positions

Enhance patient safety

Manual Override System

Manual Override System allows table articulation movements and release the brake in the event of primary control or power/battery malfunction, through an independent hydraulic circuit. Manual operations of all basic electro-hydraulic functions can be actuated by using a foot pump combined with the knob next to it.





Step 1: Rotate the function icon to the position indicated by the arrow.





Step 2: Adjust the operating table by stepping on the foot pump.

Multi-layer Decompression Pad

Mindray's HyBase V series operating table introduces a new decompression mattress. The multi-layer composite mattress technology fully considers the body shape of the human and can be applied to patients with BMI between 17-35. Especially for heavy patients, the buffer layer can greatly reduce the contact between the body and the support layer, improve the support performance and further balance the pressure.



Compared with current mainstream products, our mattress has a more than 10% optimizing decompression effect, which can better disperse the pressure and reduce the risk of pressure ulcer.

Normal pad





BMI=35.8

Color-coded Indication Technology

During the tilt or slope adjustment, a color-coded indicator appears on the screen to show the angle and issue a timely warning when it comes to excessive Trendelenburg and reverse Trendelenburg positions.



HyBase V series operating table pad

Efficiency in every step

Improve efficiency in the operating room

Intuitive icons with reasonable layouts





Corded/wireless hand control



Override panel (Backup control panel)

Easy to Adjust



Featherweight head plate to lighten your work.

Easy to Use



With the One-button design, it's convenient to remove or install the head and leg plates.

Four double-swivel castors make it easier to move the table toward any directions.

Easy to Clean



Bellows-free design.



Gas-spring assisted leg plates with foldable design.





Flat stainless-steel base cover.

HyBase V8 Classic Operating table

Flexibility for multiple disciplines, the modular design of the tabletop is tailored for various surgical discipline needs.

Safety for various positions, meeting the requirements for increasing patients safety.

Efficiency in every step, four double-swivel castors make it easier to move the table in any direction.

- An extreme load capacity of up to 460kg.
- All positions are permitted up to overall load of 250kg.
- The longitudinal shift up to 350mm for free to access to C-arm.
- 600mm lowest table position for neurosurgery, while 1,050mm highest table position for orthopedics.
- 12 electric movements, such as table top up/down, slope and tilt, adapting to various surgical needs.
- Bluetooth hand control for unobstructed control in operating rooms.
- Independent manual override system for dual safety.
- Triple-layer decompression pad to redistribute the patient's weight.





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_	410mm 625mm	

Dimensions

Length (Main unit)		Base cover	
Height without padding	600-1050mm	Base thickness	140mm
		Length	1100mm





Head plate		Back plate	
Head plate up	Max. 45°	Back plate up	Max. 80°
Head plate down	Max. 90°	Back plate down	Max. 40°





Trendelenburg Position		Lateral tilt	
Trendelenburg	Max. 36°	Left tilt	Max. 26°
Reverse Trendelenburg	Max. 36°	Right tilt	Max. 26°





Leg plate		Manual body elevator(optional)	
Leg plate up	Max. 20°	Body elevator up	Max.120mm
Leg plate down	Max. 90°	Body elevator width	86mm





	Longitudinal slide(optional)		Four double-swivel/universal castors	
	Longitudinal shift toward head end	Max. 160mm	Diameter	75mm
	Longitudinal shift toward foot end	Max. 190mm		
ĺ	Maximum longitudinal shift	350mm		





Flex Position		Reflex Position	
Flex Position	220°	Reflex Position	110°
Reverse Trendelenburg	20°	Trendelenburg	30°
Back plate down	40°	Back plate up	70°



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HyBase V8 Classic

Operating Table

Operator's Manual

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- all installation operations, expansions, changes, modifications and repairs of this product are conducted by Mindray authorized personnel;
- 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.

WARNING

- This equipment must be operated by skilled/trained clinical professionals.
- It is important for the hospital or organization that employs this equipment to carry out a reasonable service/maintenance plan. Neglect of this may result in machine breakdown or personal injury.

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- Malfunction of the instrument or part whose serial number is not legible enough.
- Others not caused by instrument or part itself.

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Preface

Manual Purpose

This manual contains the instructions necessary to operate the product safely and in accordance with its function and intended use. Observance of this manual is a prerequisite for proper product performance and correct operation and ensures patient and operator safety.

This manual is based on the maximum configuration and therefore some contents may not apply to your product. If you have any question, please contact us.

This manual is an integral part of the product. It should always be kept close to the equipment so that it can be obtained conveniently when needed.

Intended Audience

This manual is geared for clinical professionals who are expected to perform daily operations of the product, maintain and troubleshoot the product and learn how to use the product.

Illustrations

All illustrations in this manual serve as examples only. They may not necessarily reflect the setup or data displayed on your equipment.

Conventions

- Italic text is used in this manual to quote the referenced chapters or sections.
- Bold text is used to indicate the screen texts.
- \blacksquare \rightarrow is used to indicate operational procedures.

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1.1 Safety Information

WARNING

 Indicates a potential hazard or unsafe practice that, if not avoided, could result in death or serious injury.

CAUTION

• Indicates a potential hazard or unsafe practice that, if not avoided, could result in minor personal injury or product/property damage.

NOTE

 Provides application tips or other useful information to ensure that you get the most from your product.

1.1.1 Warning

WARNING

- The equipment is to be used for its intended purposes only. Do not use it for other purposes.
- The equipment is to be installed by personnel authorized by Mindray only.
- The equipment is to be operated by trained personnel only.
- Before using the equipment, be sure to read and fully understand this manual.
- Keep this manual in the vicinity of the equipment so that it can be obtained conveniently when needed.
- Ensure the ground is flat and level before installing the operating table.
- Route, wrap and secure the cables to avoid inadvertent disconnection, stumbling and entanglement.
- Loosely laid cables may cause stumbling. Fix the cables reliably when laying them.

- The overall load of this operating table is 460kg in normal patient orientation. Be sure to take those conditions into consideration when using the operating table.
- The overall load of this operating table is 250kg in reverse patient orientation. Be sure to take those conditions into consideration when using the operating table.
- If accessories are used, ensure the weights of the accessories installed on the operating table and of the patient add up no more than 460kg in normal patient orientation or 250kg in reverse patient orientation.
- The overall load of this operating table is 460kg (250kg in reverse patient orientation). However, in case of certain special conditions, the load capacity is further limited. Be sure to take those conditions into consideration when using the operating table. If you have any questions, contact the customer service department authorized by Mindray or your local distributor.
- Transferring the patient inappropriately may tip over the operating table. Be sure to transfer the patient from either side of the operating table. Do not transfer the patient from either the head or the foot end.
- Be sure to transfer or position the patient under medical staff's direction.
- Before positioning the patient, ensure the patient's body width is no greater than the width of the table top. Neglect of this may cause personal injury. Positioning the patient improperly may tip over the operating table. Be sure to place the patient in the right direction. Do not place the patient's trunk on the leg plate.
- When placing the patient, ensure the patient's center of gravity is as close to the column as possible. Otherwise, the operating table may tip over.
- Improper positioning may damage the patient, especially in case of excessive tilted or sloped position. Ensure the patient is properly positioned and check if the position deviates at any moment when using the operating table.
- Ensure the patient is properly positioned and keep a close eye on the patient during the whole surgery to avoid damaging the patient's respiratory system, nervous system or circulating system.
- Before positioning the patient, ensure the castors are properly locked.
- Do not use such devices as cell phones, radio equipment and MR equipment around the operating table.
- The operating table cannot be used in the presence of a flammable anesthetic mixture with air or with oxygen or with nitrous oxide.
- To avoid explosion hazard, do not use the equipment in the presence of oxygen-rich atmospheres, flammable anesthetics, or other flammable agents.
- Use and store the equipment within the specified temperature, humidity, and altitude ranges.

- Before putting the equipment into operation, the user must verify that the equipment, connecting cables and accessories are in correct working order and operating condition.
- If the integrity of the external protective conductor in the installation or its arrangement is in doubt, run the operating table from its internal batteries.
- Do not touch the patient and live parts simultaneously. Otherwise, patient injury may occur.
- To avoid infection, be sure to cover the table-top pad with a sterile surgical drape before using the operating table.
- Do not use wet surgical drapes on the operating table.
- Each time before using the operating table, be sure to select the proper function keys according to the patient orientation. Otherwise, personal injury may occur.
- Be sure to secure the patient before moving or adjusting the operating table when a patient lying on the operating table.
- When lowering the table top, ensure that there are neither the operator's feet nor any objects on the base (underneath the column). Otherwise, the operator may be pinched or the equipment may be damaged.
- Before lowering the table top, ensure that there are neither the operator's feet nor any objects underneath the column.
- Before locking the operating table, ensure that there are neither the operator's feet nor any objects underneath the base. Otherwise, personal injury may occur.
- Before unlocking the castors, grasp the operating table to avoid unexpected movement.
- Before moving the mobile operating table, ensure that the operating table is not bent and the longitudinal shift is cancelled.
- The speed of table-top movement can change depending on the patient's weight. Closely watch the table top during adjustment.
- If the patient weight exceeds 250kg, do not slope the table top by more than 15° and do not tilt the table top to the left or right by more than 5°. Do not adjust the back plate up by more than 70° and down more than 10°. Otherwise, the patient may slip off the table and cause injury.
- Ensure the patient is properly secured especially in case of excessive tilted or sloped position when adjusting the operating table or patient position.
 Otherwise, personal injury or device damage may occur.
- If the longitudinal shift is operated when using an x-ray, be sure to restore the operating table to 0 position before the surgery.
- Be sure to observe the joints when adjusting the head plate, back plate and leg plate. Otherwise, the patient or medical staff may be pinched.

- Be sure to observe the positions of the operating table, accessories, patient and other objects when adjusting or moving the operating table to avoid personal injury or device damage resulting from collision.
- When using such medical devices as electrosurgical units, defibrillators or defibrillator monitors, be sure to keep the patient from touching the metallic parts of the operating table or of the accessories. Moreover, ensure the patient is not lying on a wet surgical drape or a conductive pad. Otherwise, the patient may be burned.
- Devices connected to the equipment must meet the requirements of the applicable IEC standards (e.g. IEC 60950 safety standards for information technology equipment and IEC 60601-1 safety standards for medical electrical equipment). The system configuration must meet the requirements of the IEC 60601-1 medical electrical systems standard. Any personnel who connect devices to the equipment's signal input/output port are responsible for providing evidence that the safety certification of the devices has been performed in accordance to the IEC 60601-1. If you have any questions, please contact Mindray.
- Make sure the electrical installation of building complies with the requirements of IEC 60364-7-710.
- Be sure to operate the equipment as instructed in this manual. Warranty does not cover damage caused by non-compliance.
- Device service or maintenance should be completed by the authorized personnel only. Warranty does not cover damage caused by unapproved service or maintenance.
- The operating table can function safely and effectively only if it is maintained properly and regularly. Be sure to maintain the operating table as instructed in this manual.
- Be sure to only use accessories made by Mindray on the operating table. Accessories produced by other manufacturers must not be used as they may cause personal injury. If accessories of other manufacturers must be used, their use should be expressly permitted by Mindray.
- Before using accessories, be sure to read the operator's manual thoroughly.
- Be sure to use the pad provided by Mindray only. Otherwise, personal injury or device damage may occur.
- Disconnect the operating table from the mains power before opening the housing.
- No modification of this device is allowed.
- Do not use agents containing alcohol where high-frequency equipment is being used. Neglect of this may cause fire hazard.
- There may be a risk of infection if the operating table is under the complicated and changeable medical conditions. Follow the hospital

requirements and refer to "Cleaning and Disinfection" chapter to clean/ disinfect the operating table.

- The software was developed in compliance with IEC 62304. The possibility of hazards arising from software errors is minimized.
- The equipment software copyright is solely owned by Mindray. No
 organization or individual shall resort to modifying, copying, or exchanging
 it or to any other infringement on it in any form or by any means without due
 permission.
- At the end of its service life, the equipment, as well as its accessories, must be disposed of in compliance with the local regulations. If you have any questions concerning disposal of the equipment, please contact Mindray.
- For disposal of parts and accessories, unless otherwise specified, follow local regulations regarding disposal of hospital waste.
- Inspect and replace the battery at regular service intervals. If you have any questions, please contact Mindray.
- Dispose of the package material as per the applicable waste control regulations. Keep it out of children's reach.

1.1.2 Caution

CAUTION

- Exercise caution when transporting or moving the equipment. Do not damage it during transportation or movement.
- Always install or carry the equipment or accessories properly to avoid damage caused by drop, impact, strong vibration or other mechanical force.
- Before moving the operating table, ensure the path is clear. Do not damage the castors or other parts.
- When moving the operating table, ensure there is not any object on the table top.
- When moving the operating table, be sure to use both hands to adjust/ control the movement. Keep the operating table from colliding with other things.
- If the table top has descended to a low position, exercise caution when adjusting the table top. Keep the table top from colliding with the base or the ground.
- If the head/leg plate has been positioned downward, exercise caution when adjusting the table top. Keep the head/leg plate from colliding with the base or the ground.
- When sloping the table top or adjusting the leg plate, ensure they do not collide with the column or the base.

- Do not put any object on the operating table base. Neglect of this may cause equipment damage.
- When adjusting the table top, ensure the cord of the corded hand control is not jammed by the joints.
- Be sure to lock the operating table and adjust the table top to 0 position when it is not in use.
- When adjusting the operating table, ensure the suspended parts (such as leg plates, arm boards) do not collide with other things inside the operating room.
- When adjusting the operating table, ensure the accessories or the surgical drapes are not jammed by the joints. Otherwise, equipment damage may occur.
- Check the screws for components and parts regularly to prevent them falling off.
- Be sure to use spare parts supplied by Mindray only.
- Connector pins with an ESD warning symbol should not be touched and no connections should be made between these connectors without implementing ESD protective measures. Such precautionary procedures may include antistatic clothing and shoes, the touch of a ground stud before and during connecting the pins or the use of electrically isolating and anti-static gloves. All staff involved in the above shall receive instruction in these ESD precautionary procedures.

1.1.3 Note

NOTE

• Contents of this manual are based on the full configuration of the operating table. Some of them may not apply to your operating table. Use your operating table according to the actual configuration. If you have any questions, contact the customer service department authorized by Mindray or your local distributor.

1.2 Equipment Symbols

Some symbols may not appear on your equipment.

Symbol	Description	Symbol	Description
(Blue)	Refer to instruction manual/booklet	\triangle	Caution
UDI	Unique Device Identification		Class II equipment
	Protective earth (ground)	$\stackrel{\frown}{\forall}$	Equipotentiality
IPX5	Protection against water jets	IPX8	Protection against the effects of continuous immersion in water
+	Body elevator	⊙/Ò	On/Off
(Red)	No sitting	SN	Serial number
REF	Catalogue number	\sim	Alternating current
	Direct current		Anti-static
M	Date of manufacture		Manufacturer
<u> </u>	This way up	Ť	Keep dry

Symbol	Description	Symbol	Description
Ţ	Fragile, handle with care		Stacking limit by number
kg max	Stacking limit by mass		Temperature limit
	Humidity limitation		Atmospheric pressure limitation
£\$	Recovery/recyclable	((😦))	Non-ionizing electromagnetic radiation
★	Type B applied part	X	Separate collection for electrical and electronic equipment
(6	This product is provided with a CE marking in accordance with the regulations stated in Regulation(EU) 2017/745 concerning Medical Devices.	MD	Medical Device
EAC	Unified circulation mark indicates that products marked them passed all specified in the technical regulations of the Customs Union of the procedure for the assessment (confirmation) of conformity and complies with the requirements applicable to all the products technical regulations of the Customs Union.	EC REP	Authorized representative in the European Community
	General warning sign		
(Yellow)			

1.3 Warning Label

Warning label for longitudinal shift: The label is located above the column cover. With an overall load of more than 250kg, do not operate the longitudinal shift.



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2.1 Intended Use

2.1.1 Intended Purpose

The operating table is intended to support and position patients immediately before, during and after surgical procedures, as well as for examination and treatment.

2.1.2 Intended Users

The operating table may only be operated by medically trained staff.

2.1.3 Intended Patient Population

The operating table is intended for patients who are in surgical, examination and treatment procedures.

2.1.4 Intended Medical Conditions

The operating table is used in healthcare facility environment.

2.1.5 Contraindications

None.

2.1.6 Side-effects

According to the conclusion of clinical evaluation and residual risk evaluation, for the intended patients, there are no known side effects that can occur during or after the use of the medical device. And there is no need for the operator to make extra preparations. Thus, no residual risk associated with using the medical device should be disclosed.

WARNING

- Contents of this manual are based on the full configuration of the operating table. Some of them can not apply to your operating table. Use your operating table according to the actual configuration. If you have any questions, contact the customer service department authorized by Mindray or your local distributor.
- The operating table is to be operated by trained personnel only. Nonauthorized or untrained personnel cannot operate the operating table.

2.2 Main Components

The operating table consists of the main unit (including the table top, column, base, hydraulic system, electric system), pad and control unit, and has optional head plate, leg plate and upper back plate.

2.2.1 Main Unit



2.2.2 Control Unit

(1)Corded hand control

(2)Wireless hand control (3)Foot switch





2.3 Definition

2.3.1 Slope/Tilt/Longitudinal Shift



2.3.2 0 Position

0 position: the table top is horizontal and does not have any longitudinal shift.

2.3.3 Zero Longitudinal Shift

The longitudinal shift distance of the table top is zero.

2.3.4 Adjustable Part

Adjustable parts of the product, including release lever, release handle, grip of the release bracket and so on, are indicated in blue which are easy for users to recognize.

2.3.5 Overall Load

The "overall load" results from putting together patient weight and accessory weight. The "overall load" is the load which can be placed on the table top. Restrictions might result from the components or accessories for which other overall loads may be applicable, or from special patient positions.

2.3.6 Normal/Reverse Patient Orientation

The patient orientation (normal or reverse) depends on the position of the patient on the table top with regard to the operating table base (1).

WARNING

- Incorrect patient orientation may cause adjustment of the operating table in a direction that is not intended. Check if the patient orientation is correct prior to making any adjustments.
- The patient orientation is indicated in the status bar on the screen of the hand control. See 3.3.5.13 Status Indicator.

2.3.6.1 Normal Patient Orientation

The upper part of the patient's body is located above the operating table base (1).



2.3.6.2 Reverse Patient Orientation

The patient's legs are located above the operating table base (1).



2.4 Prompt Tone

Acoustic signal of the operating table is for prompt, detailed types and cases are listed below:

Prompt Tone	Acoustic Signal Type	Case
Normal prompt tone	Beep once	e.g. desired movement completed, normal operation
Arrival confirmation tone	Beep twice	e.g. final position reached, wireless hand control successfully connected
Risk prompt tone	Beep intermittently and slowly	e.g. workable button pressed under unlocking
Collision prompt tone	Beep intermittently and quickly	e.g. impending collision
Use-prohibited tone	Beep continuously	e.g. movement prohibited or limited
Timing tone	Beep five times	e.g. The wireless hand control is being connected after Trendelenburg button on the override panel has been pressed.

2.5 Abbreviations

Abbreviations	In full
AC	Alternating current
DC	Direct current
EMC	Electromagnetic compatibility
EU	European Union
Info.	Information
RF	Radio frequency
UPS	Uninterrupted power supply
3.1 Connecting Potential Equalization Pin

WARNING

 When using the operating table together with other devices, connect their potential equalization pins together to eliminate the potential difference between them. Otherwise, personal injury or device damage may occur.



To connect the potential equalization pin:

- 1. Open the waterproof cover (1) on the base.
- 2. Connect one end of the equipotential cable to the potential equalization pin (2) of the operating table.
- 3. Connect the other end of the equipotential cable to the potential equalization pin of another device.

• Disconnect the equipotential cable before moving the operating table.

3.2 Connecting to Mains Power

3.2.1 Power Button



Press the power button (1) to control the power-on or power-off of the operating table.

WARNING

 Press the power button to power off the operating table in case of an emergency.

CAUTION

• Power off the operating table by pressing the power button if the operating table is not to be used for a long time (e.g. for weekend or holidays). Otherwise, battery damage may occur.

3.2.2 Charging Batteries



To keep good performance, fully or nearly fully discharged batteries should be charged as soon as possible. The operating table is charged automatically once connected to AC power.

For new batteries, it takes about 10 hours to fully charge them. Fully charged rechargeable batteries are sufficient for about one week of operating table operation.

WARNING

- Check if the power cord has any sign of wear or damage before use. Damaged cord must be replaced immediately.
- Always use the accompanying power cord delivered with the operating table.
- Before connecting the equipment to AC power, check whether the voltage and frequency ratings are the same as those indicated on the AC adapter and manual.
- Do not touch the power connector with the wet hand. Eliminate the liquid residue inside of or at the surroundings of the AC power connector of the equipment and power cord connectors. Otherwise, personal injury or equipment damage may occur.
- If the integrity of the external protective conductor in the installation or its arrangement is in doubt, run the operating table from its internal batteries.

• To avoid the risk of electric shock, the operating table must only be connected to mains power with protective earth.

NOTE

- The operating table is disconnected from the mains power by means of detaching the plug. Do not locate the operating table in a place where it is difficult to operate the mains plug.
- It is recommended that the operating table be run from its internal batteries.
- In order to prevent the charging of batteries during surgeries, it is recommended the rechargeable batteries be charged after surgeries and on a daily basis. On principle, deep discharge of the rechargeable batteries should be avoided.
- Ensure the battery capacity is sufficient if the operating table is powered by internal batteries. Charge the batteries if necessary.
- If the batteries cannot work normally, contact the customer service department authorized by Mindray or your local distributor.
- If the run time of the fully charged batteries is noticeably shortened, contact the customer service department authorized by Mindray or your local distributor.
- The battery must only be installed and replaced by service personnel trained and authorized by Mindray. Installation or replacement by untrained personnel may cause personal injury (e.g. batteries overheated, ignited or exploded). Batteries are recommended to be replaced every three years under normal use.
- Disconnect the power cord before moving the operating table.

Follow the steps below to connect the power cord:

- 1. Open the waterproof cover.
- 2. Connect the power cord to the power connector (2) of the operating table.
- 3. Connect the power cord to the mains power outlet.

Follow the steps below to detach the power cord:

- 1. Detach the power cord from the mains power outlet.
- 2. Detach the power cord from the power connector of the operating table.

3.2.3 Indicator on the Base

AC indicator (1) and battery indicator (2) are located on the base. Indicator status is shown below:



Indicator	Status	Description
AC indicator	Green	The operating table is connected to mains power.
Battery indicator (not connected to	Four battery bars are green.	The battery capacity of the operating table is no less than 90%.
AC)	Three battery bars are green.	The battery capacity of the operating table is no less than 70% and less than 90%.
	Two battery bars are green.	The battery capacity of the operating table is no less than 35% and less than 70%.
	One battery bar is red.	The battery capacity of the operating table is less than 35%.

Indicator	Status	Description	
Battery indicator (connected to AC)	All battery bars cyclically change.	The batteries of the operating table are being charged and the charging capacity is less than 35%.	
	The first battery bar is green and other three bars cyclically change.	The batteries of the operating table are being charged and the charging capacity is no less than 35% and less than 70%.	
	The first two battery bars are green and other two bars cyclically change.	The batteries of the operating table are being charged and the charging capacity is no less than 70% and less than 90%.	
	The first three battery bars are green and the fourth bar flashes.	The batteries of the operating table are being charged and the charging capacity is no less than 90% and less than 100%.	
	Four battery bars are green.	The batteries of the operating table are fully charged.	
Battery indicator	The second battery bar is red and flashes.	The operating table has an error, contact the customer service department authorized by Mindray.	
	The third battery bar is red and flashes.		
	The fourth battery bar is red and flashes.		

NOTE

 Once the operating table is powered on by pressing the power button and not connected to mains power, the battery indicator will show current capacity and go out after 20 seconds.

3.3 Hand Control

Operate the corded hand control and wireless hand control in the same way.



(10)Trendelenburg	(11)

(11)Tilt to left

(12)Leg plate up (Reverse patient orientation)

(13)Leg plate down (Reverse (14)Longitudinal shift patient orientation) toward foot end

(15)Lock

(16)0 position

WARNING

• Ensure the operating table is in view of users when the wireless hand control is used. Otherwise, personal injury may occur.

NOTE

- When adjusting the operating table, ensure the cord of the corded hand control is not jammed by joints.
- Switch off the control unit after all the adjustments are completed to avoid misoperation.
- Ensure all functions are normal before each use.
- If different functions are triggered on the same control unit, the motion of the operating table except the switch-off will be stopped immediately.
- Do not perform adjustments with two control units at the same time. If functions are triggered by multiple control units, then all motions except the switch-off of the operating table will be stopped immediately and will resume only after none of the control units has been activated. If the override panel is triggered, it takes priority over other control units.

3.3.1 Connecting /Disconnecting Corded Hand Control

The sockets for connecting the corded hand control are located on both sides of the column.

- To connect the corded hand control: insert the plug into the socket with the red mark (1) on the corded hand control cable aligned with that (2) on the operating table socket. Ensure the plug is properly seated.
- To disconnect the corded hand control: grasp the front end of the corded hand control cable and pull it out straight down.



3.3.2 Attaching Hand Control

CAUTION

• The hand control which is attached to the side rail may slip off the side rail or catch the cable during table top adjustment. Take the hand control out of the side rail during the adjustment.

Attach the bracket (1) to the side rail (2).



The hand control can be attached onto the lug (3) of the head plate if equipped.



3.3.3 Button Operations

After the operating table is switched on, press and hold the button until the desired position is reached and then release the button.

3.3.3.1 Button Symbol

Symbol	Description	Symbol	Description
٩	Standby	0 +	One-button to 0 position
Ê	Unlock	a	Lock
i	Longitudinal shift toward head end	Ð	Longitudinal shift toward foot end
	The circled means back plate up.		The circled means back plate down.
	The circled means leg plate up.		The circled means leg plate down.
\bigcirc	Tilt to right		Tilt to left

Symbol	Description	Symbol	Description
\bigcirc	Table top up	\bigtriangledown	Table top down
	Reverse Trendelenburg	N	Trendelenburg

3.3.4 Screen Control

The screen will show movement part, direction and value when the control unit is used to adjust the table top movements. Screen interface can be operated by control buttons below.



(1)Select/Confirm

(2)Select

(3)Select

(4)Select/Confirm

3.3.4.1 Interface Symbol

Symbol	Description	Symbol	Description
Ĥ	Operating table locked		Operating table unlocked
	Battery of wireless hand control	Ţ	Operating table batteries

Symbol	Description	Symbol	Description
-	0° tilt		Tilt to left
•	Tilt to right	i	0° slope
ľ	Trendelenburg position		Reverse Trendelenburg position
←I →	Zero longitudinal shift		Longitudinal shift toward head end
→	Longitudinal shift toward foot end	ľ	0° back plate angle
	Back plate up	1	Back plate down
<	Select	>	Select
Ð	Back	i	Flex position
>>	Reflex position		Normal patient orientation
RI	Reverse patient orientation		Save
\checkmark	Select/Confirm		Delete
ବ୍ତି	User Settings		General warning sign
C D	Disconnected with operating table	æ	Connected with operating table

3.3.4.2 Main Interface-Flex Position/Reflex Position



(4)Flex position (5)Reflex position

(1)Status bar

Once the hand control is switched on, it enters the main interface and displays the following from the top down:

- Status bar: displaying the locked status of the operating table, battery status of both the operating table and wireless hand control, operating table name. See 3.3.5.13 Status Indicator.
- Parameter area: displaying real-time status including tilt angle, slope angle.
- Function selection area: main interface displays flex position and reflex position which can be accomplished by pressing and holding the button below.

3.3.4.3 Normal/Reverse Patient Orientation



(1)Normal patient orientation

(2)Reverse patient orientation

Enter the selection interface of normal or reverse patient orientation by pressing the selective button (or)) → press and hold the button below the screen for two seconds to reach the normal or reverse patient orientation.

3.3.4.4 Quick Save/Call Memory Position



(1)Save

(2)Select/Confirm

NOTE

• Before calling a memory position, store the patient position with the hand control.

3.3.4.5 User Settings



(1)User settings

- Enter the user settings interface (1) by pressing the selective button (\checkmark or >). \rightarrow press the button () to enter detailed user settings.
- You can call/save memory position, check parameters of operating table, set the screen brightness, select the language of the hand control, name the operating table, operate the bluetooth connection, enable the smart backlit keypad, set switch-on mode and foot switch function, check the operating table battery and software information and restore factory settings. Please see 3.3.7 User Settings for details.

3.3.4.6 Prompt Message

The operating table has not only prompt tone, but prompt message on the screen. See details below:

Common prompt message: indicated in white

Prompt message of normal use, e.g. position reached, normal or reverse patient orientation set, operating table locked or unlocked, operating table connected, position saved, battery status of operating table and wireless hand control.



■ Low-risk prompt message: indicated in yellow

Noticeable prompt message, e.g. final position reached, operating table disconnected, low battery, locking error, collision risk.



■ High-risk prompt message: indicated in red

Especially noticeable prompt message, e.g. excessive sloped angle.



- Error indication
 - If the following prompt message appears, contact the customer service department authorized by Mindray or your local distributor.



 If the symbol A appears on the status bar, some error has occurred. Contact the customer service department authorized by Mindray or your local distributor.

3.3.5 Functions

3.3.5.1 Switch-on

- Press the standby button U to enter the switch-on status.
- Press any button to switch on the operating table if it is configured with such a function.

3.3.5.2 Unlock /Lock

- 1. Lock: press and hold the lock button for two seconds. It takes about six seconds to lock the operating table.
- 2. Unlock: press and hold the unlock button for two seconds. It takes about six seconds to unlock the operating table.
- 3. A progress bar will appear on the screen during the process of unlocking or locking. A prompt message will also appear once unlocking or lock is accomplished.

Lock the castors:





Unlock the castors:



WARNING

- Before locking the operating table, ensure that there are neither the operator's feet nor any objects underneath the base. Otherwise, personal injury or equipment damage may occur.
- The adjustment of table top up, tilt or longitudinal shift is restricted after the operating table castors are unlocked.

NOTE

• A prompt message as [Long press 2s] will appear on the screen if you do not unlock or lock the operating table within specified time.

ri A Long press 2s

NOTE

Interruption of locking or unlocking procedure or sudden power-off may cause both locking and unlocking indicators flashing after the operating table is switched on again. Press the lock or unlock button to continue the operation.

One-button to 0 position 3.3.5.3

Press and hold 0 position button •••• to adjust the operating table to 0 position.

Longitudinal Shift 3.3.5.4

- Longitudinal shift toward foot end: press and hold the button 1.
- Longitudinal shift toward head end: press and hold the button 😇 2.

3.3.5.5 Table Top Up/Down









Lock:

Table top down: press and hold the button () ().

3.3.5.6 Tilt

- 1. Tilt to left: press and hold the button 🦉
- 2. Tilt to right: press and hold the button D

3.3.5.7 Slope

- 1. Trendelenburg position: press and hold the button
- 2. Reverse Trendelenburg position: press and hold the button

WARNING

 Ensure the patient is properly secured in case of sloped position when adjusting the operating table. Otherwise, personal injury or device damage may occur.

3.3.5.8 Leg Plate Up/Down

In the reverse patient orientation, the leg plate can be adjusted up or down.



1. Leg plate up: press and hold the button



2. Leg plate down: press and hold the button

3.3.5.9 Back Plate Up/Down

- 1. Back plate up: press and hold the button
- 2. Back plate down: press and hold the button

3.3.5.10 Smart Backlit Keypad

If the smart backlit keypad is enabled, buttons on the hand control are divided by backlit keypad. Identify the available functions by backlit buttons. See the details below:

- If a certain movement reaches its final position, its button backlight will go off.
- Under the unlocked status, function of the backlit button is enabled while function of the button without backlight is disabled.

3.3.5.11 Color-coded Indication

During the tilt or slope adjustment, a color-coded indication will respectively appear on the screen if the angle of the single movement is no less than 15° or 30°.







≥ 15°, <30°:

≥ 30°:

NOTE

• The default yellow-coded and red-coded values are 15° and 30° respectively. If you need to change the setting, see 3.3.7.8 More Settings.

3.3.5.12 Switch-off

Press the standby button 0 to enter the standby status or it will enter the standby status automatically when the hand control is idle for more than 20 seconds.

NOTE

The default automatic switch-off time for the hand control is 20 seconds. If you need to change the time setting, contact the customer service department authorized by Mindray or your local distributor.

3.3.5.13 Status Indicator

Observe the battery status of the wireless hand control or operating table, locking/ unlocking status, normal/reverse patient orientation in the status bar.

Indicator	Status	Description
Battery indicator of the wireless hand control		Red. The battery capacity of the wireless hand control is no more than 10%, which indicates the wireless hand control needs to be charged immediately.
		Red. The battery capacity of the wireless hand control is in the range of approximately 10%-25%. The screen will prompt low battery message.
		Grey. The battery capacity of the wireless hand control is in the range of approximately 25%-50%.
		Grey. The battery capacity of the wireless hand control is in the range of approximately 50%-75%.
		Grey. The battery capacity of the wireless hand control is in the range of approximately 75%- 100%.
		Battery bars cyclically change in green during charging, which will stop if the battery is fully charged.

Indicator	Status	Description
Operating table battery indicator	Ţ	Red. The battery capacity of the operating table is less than 35%. The screen will prompt low battery message.
	T	Grey. The battery capacity of the operating table is no less than 35% and less than 70%.
	Ţ	Grey. The battery capacity of the operating table is no less than 70% and less than 90%.
	T	Grey. The battery capacity of the operating table is no less than 90%.
Operating table battery indicator		All battery bars are green and cyclically change.The batteries of the operating table are being charged and the charging capacity is less than 35%.
		The first battery bar is green and other three bars cyclically change. The batteries of the operating table are being charged and the charging capacity is no less than 35% and less than 70%.
		The first two battery bars are green and other two bars cyclically change. The batteries of the operating table are being charged and the charging capacity is no less than 70% and less than 90%.
		The first three battery bars are green and the fourth bar flashes. The batteries of the operating table are being charged and the charging capacity is no less than 90% and less than 100%.
		Green. The batteries of the operating table are fully charged.
Operating table locking indicator	đ	The operating table is locked.

Indicator	Status	Description
Operating table unlocking indicator	C	The operating table is unlocked.
Normal patient orientation indicator	-	The operating table is set to normal patient orientation.
Reverse patient orientation indicator	RI	The operating table is set to reverse patient orientation.

3.3.6 Wireless Hand Control

3.3.6.1 Bluetooth Connection of Wireless Hand Control

If the wireless hand control switched on is not connected to the operating table, follow the steps on the screen to continue the connection. Once the wireless hand control is connected, the screen shows the message [Connected].







The connection may fail and a prompt message below will appear. Repeat the connection procedure until it succeeds.



The connection may disconnect after a successful connection. Repeat the connection procedure until it succeeds.

3.3.6.2 Charging Wireless Hand Control

Observe the current battery capacity of the wireless hand control by the battery indicator in the status bar. See details in **3.3.5.13 Status Indicator**. Follow the steps below to charge the battery if it is low. It takes about 10 hours to fully charge an automatically switched-off wireless hand control.



(1) Mobile charging station

(2) Indicator of mobile charging station

- 1. Connect the mobile charging station to the mains power.
- 2. Switch off the wireless hand control and insert it into the mobile charging station (1) and the wireless hand control is automatically charged. Observe the charging status by the indicator (2):

Indicator	Status	Description
Indicator of mobile charging station	Red	As soon as the mobile charging station is connected to AC power, it is dangerous to let the metal get close.
	Green	The wireless hand control is being charged.

The run-time of the battery depends on the equipment operation. The fully charged wireless hand control is sufficient for four hours of continuous operation under normal use.

NOTE

- In order to avoid empty battery of the wireless hand control during the surgery, we recommend you store the wireless hand control in the mobile charging station when it is not in use.
- Be sure to use the mobile charging station to charge the wireless hand control.
- Ensure the wireless hand control is reliably placed into the mobile charging station, which means the indicator of mobile charging station is green.

- Do not place the objects such as metal to the charging area.
- Do not use the wireless hand control during charging to adjust the operating table.
- Ensure the battery power is sufficient if the wireless hand control is to be used. Charge the battery if necessary.
- Use the battery in the wireless hand control at least once every month to extend battery life. Charge the battery before its capacity is depleted.
- Inspect and replace the battery at regular service intervals. Life expectancy
 of a battery depends on how frequent and how long it is used. For a properly
 maintained and stored lithium-ion battery, its life expectancy is
 approximately three years. More aggressive usage will shorten the life
 expectancy. Replacing the lithium-ion battery every three years is
 recommended.
- In case of a battery failure, contact Mindray service personnel for battery replacement.

3.3.7 User Settings

3.3.7.1 Calling Memory Position

Select [Call Memory Position] via the selective button (< or) press the button

 $oldsymbol{O}$ below the symbol $oldsymbol{V}$ to enter the interface—select the desired position— press

and hold the button \mathbf{O} \rightarrow the operating table is to be adjusted until the desired memory position is reached.



NOTE

 If the patient orientation of the selected memory position does not match that of the current operating table, adjust it to the right patient orientation before calling memory position.

3.3.7.2 Saving Memory Position

Select [Save Memory Position] via the selective button (✓ or ➤)→press the

button **O** below the symbol **I** to enter the desired interface.

TI D	,
Call Memory Position	
Save Memory Position	
Operating Table Parameters	
Screen Brightness	
Language	
Operating Table Name	
Bluetooth Connection	
More Settings	
Operating Table Battery Info.	
Software Version	
Restore Factory Settings	
5 < > ·	~

TÎ 🖬			 •
	4		<u>••</u> 0° 0°
R	eal-time pos	sition previ	0° ew
Add Memo	ory Position		
< 13	14	+	— >
Ð	<	>	\checkmark

◆ Delete a memory position: select the button via the selective button
 (or >) → press the button oblow the symbol of to delete the position.



Overwrite a memory position: select the position to be overwritten via the selective button (✓ or) → press the button (below the symbol ↓ to overwrite it.



• The status bar will display the corresponding icon if the memory position is saved under the reverse patient orientation.

3.3.7.3 Checking Parameters of Operating Table

Select [**Operating Table Parameters**] via the selective button (\checkmark or \rightarrow) \rightarrow press the button \bigcirc below the symbol \checkmark -check the parameters.

TI D	,			
Call Memory Position				
Save Memory Position				
Operating Table Parameters				
Screen Brightness				
Language				
Operating Table Name				
Bluetooth Connection				
More Settings				
Operating Table Battery Info.				
Software Version				
Restore Factory Settings				
5 < > ·	~			



3.3.7.4 Selecting Screen Brightness

Select [Screen Brightness] via the selective button (\checkmark or >) \rightarrow press the button **O** below the symbol \checkmark \rightarrow select one of the four levels: 25%, 50%, 75%,100%.

TI D				
Call Memory Position				
Save Memory Position				
Operating Table Parameters				
Screen Brightness				
Language				
Operating Table Name				
Bluetooth Connection				
More Settings				
Operating Table Battery Info.				
Software Version				
Restore Factory Settings				
5 < >	~			



3 - 35

3.3.7.5 Selecting Language

Select [Language] via the selective button (\checkmark or >) \rightarrow press the button \bigcirc below

the symbol \checkmark to enter the language selection interface \rightarrow select the desired language.

±∎ A					
Call Memory Position					
Save Memory Position					
Operating Table Parameters					
Screen Brightness					
Language					
Operating Table Name					
Bluetooth Connection					
More Settings					
Operating Table Battery Info.					
Software Version					
Restore Factory Settings					
5 < >	\checkmark				

TI 🖬			.			
Language						
中文 Chinese			0			
English English			•			
Español Spanish						
Franç. French						
Italiano Italian						
~						
A	<	>	\checkmark			
3.3.7.6 **Naming Operating Name**

Select [**Operating Table Name**] via the selective button (\checkmark or >) \rightarrow press the button **O** below the symbol **I** to enter the name interface.

The name of the operating table can have four characters at most. Save the name by pressing [Save] and it will show on the status bar.

TI 🖬 📖			
Call Memory Position			
Save Memory Position			
Operating Table Parameters			
Screen Brightness			
Language			
Operating Table Name			
Bluetooth Connection			
Bluetooth Connection More Settings			
Bluetooth Connection More Settings Operating Table Battery Info.			
Bluetooth Connection More Settings Operating Table Battery Info. Software Version			
Bluetooth Connection More Settings Operating Table Battery Info. Software Version Restore Factory Settings			



3.3.7.7 Bluetooth Connection

Select [**Bluetooth Connection**] via the selective button (✓ or)→press

the button \bigcirc below the symbol \checkmark to enter the bluetooth connection interface \rightarrow confirm whether to be connected to a new operating table \rightarrow connect it by following the steps in **3.3.6.1** Bluetooth Connection of Wireless Hand Control after confirmation.





3.3.7.8 More Settings

Select [**More Settings**] via the selective button (✓ or)→press the

button **O** below the symbol **I** to enter the interface to enable or disable the smart

backlit keypad, set switch-on mode and foot switch function.



Smart Backlit Keypad



-xâ â	.
Smart Backlit Keypad	
Enabled	\circ
Disabled	
5 < >	\checkmark

Switch-on Mode

Enter [Switch-on Mode] via the selective button (\checkmark or >) \rightarrow select [Any button] or [Standby button] \rightarrow press the button \bigcirc below the symbol \checkmark .

TÎ			
	Switch-	on Mode	
Any butto	n		
Standby I	button		
A	<	>	\checkmark

Foot Switch Function

See the operator's manual of foot switch for details.

Color-coded Indication

Enter [Color-coded Indication] via the selective button (\checkmark or \triangleright).

- 1. Select [Enabled] or [Disabled] to enable or disable the function.
- 2. Once enabled, press [Parameter Setting] to access the indication value setting.



TE B	
Parameter Sett	ing
Tilt to Left/Right (0*-26*) Yellow-coded value	≥ 1 5 °
Trend./Reverse Trend. (0) Yellow-coded value)*-36*) ≥ 1 5 °
Red-coded value	≥ 3 0 °
0 1 2 Resto	re defaults
• ১	\checkmark

3.3.7.9 Operating Table Battery Information

Select [**Operating Table Battery Info.**] via the selective button (\checkmark or \triangleright) \rightarrow press the button \bigcirc below the symbol \checkmark to check the battery status: [**Healthy**] or

[Replacement recommended].



NOTE

• When the screen shows the message [Replacement recommended], contact the customer service department authorized by Mindray or your local distributor.

3.3.7.10 Software Version

Select [Software Version] via the selective button (✓ or >)→press the button

O below the symbol it to check the software information.

TE D	.		
Call Memory Position			
Save Memory Position			
Operating Table Parameters			
Screen Brightness			
Language			
Operating Table Name			
Bluetooth Connection			
More Settings			
Operating Table Battery Info.			
Software Version			
Restore Factory Settings			
5 < > ·	~		



3.3.7.11 Restore Factory Settings

Select [**Restore Factory Settings**] via the selective button (< or) press the

button \bigcirc below the symbol \checkmark \rightarrow the screen prompts the message [Restore]

factory settings?]. Restore factory setting after the confirmation.

TÍ 🖬			
Call Memory Position			
Save Memory Position			
Operating Table Parameters			
Screen Brightness			
Language			
Operating Table Name			
Bluetooth Connection			
More Settings			
Operating Table Battery Info.			
Software Version			
Restore Factory Settings			
5 < > ·	~		



Override Panel 3.4

The override panel is located on the column, of which the buttons are the same as those

of the hand control except the switch-on button 🧿.



After the operating table is switched on, press and hold the switch-on button and the desired adjustment button at the same time and do not release them until the desired position is reached. Detailed buttons are shown below:



NOTE

- The override panel is for emergency operation.
- The normal/reverse patient orientation is not effective when the override panel is used. If the table top is set to reverse patient orientation, the button operation is the same as that of normal patient orientation. Watch every button adjustment.
- Do not perform adjustments with two control units at the same time. If functions are triggered by multiple control units, then all motions except the

switch-off of the operating table will be stopped immediately and will resume only after none of the control units has been activated. If the override panel is triggered, it takes priority over other control units.

3.4.1 Functions

3.4.1.1 Switch-on

Press the button 🕑 to switch on the operating table.

3.4.1.2 Lock/Unlock

- 1. Lock: press and hold the **o** and **b** buttons for two seconds. It takes about six seconds to lock the operating table.
- 2. Unlock: press and hold the O and D buttons for two seconds. It takes about six seconds to unlock the operating table.

WARNING

- Before locking the operating table, ensure that there are neither the operator's feet nor any objects underneath the base. Otherwise, personal injury or equipment damage may occur.
- The adjustment of table top up, tilt or longitudinal shift is restricted after the operating table is unlocked.

3.4.1.3 One-button to 0 position

Press and hold both 💿 and 😶 buttons to restore the operating table to 0 position.

3.4.1.4 Longitudinal Shift

- 1. Longitudinal shift toward head end: press and hold 🧿 and 🚭 buttons.
- 2. Longitudinal shift toward foot end: press and hold 💿 and 😜 buttons.

3.4.1.5 Table Top Up/Down



WARNING

 During the table top down process, there is a risk of crushing and shearing to the feet or objects placed on the base. Before lowering the table top, ensure that there are neither the operator's feet nor any objects underneath the column.

3.4.1.6 Tilt

3.4.1.7



- 1. Trendelenburg: press and hold O and S buttons.
- 2. Reverse Trendelenburg: press and hold 🗿 and 🥏 buttons.

WARNING

• Ensure the patient is properly secured in case of sloped position when adjusting the operating table. Otherwise, personal injury or device damage may occur.



3.4.1.9 Indicator on Override Panel

Indicators on the override panel are described below:

Indicator	Status	Description
Switch-on indicator	Green	The operating table is switched on.
Locking indicator	Green and flashing	Locking in progress
	Green	Locked
Unlocking indicator	Red and flashing	Unlocking in progress
	Red	Unlocked
Locking indicator and unlocking indicator	Locking indicator (green) and unlocking indicator (red) flash at the same time.	The operating table is under the intermediate status of locking and unlocking.

3.5 Foot Pump (optional)

Foot pump can be used to adjust the table-top movements. The function can be realized by the pedal (1) and knob (2) at the base. 12 movements at most can be accomplished by the foot pump.



To use the foot pump,

- 1. Turn the knob until the arrow above is aligned with the desired movement label.
- 2. Step the pedal repeatedly until the desired function has been reached.

NOTE

• When using the foot pump to lock the castors, step the pedal and do not release your foot until the locking is finished. It takes about six seconds to fully lock the operating table.



3.6 Body Elevator with Hand Crank (optional)

To use the body elevator with the hand crank:

- 1. Install the hand crank (1) onto either side of the hole on the operating table.
- 2. Raise or lower the body elevator by rotating the hand crank.



3.7 Remote Control Interface (optional)

The operating table can provide a remote control interface to realize the remote control function. See the communication protocol for details.

3.8 Manual Brake Release

In case of the control unit and override panel failure, execute the manual brake release.

- 1. Loosen two screws (2) fixing the base cover on both sides. Lift the base cover (1).
- 2. Insert the proper Allen key into two adjustment holes (3) of one castor and turn it clockwise until it can no longer be turned. Then insert the Allen key into two adjustment holes that are in a diagonal castor and then turn it clockwise. Finally adjust the holes of other two castors until the operating table is unlocked.



NOTE

- Keep the screws fixing the base cover for reinstallation use.
- 3. Reinstall the base cover after the operating table is unlocked.

3.9 Foot Switch

Connect or disconnect the foot switch in the same way as the corded hand control. See **3.3.1** Connecting /Disconnecting Corded Hand Control. For more details, see the operator's manual of foot switch for details.

3.10 Table-top Modules

WARNING

- When removing/installing the head plate, upper back plate or leg plate, be sure to grasp it with both hands. Otherwise, it may fall and cause personal injury and/or property damage.
- When using the head plate, upper back plate or leg plate, be sure to cover it with a sterile surgical drape to avoid infection.

Be sure to use the pad delivered with head plate, upper back plate or leg
plate. Otherwise, patient injury may occur during the table top movement.

3.10.1 Head Plate

3.10.1.1 Installation

- 1. Hold the head plate (1) with both hands. Insert both joints (2) fully into the mounting points (3) until a click is heard.
- 2. Slightly pull the head plate and check if it is reliably installed.



3.10.1.2 Removal

- 1. Hold the head plate with both hands and press the release buttons (4) on both front ends.
- 2. Pull the head plate out slowly without tilting.



3.10.1.3 Angle Adjustment

- 1. Hold the head plate with one hand and pull the grip (5) in the middle of the release bracket with the other hand. Raise or lower the head plate to the desired position.
- 2. Release the bracket.
- 3. Pull the head plate to check whether it has been adjusted properly.



WARNING

 When lowering the head plate, be sure to adjust it slowly and carefully. Ensure the head plate keeps stable during the adjustment to avoid patient injury.

NOTE

 Do not install such devices as arm boards, anesthesia frames and infusion holders on the head plate. Otherwise, the head plate may move unexpectedly and cause personal injury or equipment damage.

3.10.2 Upper Back Plate

3.10.2.1 Installation

- 1. Hold the upper back plate (1) with both hands. Insert both joints (2) fully into the mounting points (3) until a click is heard.
- 2. Slightly pull the upper back plate and check if it is reliably installed.



3.10.2.2 Removal

- 1. Hold the upper back plate with both hands and press the release levers (4) on both front ends.
- 2. Pull the upper back plate out slowly without tilting.



3.10.3 Leg Plate

3.10.3.1 Installation

- 1. Hold the leg plate (1) with both hands. Insert the joint (2) fully into the mounting point (3) until a click is heard.
- 2. Slightly pull the leg plate and check if it is reliably installed.



3.10.3.2 Removal

- 1. Hold the leg plate with both hands and press the release button (4).
- 2. Pull the leg plate out slowly without tilting.



3.10.3.3 Adjustment

- To adjust the leg plate up/down:
- 1. Hold the leg plate and press the release bracket (5) upward.
- 2. Raise or lower the leg plate to the desired position.
- 3. Release the leg plate and the release bracket (5).



- To abduct the leg plate:
- 1. Loosen the handle screw (6).
- 2. Abduct the leg plate to the desired position.

3. Tighten the handle screw (6).



3.10.4 Pad

WARNING

- Each time before using the pad, check it carefully. Damaged pad must be replaced immediately.
- Do not use the worn or damaged pad. Liquid can penetrate damaged pad and cause hygienic problems.
- The patient may slip off the operating table if the pad is not secured properly to the surface.

CAUTION

- Ensure the Velcro strap is not damaged or wet before installing the pad. Otherwise, the pad cannot be reliably installed.
- Ensure the pad is affixed flat and reliably. Otherwise, personal damage may occur.
- Do not place sharp objects on the pad. Otherwise, the pad may be damaged.

NOTE

• Pad should be stored lying flat and horizontal.

• When removing the pad, be sure to grasp it with both hands.

3.10.4.1 Installation

- 1. Ensure the pad is fully aligned with the table top.
- 2. Check if the pad is affixed flat and reliably.



3.10.4.2 Removal

When removing the pad, be sure to grasp it with both hands.



3.11 Equipment Preparation

Check the following items before use:

- The operating table is placed on a smooth ground.
- When running on mains power, ensure the equipment is connected to AC power outlet with protective earth.
- When running on battery power, ensure the battery capacity is sufficient.
- The operating table is locked.

WARNING

- Before using the equipment, the user must verify that the equipment, connecting cables and accessories are in correct working order and operating condition.
- Do not use the operating table if it is damaged or cannot work properly. Contact service personnel or Mindray.

4.1 General

The operating table can be used together with different modular table tops and accessories, which makes different positioned patients suitable for different surgeries.

The table top is radiolucent and enables the intraoperative use of x-ray equipment.

WARNING

• Be sure to select the proper patient orientation and accessories according to patient weight and overall load.

NOTE

• Position patients preferably in normal patient orientation on the table top.

4.2 With an Overall Load between 250kg and 460kg

With an overall load of more than 250kg, the operating table should be used with the following restrictions:

- 1. Lock the castors.
- 2. Position the patient to the normal patient orientation.
- 3. Do not operate the longitudinal shift.
- 4. Trendelenburg/reverse Trendelenburg: ≤15°
- 5. Tilt to left/right: \leq 5°
- 6. Back plate up: $\leq 70^{\circ}$
- 7. Back plate down: $\leq 10^{\circ}$

Accessories used together with the table top are shown below:



(1)Upper back plate

(2)Head plate

(3)2-joint head plate

(4)Leg plate

4.3 With an Overall Load between 155kg and 250kg

With an overall load of more than 155kg and no more than 250kg, the operating table should be used with the following restrictions:

1. Lock the castors.

Accessories used together with the table top are shown below:

Normal patient orientation



Reverse patient orientation





(2)4-part leg plate

(3)One-section leg plate

(4)Extension plate

(5)Upper back plate

(6)Head plate

(7)2-joint head plate

4.4 With an Overall Load Less Than 155kg

With an overall load of less than 155kg, the table top should be used with the following accessories when castors are locked:

Normal patient orientation



Reverse patient orientation



5.1 Safety Information

The product must be cleaned and disinfected after every use.

WARNING

- Use only Mindray approved cleaners and disinfectants and methods listed in this chapter to clean or disinfect your equipment. Warranty does not cover damage caused by unapproved substances or methods.
- Do not mix disinfecting solutions as hazardous gases may be generated.
- We make no claims regarding the efficacy of the listed chemicals or methods as a means for controlling infection. For the method to control infection, consult your hospital's infection control officer or epidemiologist.
- Agents containing alcohol can form explosive vapor mixtures and ignite where high-frequency equipment is being used. Do not use the cleaners or disinfectants containing alcohol where high-frequency surgical equipment may be used.
- Always wear gloves for cleaning and disinfection. Neglect of this may cause infection.
- Be sure to disconnect the power supply before cleaning and disinfection.
- Do not put hands into the gaps when cleaning /or disinfecting the table top. Otherwise, you may be pinched.
- The responsible hospital or institution shall carry out all cleaning and disinfection procedures specified in this chapter.

CAUTION

- Do not clean/disinfect the operating table mechanically.
- Never immerse any part of the equipment or accessories in liquids.
- Particles of grime may become encapsulated and lead to the product not reaching the desired germ-reduction after disinfection. Before disinfection, the product must be cleaned thoroughly of contamination and encapsulated particles of grime and then be dried.

- Any contact of cleaners or disinfectants with connectors may cause corrosion.
- Do not pour or spray any liquid directly on the equipment or accessories or permit fluid to seep into live parts inside the equipment.
- If you spill liquid on the equipment or accessories, disconnect the power supply, dry the equipment and contact your service personnel.
- Improper cleaning/disinfection can damage the operating table. Do not spray the cleaning agent/disinfectant directly into the joints or gaps. Do not use high pressure to clean/disinfect components.
- Never use abrasive materials (such as steel wool or silver polish), or strong solvent (such as acetone or acetone-based cleaners) for cleaning.
- Do not use polish or steel cleaner to clean or disinfect stainless steel surfaces.
- Dilute and use the cleaners or disinfectants according to the manufacturer's instructions.

NOTE

- Be sure to follow the related national hygiene and disinfection regulations to carry out cleaning and disinfection procedures.
- Check the equipment or accessories after cleaning and disinfection. Stop the use if there is any sign of wear or damage.

5.2 Cleaning

5.2.1 Preparation before Cleaning

- 1. Adjust the operating table to 0 position.
- 2. Lift the table top to the highest position.
- 3. Switch off the operating table and disconnect the power cord.

5.2.2 Cleaning Procedure

- 1. Use a piece of lint-free clean cloth to wipe off the dust gently and thoroughly.
- 2. Use a piece of lint-free cloth moistened with cleaner (clean water or soap solution) to wipe the surface.
- 3. Use a piece of disposable lint-free cloth moistened with water to clean the surface.
- 4. Use a piece of dry lint-free cloth to wipe the surface dry.

- Use proper amount of the cleaner to clean the operating table, and remove any excessive agent by the dry cloth.
- Be sure to disinfect the equipment after cleaning.

5.3 Disinfection

5.3.1 Recommended Disinfectants

Disinfect the equipment as required in your hospital's service schedule. Cleaning the equipment before disinfection is recommended. Always dilute and use disinfectants according to the manufacturer's instructions. The following table lists approved disinfectants:

Product name	Product type	Manufacturer
Alpet® D2 Surface Sanitizing Wipes	Wipes	BEST SANITIZERS INC™.
CIDEX [®] OPA	Liquid	Gilag GmbH International Advanced Sterilization products
Clorox Dispatch® Hospital Cleaner Disinfectant Towels with Bleach	Wipes	Clorox professional products company
Clorox Healthcare® Bleach Germicidal Wipes	Wipes	Clorox professional products company
Metrex CaviCide1™	Liquid, spray	METERX® RESEARCH
Metrex CaviWipes™	Wipes	METERX® RESEARCH
PDI Sani-Cloth [®] AF3 Germicidal Disposable Wipe	Wipes	PDI Inc.
PDI Sani-Cloth® HB Germicidal Disposable Wipe	Wipes	PDI Inc.
PDI Sani-Cloth [®] Plus Germicidal Disposable Cloth	Wipes	PDI Inc.
PDI Super Sani-Cloth® Germicidal Disposable Wipe	Wipes	PDI Inc.
Virex [®] II 256 (1:256)	Liquid	Diversey Inc

Product name	Product type	Manufacturer
Rely+On™ Virkon [®] High Level surface Disinfectant, 1%	Powder	Antec International Ltd
Clinell [®] Universal Wipes	Wipes	GAMA Healthcare Ltd
Clinell [®] Sporicidal Wipes	Wipes	GAMA Healthcare Ltd
Tristel Duo™	Liquid, foam	Tristel solutions Limited
Tristel Jet	Liquid, spray	Tristel solutions Limited
Surfanios Premium, 0.25%	Liquid	ANIOS LABORATORIES
Surfa 'safe	Liquid, spray	ANIOS LABORATORIES
Aniosurf ND premium, 0.25%	Liquid	ANIOS LABORATORIES
mikrozid® Sensentive Wipes	Wipes	Schülke & Mayr GmbH
mikrozid® AF Wipes	Wipes	Schülke & Mayr GmbH
Terralin [®] Liquid	Liquid	Schülke & Mayr GmbH
JIAN ZHI SU Disinfectant Tablets	Tablet	Beijing ChangJiangMai Medical Science Technology Co. Ltd
JIAN ZHI SU Surface Disinfectant Spray	Liquid, spray	Beijing ChangJiangMai Medical Science Technology Co. Ltd
DIAN'ERKANG Surface Disinfectant	Liquid	Shanghai Likang Disinfectant Hi-Tech Co., Ltd
DIAN'ERKANG Surface Wipes	Wipes	Shanghai Likang Disinfectant Hi-Tech Co., Ltd
DIAN'ERKANG Disinfectant Spray	Liquid, spray	Shanghai Likang Disinfectant Hi-Tech Co., Ltd
Glutaraldehyde, 2%	Liquid	/
Ethanol, 70%	Liquid	/
Isopropanol, 70%	Liquid	/

Product name	Product type	Manufacturer
Hydrogen peroxide, 3%	Liquid	/
1-Propanol, 50%	Liquid	/
Domiphen, 2000mg/L	Liquid	/

5.3.2 Disinfection Procedure

- 1. Use a piece of lint-free cloth moistened with disinfectant to wipe the surface.
- 2. Use a piece of disposable lint-free cloth moistened with water to clean the surface.
- 3. Use a piece of dry lint-free cloth to wipe the surface dry.

CAUTION

- Do not disinfect the equipment by fumigation methods. Otherwise, the equipment may be damaged.
- Do not use the disinfectants containing chlorine or compounds which can release chlorine to disinfect the metal parts. Long-term use of these disinfectants may corrode the metal parts, hence compromising safety and effectiveness of the device. It is recommended that metal parts be disinfected by 2% alkaline glutaraldehyde.

5.4 Pad Cleaning and Disinfection

WARNING

• Do not use the worn or damaged pad. Liquid can penetrate damaged pad and cause hygienic problems.

CAUTION

- Be sure to replace damaged pad immediately.
- Moisture may cause damage to pad.
- Do not use the disinfectant containing chlorine or compounds which can release chlorine.
- The adhesive force of the Velcro strap may be compromised due to contamination. It is recommended a plastic soft brush or a plastic comb be used to remove the contamination.

5.4.1 Cleaning Procedure

- 1. Remove the pad.
- 2. Use a piece of lint-free cloth moistened with cleaner (clean water or soap solution) to wipe the pad and then dry the surface.

5.4.2 Disinfection Procedure

- 1. Use a piece of lint-free cloth moistened with disinfectant (as described in **5.3.1 Recommended Disinfectants**) to wipe the surface.
- 2. Use a piece of disposable lint-free cloth moistened with water to clean the surface.
- 3. Use a piece of dry lint-free cloth to wipe the surface dry.

NOTE

- When removing the pad, be sure to grasp it with both hands.
- Pad should be stored lying flat and horizontal.
- Before installing the pad, ensure that the table top, pad and Velcro Strap are dry.

Regular maintenance is essential to ensure that the equipment functions properly. This chapter contains information on equipment maintenance method and frequency.

6.1 Maintenance Safety Information

WARNING

- To avoid electric shock, stop using the equipment if you find the housing has signs of damage. Contact the service personnel for help in that case.
- Failure on the part of the responsible individual hospital or institution using this equipment to implement a recommended maintenance schedule may cause undue equipment failure and possible health hazards.
- If you have any questions during the inspection, contact the customer service department authorized by Mindray.
- The device maintenance that requires disassembling shall be performed by the professional service personnel authorized by Mindray. Otherwise, device failure or personal safety accidents may occur.
- The service personnel must be properly qualified and thoroughly familiar with the operation of the equipment.

CAUTION

- The equipment and accessories shall not be tested or maintained while in use with a patient.
- If you discover a problem with any of the equipment, contact your service personnel or Mindray.

NOTE

- This equipment contains no user serviceable parts. Refer servicing to qualified service personnel.
- If needed, contact Mindray for circuit diagrams, component part lists, descriptions, calibration instructions, or other information concerning the repair of the equipment.

6.2 Maintenance Schedule

Follow the maintenance schedule or local regulations to perform maintenance. Be sure to clean and disinfect the equipment before test or maintenance. The following table lists the maintenance and testing schedule:

Test/Maintenance Item		Recommended Frequency
Visual inspection		Every day, before first use.
Switch-on test		Each time the operating table is switched on.
Battery maintenance	Functional test	1. When first installed. 2. When the battery is replaced.
	Performance test	Every three months or when the battery runtime reduced significantly
Hydraulic oil repl	acement	Every five years recommended

6.3 Testing Methods and Procedures

Except the following maintenance tasks, all other test and maintenance tasks should be performed by Mindray's qualified service personnel only.

- Regular check, including visual inspection and switch-on test
- Battery maintenance

If the operating table needs other test and maintenance, contact the service personnel.

6.3.1 Visual Inspection

Visually inspect the equipment before its first use every day. If you find any signs of equipment or accessory damage, stop using it and contact the service personnel.

Visual inspection items are as follows:

- Environment and power supply specifications are met.
- The equipment surface has not any stain. The override panel and hand control are free from cracks or damages.
- Mechanical parts are not damaged.
- Connectors, plugs and cables are not damaged and kinked.
- Cables are securely connected with the equipment.
- Pads are not damaged and firmly fixed on the table top.
- The operating table has no oil leakage.
- The operating table has no obvious vibration.
6.3.2 Switch-on Test

Check the following items after the operating table is switched on:

- The equipment can be switched on properly.
- The operating table works properly: table up/down, tilt, slope, longitudinal shift, back plate up/down, lock/unlock
- The hand control displays properly.

6.3.3 Battery Maintenance

6.3.3.1 Battery Safety Information

WARNING

- Use specified batteries only. Use of different batteries may present a risk of fire or explosion.
- Do not crush, drop or puncture the battery. Mechanical abuse may lead to internal damage and internal short circuits. If a battery has been dropped or banged against a hard surface, whether damage is externally visible or not, remove the battery from use and dispose of it properly.
- If the battery shows signs of damage or leakage, replace it immediately.
- The battery must only be installed and replaced by service personnel trained and authorized by Mindray. Installation or replacement by untrained personnel may cause personal injury (e.g. batteries overheated, ignited or exploded).
- Extremely high ambient temperature may cause battery overheat protection, resulting in interruption of power supply.
- Do not open the battery, heat the battery above 60 °C, incinerate battery, or short battery terminals. They may ignite, explode, leak or heat up, causing personal injury.

CAUTION

- Remove the battery if it will not be used for an extended period of time. Contact the customer service personnel authorized by Mindray or local distributor.
- Storing the battery at high temperature for a long period of time will significantly shorten its life expectancy.

 Battery runtime depends on equipment configuration and operation. For example, high brightness of the screen or repeated operation will shorten the battery runtime.

6.3.3.2 Functional Test

The built-in rechargeable lead-acid battery ensures normal use of the operating table without external power supply. Once the external power supply is connected, the battery will be charged automatically. See **3.2.2** Charging Batteries for details.

The built-in rechargeable lithium-ion battery ensures normal use of the wireless hand control without external power supply. Use the mobile charging station to charge the wireless hand control. See **3.3.6.2 Charging Wireless Hand Control** for details.

6.3.3.3 Optimizing Batteries

Life expectancy of a battery depends on how frequent and how long it is used. The performance of a rechargeable battery deteriorates over time.

It is recommended the battery be optimized every three months. Use the battery with its remaining capacity indicated the last bar before optimization. Optimize the battery as shown below:

- Operating table battery
- 1. Switch off the operating table and connect it to AC power.
- 2. Charge the battery uninterruptedly until it is fully charged. It takes about 10 hours to fully charge the battery.
- 3. Then disconnect it from AC power.

NOTE

- If the battery is not maintained for a prolonged time, its capacity indication may not be accurate and you may wrongly evaluate the remaining battery runtime. In addition, the battery is subject to regular maintenance. Otherwise, aging may be accelerated and unexpected failure may occur.
- Do not use the operating table during battery optimization.
- Do not interrupt the battery optimization.
- Battery of wireless hand control

It is recommended the battery be optimized every three months if the wireless hand control has not been used for a long period. Optimize the battery as shown below:

- 1. Connect the mobile charging station to AC power.
- 2. Switch off the wireless hand control and insert it into the mobile charging station and charge the wireless hand control uninterruptedly until it is fully charged.

- Disconnect the wireless hand control from the external AC power and switch it on. Use the wireless hand control to adjust the operating table until the battery of the wireless hand control is completely discharged and the wireless hand control automatically shuts down.
- 4. Recharge the wireless hand control fully if it is to be used. If the wireless hand control is to be stored, place the battery with 40% to 60% charging capacity.

6.3.3.4 Checking Battery Performance

See steps in **6.3.3.3 Optimizing Batteries** to check the battery performance. The runtime of the battery reflects their performance directly. If the runtime of a battery is noticeably shorter than that stated in the specifications, the battery may reach its service life or malfunction. Contact your service personnel.

6.4 Disposal

Dispose of the equipment when its service life is reached.

WARNING

- At the end of its service life, the equipment, as well as its accessories, must be disposed of in compliance with the local regulations. If you have any questions concerning disposal of the equipment, please contact Mindray.
- For disposal of parts and accessories, unless otherwise specified, follow local regulations regarding disposal of hospital waste.
- Dispose of the package material as per the applicable waste control regulations. Keep it out of children's reach.

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7.1 Common Error

WARNING

- The chapter is meant to help you solve common errors only. In case you
 encounter problems not included in this chapter or following the introduced
 methods cannot solve the problem, contact the customer service department
 authorized by Mindray or your local distributor for help. Unauthorized device
 servicing is not allowed.
- Device servicing is to be performed by personnel authorized by Mindray only. Unauthorized device servicing may cause personal injury and/or property damage.
- Be sure to disconnect the equipment from AC power when device servicing.
- Device servicing should be strictly based on technical data supplied by Mindray. If you are in need of more technical data, contact the customer service department authorized by Mindray or your local distributor.

Error	Cause	Solution	
Bluetooth connection of the wireless hand control fails.	1. Fail to start the connection within 20 seconds.	1. Press and hold the Trendelenburg button on the override panel and start the bluetooth connection within 20 seconds.	
	2. The operating table is powered off.	2. Press the power button on the base to power on the operating table.	
	3. The wireless hand control stays far away from the operating table or there are obstacles between the wireless hand control and operating table.	3. The distance between the wireless hand control and the operating table is no more than 3.5 meters, and there are not any obstacles between the wireless hand control and operating table.	
	4. The operating table has been connected with another wireless hand control.	4. Reconnect by following steps of the bluetooth connection.	
	5. Bluetooth module fails.	5. Contact the customer service department authorized by Mindray or your local distributor.	
Unlock fails.	The unlock button has not been pressed for 2 seconds.	Press and hold the unlock button for two seconds. See 3.3.5.2 Unlock / Lock .	
	The operating table fails.	Perform the manual brake release. See 3.8 Manual Brake Release .	
Locking indicator (green) and unlocking indicator (red) on the override panel flashes at the same time.	The operating table is in the intermediate state of locking and unlocking.	Perform locking or unlocking again.	
Battery indicator on the base is red and flashes.	The second battery bar is red and flashes.	Contact the customer service department authorized by Mindray	
	The third battery bar is red and flashes.	or your local distributor.	
	The fourth battery bar is red and flashes.		

Error	Cause	Solution
The icon appears on the screen.	The operating table may have any hardware, sensor or motor errors.	Contact the customer service department authorized by Mindray or your local distributor.

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A.1 Safety Specifications

A.1.1 Product Classification

According to the protection against electrical shock	Class I, internal electrical power source. Type B applied part.
According to the protection against harmful ingress of water or particulate matter	Operating table: IPX5 Mobile charging station (except the adapter): IPX5 Foot switch: IPX8
According to the method(s) of sterilization	Non-sterilization.
According to the suitability for use in an oxygen rich environment	The operating table cannot be used in an oxygen rich environment.
According to the mode of operation	Non-continuous operation: 10 minutes on /20 minutes off

WARNING

 To avoid the risk of electric shock, this equipment must only be connected to a supply mains with protective earth. Before installing it, check and ensure that the protective earth system in the operating room is reliable and safe.

A.1.2 Environmental Specifications

ltem	Temperature (°C)	Humidity (non- condensing)	Atmospheric pressure (kPa)
Operating condition	5-40	15%-95%	70-106
Storage condition	-40-60	10%-95%	50-106

CAUTION

- During transportation, ensure that the operating table is well protected from rain, snow or mechanical collision.
- The operating table should be stored in a room that is dry, draughty and without caustic gas.

A.2 Power Supply Specifications

A.2.1 AC Power

A.2.2 Fuse

Fuse	250V T8AH

A.2.3 Operating Table Batteries

Туре	Internal batteries
Voltage	12V×2
Runtime	About one week
Charge time	About 10h
Power-off delay	About 10 minutes of continuous operation after the low battery prompt first occurs

A.3 Wireless Specifications

WARNING

- Do not connect non-medical devices to the operating table network.
- If wireless network signal is poor, there may be a risk of data loss.
- RF interference may result in wireless network disconnection.

A.3.1 Bluetooth

Work frequency	2402-2480MHz	
Modulation mode	GFSK	
Transmission power	<20dBm	

A.3.2 Mobile Charging Station

Work frequency	112-205KHz
Modulation mode	AM
Transmission power	<20dBm

A.4 Load

- 1. Maximum overall load: 460kg
- 2. With an overall load of more than 250kg, the operating table should be used with the following restrictions:
 - a Lock the castors.
 - b Position the patient to the normal patient orientation.
 - c Do not operate the longitudinal shift.
 - d Trendelenburg/reverse Trendelenburg: $\leq 15^{\circ}$
 - e Tilt to left/right: $\leq 5^{\circ}$
 - f Back plate up: $\leq 70^{\circ}$
 - g Back plate down: $\leq 10^{\circ}$
- 3. With an overall load of more than 155kg and no more than 250kg, the operating table should be used with the following restrictions:
 - a Lock the castors.

A.5 Performance Specifications

1. Base thickness: 140mm±10mm



- 2. Radiolucent distance
 - Maximum radiolucent distance with longitudinal shift toward head end: 1007mm±10mm



 Maximum radiolucent distance with longitudinal shift toward foot end: 1013mm±10mm



3. Maximum Trendelenburg/reverse Trendelenburg: 36°±3°



4. Maximum tilt angle: 26°±2°



5. Head plate: head plate up: $45^{\circ}\pm5^{\circ}$, head plate down: $90^{\circ}\pm5^{\circ}$, detachable



6. Back plate: back plate up: $80^{\circ}\pm5^{\circ}$, back plate down: $40^{\circ}\pm5^{\circ}$



7. Leg plate: leg plate up: $20^{\circ}\pm5^{\circ}$, leg plate down: $90^{\circ}\pm5^{\circ}$, detachable



- 8. Maximum table top lifting height: 450mm±20mm
 - Maximum table top height: 1050mm±20mm
 - Minimum table top height: 600mm±50mm



- 9. Maximum longitudinal shift: 350mm±10mm
 - Longitudinal shift toward head end: 160mm±10mm
 - Longitudinal shift toward foot end: 190mm±10mm



10. Flex position: Reverse Trendelenburg: 20±3°, back plate down: 40±3°



11. Reflex position: Trendelenburg: 30±3°, back plate up: 70±3°



12. Body elevator lifting distance: 120mm±10mm Body elevator width: 86mm±10mm



A.6 Accessories and Parts List

ltem	Name
1.	Shoulder support
2.	Body support
3.	Suspended arm strap
4.	Leg holder with loops
5.	Arm board
6.	Leg support
7.	Leg holder
8.	Wristlet
9.	Leg restraint cuff
10.	Body strap
11.	Head rest system
12.	Head plate
13.	Table extender
14.	Leg plate
15.	Rectal positioning device
16.	Arm extension device
17.	Foot plate
18.	Leg holder
19.	Stirrup
20.	Meniscus positioning device
21.	Back plate for shoulder operation
22.	Bow frame for spine operation
23.	Orthopaedic extension device
24.	Position pad
25.	Hand grips
26.	Drainage bowl
27.	Side rail extension

ltem	Name
28.	Instrument tray
29.	Cassette fixer
30.	Infusion holder
31.	Accessory cart
32.	Anesthesia frame
33.	Clamp
34.	Foot switch
35.	Remote control module

A.7 Electronic Interface

You may see the following electronic interfaces on the operating table. The technical descriptions of the electronic interfaces are as follows:

Electronic Interface	Specification
	Used for transmitting control commands and operating table status
Corded hand control interface	Connected with the corded hand control
	Mindray internal protocol
	Push-pull self-locking connector
	Used for transmitting control commands and operating table status
Foot switch interface	Connected with the foot switch
	Mindray internal protocol
	Push-pull self-locking connector

The device meets the requirements of IEC 60601-1-2. The accessories listed in this manual meet the requirements of IEC 60601-1-2 when used with the device.

WARNING

- Use of accessories, transducers 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.
- Use of this device adjacent to or stacked with other device should be avoided because it could result in improper operation. If such use is necessary, this device and the other device should be observed to verify that they are operating normally.
- 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 HyBase V8 Classic operating table, including cables specified by the manufacturer. Otherwise, degradation of the performance of this device could result.
- The non-ME EQUIPMENT (e.g. ITE) that is a part of an ME SYSTEM may be disrupted by the electromagnetic interference of nearby equipment. It may be necessary to take mitigation measures, such as re-orienting or relocating the non-ME EQUIPMENT or shielding the location.
- This device is intended for use in professional healthcare facility environment only. If it is used in special environment, such as magnetic resonance imaging environment, the equipment/system may be disrupted by the operation of nearby equipment.

Guidance and Declaration - Electromagnetic Emissions

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

Emission test	Compliance	Electromagnetic environment - guidance
Conducted and radiated RF EMISSIONS CISPR 11	Group 1	The device uses RF energy only for its internal function. Therefore, its RF emissions are very low and are not likely to cause any interference in nearby electronic device.
Conducted and radiated RF EMISSIONS CISPR 11	Class A	The device is suitable for use in all establishments other than domestic and those directly connected to the public
Harmonic distortion IEC 61000-3-2	Not applicable	supplies buildings used for domestic purposes.
Voltage fluctuations and flicker IEC 61000-3-3	Not applicable	

NOTE

- The device needs special precautions regarding EMC and needs to be installed and put into service according to the EMC information provided below.
- Other devices may affect this device even though they meet the requirements of CISPR.
- The EMISSIONS characteristics of this device make it suitable for use in industrial areas and hospitals (CISPR 11 class A). If it is used in a residential environment (for which CISPR 11 class B is normally required) this device might not offer adequate protection to radio-frequency communication services. The user might need to take mitigation measures, such as relocating or re-orienting the device.
- If the essential performance is lost or degraded, it may be necessary to take mitigation measures, such as re-orienting or relocating the ME EQUIPMENT or ME SYSTEM or shielding the location or stopping using the device and contact the service personnel.

If the device is operated within the electromagnetic environment listed in Table Guidance and Declaration —Electromagnetic Immunity, the system will remain safe and provide the following essential performance: the normal operating table movement realized by electric control of the control unit; supporting a patient without unwanted movement in a single fault condition.

The device is int customer or the	ended for use in the ele user of the device show	ectromagnetic environ uld assure that it is used	ment specified below. The d in such an environment.
lmmunity test	IEC 60601 test level	Compliance level	Electromagnetic environment - guidance
Electrostatic discharge (ESD) IEC 61000-4-2	±8 kV contact ±15kV air	±8 kV contact ±15kV air	Floors should be wood, concrete or ceramic tile. If floors are covered with synthetic material, the relative humidity should be at least 30%.
Electrical fast transient/ burst IEC 61000-4-4	±2 kV for power supply lines ±1 kV for input/ output lines (length greater than 3 m)	±2 kV for power supply lines ±1 kV for input/ output lines (length greater than 3 m)	Mains power quality should be that of a typical commercial or hospital environment.
Surge IEC 61000-4-5	± 1 kV line(s) to line(s) ± 2 kV line(s) to earth	± 1 kV line(s) to line(s) ± 2 kV line(s) to earth	
Voltage dips and voltage interruptions IEC 61000-4- 11	$0 \% U_T$ for 0,5 cycle $0 \% U_T$ for 1 cycle and 70 % U_T for 25/ 30 cycles $0 \% U_T$ for 250/300 cycle	$0 \% U_{T}$ for 0,5 cycle $0 \% U_{T}$ for 1 cycle and 70 % U _T for 25/ 30 cycles $0 \% U_{T}$ for 250/300 cycle	Mains power quality should be that of a typical commercial or hospital environment. If the user of our product requires continued operation during power mains interruptions, it is recommended that our product be powered from an uninterruptible power supply or a battery.
RATED power frequency magnetic fields IEC 61000-4-8	30 A/m 50 Hz / 60 Hz	30 A/m 50 Hz / 60 Hz	Power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercial or hospital environment.

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Guidance and Declaration - Electromagnetic Immunity

The device is intended for use in the specified electromagnetic environment. The customer or the user of the device should assure that it is used in such an environment as described below.

lmmunity test	IEC 60601 Test level	Compliance level	Electromagnetic environment - guidance
Conducted disturbances induced by	3 Vrms 150 kHz to 80 MHz	3 Vrms	Portable and mobile RF communications equipment should be used no closer to any part of the device, including cables,
IEC61000-4-6	6 Vrms in ISM bands ^a between 0,15 MHz and 80 MHz	6 Vrms	than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter. Recommended separation distance: $d = \left[\frac{3.5}{V}\right]\sqrt{P}$ 150kHz to 80 MHz
Radiated RF EM fields IEC61000-4-3	3 V/m 80 MHz to 2.7 GHz	3V/m	[0.c]
Proximity fields from RF wireless	27 V/m 380–390 MHz	27 V/m	$d = \left\lfloor \frac{3.3}{E} \right\rfloor \sqrt{P}$ 80 MHz to 800 MHz
communicati ons equipment IEC61000-4-3	28 V/m 430-470 MHz, 800- 960 MHz, 1700-1990 MHz, 2400- 2570 MHz	28 V/m	$d = \left\lfloor \frac{r}{E} \right\rfloor \sqrt{P}$ booming to 2.7 GHz Where P is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer and d is the
	9 V/m 704–787 MHz, 5100– 5800 MHz	9 V/m	recommended separation distance in meters (m). Field strengths from fixed RF transmitters, as determined by an
	5000 111 12		electromagnetic site survey ^b , should be less than the compliance level in each frequency range ^c . Interference may occur in the vicinity of equipment
			marked with the following symbol: $(())$
Note 1: At 80 Mi Note 2: These gu	Hz and 800 MHz, uidelines may no	the higher frequ t apply in all situ	iency range applies. iations. Electromagnetic propagation is

affected by absorption and reflection from structures, objects and people.

^a The ISM (industrial, scientific, and medical) bands between 150 kHz and 80 MHz are 6.765 MHz to 6.795 MHz; 13.553 MHz to 13.567 MHz; 26.957 MHz to 27.283 MHz; and 40.66 MHz to 40.70 MHz.

^b 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 device is used exceeds the applicable RF compliance level above, the device should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as re-orienting or relocating the device.

^c Over the frequency ranges 150 kHz to 80 MHz, field strengths should be less than 3V/m.

Recommended Separation Distances between Portable and Mobile RF, Communications Equipment and This Equipment

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

Rated Maximum Output power of	Separation Distance	According to Frequenc	y of Transmitter (m)
Transmitter Watts (W)	150 kHz to 80 MHz $d = \left[\frac{3.5}{V}\right]\sqrt{P}$	80 MHz to 800 MHz $\vec{a} = \left[\frac{3.5}{E}\right]\sqrt{P}$	800 MHz to 2.7 GHz $d = \left[\frac{7}{E}\right] \sqrt{P}$
0.01	0.12	0.12	0.23
0.1	0.38	0.38	0.73
1	1.2	1.2	2.3
10	3.8	3.8	7.3
100	12	12	23

For transmitters at a maximum output power not listed above, the recommended separation distanced 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. Note 1: At 80 MHz and 800 MHz, the higher frequency range applies.

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

Cable:

No.	Description	Length (m)	Shielded or not
1	AC power cord	≤5	Not shielded

C Declaration of Conformity

D	eclaration of Conformity
Manufacturer:	Nanjing Mindray Bio-Medical Electronics Co., Ltd. 666# Middle Zhengfang Road, Jiangning, 211111 Nanjing, Jiangsu, P.R.China
EC-Representative:	Shanghai International Holding Corp. GmbH (Europe) Eiffestrasse 80, 20537 Hamburg, Germany (DIMDI No.: DE/0000040627)
Product Name :	Operating Table
Model:	HyBase V8, HyBase V80, HyBase V8 Classic, HyBase V80 Classic, HyBase V6, HyBase V60, HyBase V60C, HyBase S6, HyBase S60, HyBase S7, HyBase S70, HyBase S8, HyBase S80
Remark:	1
Object of the declaration: W above satisfies the provisions supporting documentation is The product complies with fi	e declare on our sole responsibility that the product mentioned of Directive 2014/53/EU concerning radio equipment. All retained under the premises of the manufacturer. Jlowine standards and/or other normative documents:
Object of the declaration: W above satisfies the provisions supporting documentation is The product complies with fe	e declare on our sole responsibility that the product mentioned of Directive 2014/53/EU concerning radio equipment. All retained under the premises of the manufacturer. ollowing standards and/or other normative documents:
Object of the declaration: W above satisfies the provisions supporting documentation is The product complies with fo Standards Applied:	e declare on our sole responsibility that the product mentioned of Directive 2014/53/EU concerning radio equipment. All retained under the premises of the manufacturer. Illowing standards and/or other normative documents: EN 60601-1:2006+A1:2013; EN 60601-1:2:00+2:015;
Object of the declaration: W above satisfies the provisions supporting documentation is The product complies with fo Standards Applied:	e declare on our sole responsibility that the product mentioned of Directive 2014/53/EU concerning radio equipment. All retained under the premises of the manufacturer. ollowing standards and/or other normative documents: EN 60601-1:2006+A1:2013; EN 60601-1-2:2015; EN 63011-2008:
Object of the declaration: W above satisfies the provisions supporting documentation is The product complies with fo Standards Applied:	e declare on our sole responsibility that the product mentioned of Directive 2014/53/EU concerning radio equipment. All retained under the premises of the manufacturer. allowing standards and/or other normative documents: EN 60601-1:2006+A1:2013; EN 60601-1-2:2015; EN 62311:2008; ETSI EN 801 489-1 V2.2.3;
Object of the declaration: W above satisfies the provisions supporting documentation is The product complies with fo Standards Applied:	e declare on our sole responsibility that the product mentioned of Directive 2014/53/EU concerning radio equipment. All retained under the premises of the manufacturer. ollowing standards and/or other normative documents: EN 60601-1:2006+A1:2013; EN 60601-1-2:2015; EN 62311:2008; ETSI EN 301 489-1 V2.2.3; ETSI EN 301 489-3 V2.1.1;
Object of the declaration: W above satisfies the provisions supporting documentation is The product complies with fo Standards Applied:	e declare on our sole responsibility that the product mentioned of Directive 2014/53/EU concerning radio equipment. All retained under the premises of the manufacturer. Illowing standards and/or other normative documents: EN 60601-1:2006+A1:2013; EN 60601-1-2:2015; EN 63311:2008; ETSI EN 301 489-1 V2.2.3; ETSI EN 301 489-3 V2.1.1; EN 303 417 V1.1.1;
Object of the declaration: W above satisfies the provisions supporting documentation is The product complies with fo Standards Applied:	e declare on our sole responsibility that the product mentioned of Directive 2014/53/EU concerning radio equipment. All retained under the premises of the manufacturer. ellowing standards and/or other normative documents: EN 60601-1:2006+A1:2013; EN 60601-1-2:2015; EN 63311:2008; ETSI EN 301 489-1 V2.2.3; ETSI EN 301 489-3 V2.1.1; EN 303 417 V1.1.1; ESTI EN 300 328 V2.2;
Object of the declaration: W above satisfies the provisions supporting documentation is The product complies with fo Standards Applied:	e declare on our sole responsibility that the product mentioned of Directive 2014/53/EU concerning radio equipment. All retained under the premises of the manufacturer. ollowing standards and/or other normative documents: EN 60601-1:2006+A1:2013; EN 60601-1-2:2015; EN 62311:2008; ETSI EN 301 489-1 V2.2.3; ETSI EN 301 489-3 V2.1.1; EN 103 417 V1.1.1; ESTI EN 303 228 V2.2.2; ETSI EN 301 489-17 V3.1.1;
Object of the declaration: W above satisfies the provisions supporting documentation is The product complies with fo Standards Applied:	e declare on our sole responsibility that the product mentioned of Directive 2014/53/EU concerning radio equipment. All retained under the premises of the manufacturer. ollowing standards and/or other normative documents: EN 60601-1:2006+A1:2013; EN 60601-1-2:2015; EN 60301-12:2015; ETSI EN 301 489-1 V2.2.3; ETSI EN 301 489-1 V2.2.3; ETSI EN 301 489-3 V2.1.1; EN 103 417 V1.1.1; ESTI EN 300 328 V2.2.2; ETSI EN 301 489-17 V3.1.1; EN 300 330 V2.1.1
Object of the declaration: W above satisfies the provisions supporting documentation is The product complies with fo Standards Applied: Place, Date of Issue:	e declare on our sole responsibility that the product mentioned of Directive 2014/53/EU concerning radio equipment. All retained under the premises of the manufacturer. ellowing standards and/or other normative documents: EN 60601-1:2006+A1:2013; EN 60601-1-2:2015; EN 6231:12008; ETSI EN 301 489-1 V2.2.3; ETSI EN 301 489-3 V2.1.1; EN 303 417 V1.1.1; ESTI EN 300 328 V2.2.2; ETSI EN 301 489-17 V3.1.1; EN 300 330 V2.1.1 Nanjing, July 7, 2021
Object of the declaration: W above satisfies the provisions supporting documentation is The product complies with fo Standards Applied: Place, Date of Issue: Signature:	e declare on our sole responsibility that the product mentioned of Directive 2014/53/EU concerning radio equipment. All retained under the premises of the manufacturer. ollowing standards and/or other normative documents: EN 60601-1:2006+A1:2013; EN 60601-1-2:2015; EN 63311:2008; ETSI EN 301 489-1 V2.2.3; ETSI EN 301 489-1 V2.2.3; ETSI EN 301 489-3 V2.1.1; EN 103 437 V1.1.1; ESTI EN 300 328 V2.2.2; ETSI EN 301 489-17 V3.1.1; EN 300 330 V2.1.1 Nanjing, July 7, 2021 Mark Park
Object of the declaration: W above satisfies the provisions supporting documentation is The product complies with fo Standards Applied: Place, Date of Issue: Signature: Name of Authorized Signato	e declare on our sole responsibility that the product mentioned of Directive 2014/53/EU concerning radio equipment. All retained under the premises of the manufacturer. Illowing standards and/or other normative documents: EN 60601-1:2006+A1:2013; EN 60601-1-2:2015; EN 6301-2:2015; EN 5301 489-1 V2.2.3; ETSI EN 301 489-1 V2.2.3; ETSI EN 301 489-3 V2.1.1; EN 303 417 V1.1.1; ESTI EN 300 328 V2.2.2; ETSI EN 301 489-17 V3.1.1; EN 300 310 V2.1.1 Nanjing, July 7, 2021 Mark Led
Object of the declaration: W above satisfies the provisions supporting documentation is The product complies with for Standards Applied: Place, Date of Issue: Signature: Name of Authorized Signato Position Held in Company:	e declare on our sole responsibility that the product mentioned of Directive 2014/53/EU concerning radio equipment. All retained under the premises of the manufacturer. ollowing standards and/or other normative documents: EN 60601-1:2006+A1:2013; EN 60601-1:2015; EN 62311:2008; ETSI EN 301489-1 V2.2.3; ETSI EN 301489-3 V2.1.1; EN 10489-3 V2.1.1; ESTI EN 300 328 V2.2.2; ETSI EN 301 489-17 V3.1.1; EN 300 330 V2.1.1 Nanjing, July 7, 2021 Multi Pei Manager, Technical Regulation

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Accessories for operating tables



www.mindray.com

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HyBase V8 Electro-Hydraulic Operating Table

With increasingly complex surgical procedures, today's surgical environment requires a table that instantly adapts to all versatility to provide the highest level of care. HyBase V8 comes with an extreme weight capacity of up to 460 kg, delivering the highest safety and stability in any position.



4 swivel castor





Intelligent Collision Protection System

HyBase V8 Classic Electro-Hydraulic Operating Table

HyBase V8 Classic, the sequel of HyBase V family, is a new competitive model of Mindray operating family. It not only inherits the highlights of HyBase V family, but also conveys new function which is able to upgrade the safety in OR. As surgical procedures become increasingly complex, the expanding HyBase V series family with different models has been specifically designed to address various customer needs.









Built-in Body Elevato
The Mindray table accessory portfolio provides optimal patient positioning and ideal working conditions for the medical staff. With robust design, ease-to-use and ergonomic functionality, Mindray offers standard table accessories that meets the requirements of nearly every surgical discipline.



0092-30-102657 A33XX Light quadrate clamp

Accessory Name: Universal clamp (Light, one-piece, quadrate)

Application:



Accessory Name: Universal clamp (Heavy, one-piece, quadrate)

Application:



Accessory Name: Universal clamp (Light, one-piece, radial)

Application:



General Accessories

To attach to the side rail for connecting accessories with the table

Compatible with:



0092-30-102655 A32XX Heavy quadrate clamp

To attach to the side rail for connecting accessories with the table



0092-30-102658 A34XX Light radial clamp

To attach to the side rail for connecting accessories with the table





0092-30-102656 A35XX Heavy radial clamp

Accessory Name: Universal clamp (Heavy, one-piece, radial)

Application: Attach to the side rail for connecting accessories with the table





Accessory Name:

Application:



115-038243-00 A09XX Strap clamps, a pair

Accessory Name: Strap clamps, a pair

Application: To fix the body strap in the intended position on the side rail





Accessory Name: "U" shape anesthesia frame with clamps

Application:



0092-30-102643 A60XX Infusion holder

Accessory Name: Infusion pole with one piece with radial universal clamp

Application: To hang the infusion bag in the procedure





Accessory Name:

Anesthesia frame, "L" shape design with two adjustable tubes, one piece with radial universal clamp

Application:

To hang the anesthsia screen during the surgery

03

General Accessories

0092-30-102645 A12XX Light anesthesia frame

Anesthesia Frame, one piece with quadrate universal clamp

To hang the anesthsia screen during the surgery

Compatible with:



115-005689-00 A10XX U-Shaped anesthesia Compatible with: frame V8 / V8 Classic 6100 To hang the anesthsia screen during the surgery

0092-30-102654 A11XX Heavy anesthesia frame

Compatible with: V8 / V8_{Classic} **8500** HyBase **8300** HyBase **6100** HyBase **30** UniBase



115-083097-00 A42XX Light arm board

Accessory Name:

Light arm board with rotation function in horizontal position (with integrated clamp, pad & two pieces of fasten belt for each arm board), a pair

Application:

To support and fasten the arm of the patient to avoid unexpected moving





Accessory Name:

Double layer arm support with one piece of heavy radial clamp for lateral position, with the pad

Application:

To support and fasten both of the arms of the patient as a lateral position to avoid unexpected movement



115-083163-00 A37XX Arm posturing device

Accessory Name:

Arm posturing device, a pair Note: this accessory includes side rail extension(HySRE-2)

Application:

During surgeries, it is used to allow support for patient's arm to effectively protect patient's brachial nerves



115-083165-00 A43XX Heavy arm board, SFC 115-083164-00 A43XP Heavy arm board, PU

Accessory Name:

Heavy arm board with up & down and rotation function (a heavy radial universal clamp, pad & two pieces of fasten belt for each arm board), a pair(SFC/PU)

Application:

To support and fasten the arm of the patient to avoid unexpected moving



Compatible with:



05

115-030153-00 A38XX Carbon fiber arm board, a pair

Accessory Name:

Carbon fiber arm board, with special foam core pad, a pair

Application:

A carbon fibre arm board which includes a memorised foam pad. This can work with the 1180 cardiac operating plate which provides support for the patient's upper extremities in a supine position







General Accessories

115-005693-00 A41XX Double arm board

Compatible with:







115-007051-00 A18XX Suspended arm strap

Accessory Name: Suspended arm strap, mounted on the anesthesia frame

Application: Attach to frame for hanging the hand of the patient





115-083162-00 A19XX Wristlet

Accessory Name: Wristlet with a light guadrate clamp, a pair

Application: Attach to the side rail rail for fastening the hand of the patient







Accessory Name:

Application:

movement

Accessory Name:

clamps and the pad

Application:

movement



0092-30-102640 A13XX Body strap

Accessory Name: Body strap, one-piece

Application:

To fasten the body of the patient to the table during the surgery





Accessory Name:

Heavy shoulder support (with heavy quadrate universal clamp and pad), a pair

Application:

movement

General Accessories







To support the shoulder of the patient for avoiding unexpected





0092-30-102649 A14XX Light shoulder support, a pair

Accessory Name:

Shoulder support (left and right, pair with quadrate universal clamps and pad)

Application:

To support the shoulder of the patient for avoiding unexpected movement



Compatible with:



Accessory Name:

Bow frame for spine operation Foam head block stand of bow frame Pad set for bow frame, a pair

Application:

Bow frame is adjustable for length and width according to different patients Foam head block support the head in prone position Pad set give a comfortable support for the chest



0092-30-102641 A53XX Cassette fixer

Accessory Name:

X-ray Cassette fixer with a light radial clamp (compatible with multi-size)

Application: To fix the X-ray cassette under the table





Accessory Name: Bow frame cart, four universal castors

Application:



0092-30-102639 P02XX Pad for intervertebral disc Ops

Accessory Name:

Intervertebral disc operation pad (Memorise foam, one piece)

Application:

To support the body in intervertebral operating position





Accessory Name: Disposable covers for bow frame(6 pcs)

Application:

General Accessories

115-052219-00 HyBF-1 Bow frame for spine operation 115-053656-00 A76XX Foam head block stand of bow frame 115-053654-00 A75XX Pad set for bow frame, a pair

Compatible with:



Compatible with: 115-052221-00 A73XX Bow frame cart V8 / V8 cta 6100 HyBase It is convenient for the bow frame to store and move 30 UniBase

115-052223-00 A74XX Bow frame disposable Compatible with: cover(6 sets) V8 / V8 Clas **8500** HyBase 6100 HyBase To prevent the pad getting dirty during spine operation **30** UniBase

Leg Accessories



115-035792-00 811XX One-section leg plate 115-037739-00 811CX One-section leg plate, CF

Accessory Name:

One-section leg plate, carbon fiber(optional), with special foam core pad, for HyBase 8300/8500

Application:

To support the leg with one-section plate during the surgery





Compatible with: **V8** HyBase

115-079672-00 A64XX 2-joint leg plate 115-079673-00 A64CX 2-joint leg plate, CF Accessory Name: Leg plate, dual-joint abduction, carbon fiber (optional), with special foam core pad, a pair **Application:** To support the legs of the patient during the surgery. A double-joint design helps doctors get closer to the surgical site



115-005685-00 A57XX One-section leg Plate, SFC 115-005683-00 A57XP One-section leg Plate, PU

One-section leg plate for PU or SFC (Special Form Core) mattress

To support the leg with one-section plate during the surgery

Accessory Name:

Application:



A57XP 6100 3000 Нувазе





115-079722-00 115-079723-00

Accessory Name: Leg plate, single-joint, o core pad, a pair

Application: To support the legs of t



115-085100-00 A68XX One-section leg plate 115-085101-00 A68CX One-section leg plate, CF

Accessory Name:

One-section leg plate, carbon fiber(optional), with special foam core pad

Application:

To support the leg with one-section plate during the surgery





115-085096-00 115-085097-00

Accessory Name: Leg plate, single-joint, c

core pad, a pair **Application:** To support the legs of th

Leg Accessories

V02XX Leg plate V02CX Leg plate, CF	Compatible with:
carbon fiber (optional), with special foam	
he patient during the surgery	

A85XX Leg plate A85CX Leg plate, CF	Compatible with:
arbon fiber (optional), with special foam	
he patient during the surgery	