

Assembly and operating instruction

vareflex



Dear customer,

In deciding to buy a nursing care bed from Bock you have opted for a care product that has a long service life and delivers first class functionality at the highest safety level. Our electrically adjustable care beds guarantee optimum comfort when lying, and support professional care activities. The focus is on people who need care, encouragement and protection. We have created the basic requirements for this with our care products. We urge you to prevent potential malfunction and risk of accidents by complying strictly with the safety and operating instructions and carrying out the necessary maintenance.

Yours sincerely,

Illans Rod

Klaus Bock

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1. Introduction and general information

The various bed systems that are made by Hermann Bock GmbH meet the special requirements for use in rehabilitation and therapy establishments as well as for care at home. Thereby reliable functionality and long-life-cycle are what characterize each single bed model as particularly high quality. Provided, that the bed is used in accordance with its purpose and serviced at regular intervals, the bed requires only a low level of maintenance. Each healthcare bed manufactured by the company Hermann Bock must pass a quality inspection and will be issued with a quality certificate by the TÜV before it leaves the production line. Hence, every healthcare bed meets the requirements of the directive 93/42/ EWG for medical devices (Class I). The beds have been manufactured and certified in accordance with the applicable standards for beds used for medical purposes.

Since April 2013, the standard applying to the beds has changed in line with the requirements of EN 60601-2- 52: 2010. The electrical component parts are in conformity with the safety standard EN 60601-1:2006 for medical devices.

The new standard distinguishes the beds between five different areas of application:

- 1. Intensive care in hospital, Intensive care bed
- 2. Short-term care in hospital or another medical facility, inpatient bed
- 3. Long-term care in medical environment, in-patient care bed
- 4. Home care treatment, sheer home care bed
- 5. Ambulant Care/Home care nursing service

1.1 Intended use

The health care bed has been designed for the positioning of persons in need of care or patients of medical facilities as of the year of 12 and a body height of at least 150 cm. The beds are intended for use in retirement or care retirement homes, rehabilitation facilities and with respect to home care treatment. Its purpose is to provide relief from disabilities and to facilitate the care process. Any other use is considered to be not intended; therefore all and any liability is excluded, if any damages can be attributed to any such unintended use.

The health care bed is not suitable for the use in hospitals. Besides, it has not been designed for the transport of patients. The beds must only be moved within the patient's room, for cleaning purposes or to enable the access to the patient.

Important: The beds are not equipped with any particular connections that provide for a potential equalization. Electrical medical devices connected to the patient's intravascular or intracardiac system must not be used. The operator of the medical device shall be responsible for the conformity of the combination of the devices with the requirements of DIN EN 60601-1:2006. This user's manual contains safety notes. All persons working with the beds must be acquainted with the contents of these instructions. The improper use may involve hazards.

1.2 Definition of person groups

Operator

Operators (e.g. medical supply stores, specialist dealers, facilities, and cost units) include all physical or juridical persons, who use the beds or have the beds used for medical purposes. The briefing on the use of the products shall generally be conducted by the operator.

User

Users are persons, whose training, experience, or briefing on the product allows them to operate the health care bed or carry out works on it. The user is capable of recognizing possible hazards or to prevent such from occurring and to assess the physical condition of the patient.

Patient / Resident

Persons in need of care, disabled or invalid persons lying in a care bed.

Professionals

Professionals include staff assigned by the operator, who are, owing to their training or briefing allowed to deliver, mount, dismount, and transport the bed. As a general rule, these persons must be instructed to the guidelines concerning the cleaning and disinfection of the health care bed.

1.3 Safety notes

The use of all moveable component parts in accordance with their intended use is not only crucial with respect to the hazard prevention for the patient but also when it comes to the safety of the relatives and/or the nursing staff. Another important aspect to be considered with respect to the operation of the bed is the individual physical condition of the patient and the kind and degree of their disability.

Please make sure that any hazards that might occur from unintended adjustments and incorrect operation are avoided by enabling the locking device. Whenever the operator, e.g. nursing staff or caring relatives leave the room, it is recommended to lock all operating functions of the bed; this can be done by means of the key at the hand control. For this purpose, the lying surface needs to be brought to the lowest position, and in a next step, the locking function can be enabled by means of the key, which can be found at the back side of the locking device. Just turn the key, pull it out and check, if the locking function is really working by trying the buttons of the hand control.

These recommendations are particularly important,

- > if the patient's disability hinders them to operate the hand control,
- > if the patient or nursing staff could be at risk due to unwanted adjustments,
- > if the side rails are raised, so that there is a risk of crushing or getting trapped,
- > if there are unattended children in the room.

Always pay attention that the hand control is hooked into the handle at the bed so that it cannot drop down. As a general rule, the bed should be operated by instructed nursing staff or relatives, or in attendance of instructed persons. When making adjustments to the lying surface, it should be made sure, that the patient's limbs are not positioned in the adjustment

area of the side rails. The patient's appropriate lying position is likewise important when it comes to adjustments to the side rails.

Prior to making any electrical adjustment, it should, as a general rule, be made sure that the patient's limbs are not positioned in the adjustment area between the chassis and the head- or foot board, resp. that there are no persons in the area between the floor and the raised lying surface. These areas exhibit a particular high risk of crushing injuries.

The permitted person weight depends on the total weight of the equipment that has been mounted to the bed (mattresses and other electronic medical devices). The respective max. safe capacity is specified on the name plate, which is attached to the frame of the lying surface.

1.4 Life time / guarantee

This nursing care bed has been developed, designed and constructed for a save and long use. In case of proper operation and use, the nursing care bed has an expected life time of approx. 2-10 years. The life time depends on the usage condition and frequency. Therefore a longer life time in the institutions is expected.

Attention:

In case of unauthorized technical modification of the product, all warranty claims extinguish.

This product is not approved for the North American market, especially the United States of America (USA). The distribution and use of this nursing care bed, also by third parties is prohibited by the manufacturer.

1.5 Type label (Example)

Model: xxx

Date of manufacturing: xx.xx.xxxx

 $xxx V \sim xx HZ my. xxx W or max. x A$

ED xx % (x min ON / xx min OFF)

Motor protection class OPX47

o□□□ = xxx kg

$$\frac{\triangle}{\triangle}$$
 = xxx kg

Hermann Bock GmbH - Nickelstr. 12 D-33415 Verl / Tel. +49(0)1805/262500



- (1) Model description
- (2) Date of manufacturing: Day, month, and year
- (3) Serial number: Order confirmation number serial number
- (4) Supply voltage; mains frequency; power input (Limoss) or current draw (Ilcon)
- (5) Switch on time
- (6) Drive protection type
- (7) Safe capacity/ max. person weight
- (8) Manufacturer
- (9) Symbols (on the right-hand side)

 ϵ

CE- mark in accordance with the directive for medical product

IPX4

Protection of the electrical equipment against splashing water



Medical application device type B



Only to be used in dry rooms



Protection class II (double isolation, protective insulation)



When disposed within European Union, the product disposed to a separate waste collection. The product must be disposed to the separate domestic waste



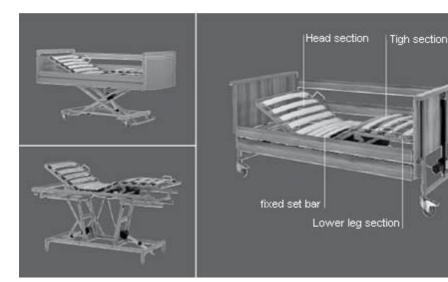
Symbol for max. person weight



Symbol for max. safe capacity



Symbol for attending the user's manual



2. General functionality description

Design configuration and functionality The lying surface and its four sections

In the standard version, the lying surface comes with comfort wooden slats (may be supplemented with a metal lying surface or special suspension systems) and is made up of four sections: head section, fixed seat support, upper - and lower leg section. The complete frame of the lying surface has been welded from steel tubes and stove-enamelled using a PES-powder coating. The electrical stepless variable height adjustment of the lying surface is controlled by means of 24 V-direct current motors and the smooth-running keys of the hand control. The head section can be electronically adjusted. The leg part consists of a two-part feet bracket. The stepless adjustment of the position can be made by means of the hand control. The control via the electronic hand control allows also for an automatic triple function for the stretched elevation of the legs towards the heart- and knee bend. In the event of a blackout, the back- and leg part can be lowered by means of a 9 V battery.

The chassis

The height adjustment of the beds can either be made via two height-adjustable actuators or a basic frame which can be operated via a single or double-drive. The surface of this steel tube construction is stove-enamelled with a PES-powder coating.

The side rails

Every health care bed comes with integrated side rails on both sides and therefore, exhibits a special degree of safety. The side rails can be raised and lowered by means of a steel bar. Owing to an integrated slider, the sliding blocks are particularly smooth and the ends are provided with a well-designed sealing cap. An ergonomically shaped trigger button allows for the easy operation of the side rails. According to the bed model, customers can choose between long and short side rails.

Bock-Hazard Notes

- Only use original bock side rails, which are available for every bock heath care bed
- Only use technical correct and undamaged side rails with the allowed distance.
- Make sure that the side rails lock in place safely.
- Control the side rails before every re-use. Make sure that the side rails and the mechanical parts of the beds are without damages.
- The operation of the side rails should always be carried out with the greatest care, in order to avoid bruises or other injuires of the fingers.

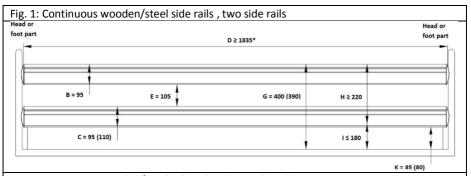


Fig. 2: Continuous wooden/steel side rails , three side rails

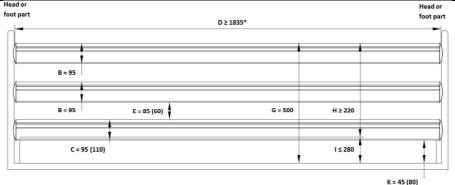


Fig. 3 Telescopic wooden side rails, duo middle bar

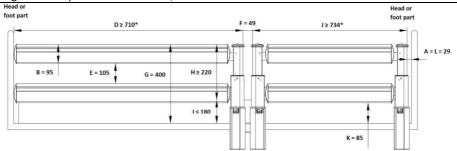


Fig 4: Telescopic wooden side rail, solo middle bar in the middle

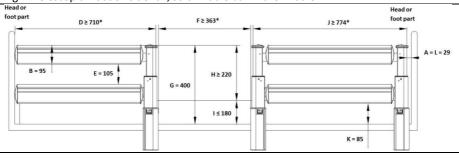
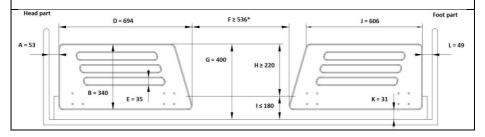


Fig. 5: Wooden plug-in/plug-on side rail



All measures in mm.

(*)Depending on the lying surface length

Single telescopic bar on the head/foot end are optional.

The measures in brackets is valid as optional.

Article numbers	
Description	Art.No.
Continuous wooden/ steel side rails	
2 side rails (Fig. 1)	
Wooden side rails (Set: 95 / 95mm)	90223
Wooden side rails (Set: 95 / 110mm)	91247
Steel side rails (Set: 95 / 110mm)	91314
Three side rails (Fig. 2)	
Wooden side rails (Set: 95 / 95mm)	91566
Wooden side rails (Set: 95 / 110mm	91531
Telescopic wooden side rails	
Duo telescopic bar in the middle (Fig. 3)	
Telescopic duo bar	91210
Wooden side rails head end :(set: 95/95mm)	80344
Wooden side rails foot end (set: 95 / 95mm)	80345
Solo telescopic bar in the middle (fig. 4)	
Solo telescopic bar (head right, foot left)	91211
Solo telescopic bar (head left, foot right)	91212
Wooden side rails (set: 95 / 95mm)	80346
Telescopic bar on head and foot end (without fig.)	
Solo telescopic bar (head right, foot left)	91211
Solo telescopic bar (head left, foot right)	91212
Continuous wooden side rails(Set:95/95mm)	80346
Plug-in/plug on wooden side rails (Fig. 5)	
Fitting plug-in	91264
Fitting plug-on	91260
Wooden side rail	80118

A: Distance between head end and side rail

B: Height 1 of the side rail

C: Height 2 of the side rails

D: Width 1 of the side rail

E: Distance between the elements

within the side rails

F: Distance between the splitted side rails

G: Distance between lying surface and upper edge of the side rail

H: Height of the upper edge of the side rail above unpressured mattress

I: Thickness of the mattress of intended use

J: Width 2 of the side rail

K: Smallest distance between side rail and lying surface (without side panneling of the frame if provided)

L: Distance between foot end and side rail

3. Electrical components

3.1 Drive unit

The drive unit consists of single drives for the electrical profiling of the moveable head and leg part. A switch-mode supply with rectifier is part of the external motor system. This switch-mode supply converts the input voltage of 110-240 V AC at 50-60 HZ at 70-180 W into a low voltage of 29 V DC. With this non-hazardous low voltage the motors and the hand control are operated. The cables are isolated twice and the power plug disposes of a primary fuse.

A performance adjustment provides a constant velocity. The safety demands therefore corresponds to the safety class II and the moisture protection IPX6.

In case that the maximum operation interval of two minutes is exceeded, due to e.g. continuous operation of the hand control, overheating of the actuators resulting in the immediate disconnection of the bed's power supply through the thermal fuse. It takes a cooling-down time of approx. one hour, until the power supply is automatically switched on again.

3.2 Locking device for all functions

The standard hand control with its 6 buttons comes with an integrated locking device enabling the nursing staff to lock all functions of the hand control by means of a key.

3.3 Level adjustment drive

The adjustment of the lifting appliance is effected through one or two integrated low-voltage direct current drives whose range of adjustment depends upon an integrated end-switch. The adjustment drive is connected with the control unit by means of a spiral cable.

3.4 The lockable hand control, fault safe operation

The extra-large, easily operable 6 buttons positioned on the ergonomically shaped hand control provide the main functions and can be controlled at the touch of a finger. Each of the operating buttons is labelled with appropriate symbols. As long as the button for the adjustment of the actuators is pushed, the actuators are operating. A spiral-shaped cable provides the necessary clearance whilst the operation is being performed. The rear side mounted clip is rotatable by 90° on both sides. The radius is exactly in line with the radius of the side rail and the lifter, so that there are no unsteady clearances. The possibly disturbing position of the hand control while performing cleaning or maintaining operations can be avoided simply by turning it to another side or easily clipping it onto any spot of the bed.

The Bock hand control

Function button 1 Head rest up
Function button 2 Head rest down
Function button 3 Foot part up

Function button 4 Foot part leg part down

Function button 5

Function button 6

Function button 7

Function button 7

Function button 8

Function button 8

Function button 9

Function button 9

Function button 10

Auto contur up

Auro contur down

Lying surface up

Anti-Trendlenburg

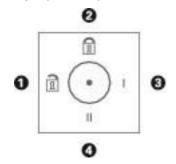
Trendelenburg



Bock Hazard note

The maximum switch-on time of 2 minutes should not be overstepped. A subsequent break of at least 18 minutes has to be observed.

Moreover, on the back side the hand control disposes of an integrated locking device. That can be activated by using the provided nurse key. For the setting of the electronic functions of the bed, just put the key into the lock on the back side and turn it to the desired function.





Switch setting 1
Switch setting 2 & 4
Switch setting 3

Hand control active Hand control inactive

Activation of Trendelenburg function

(with beds providing the Trendelenburg function)

3.5 Caution: Electrically operated drive

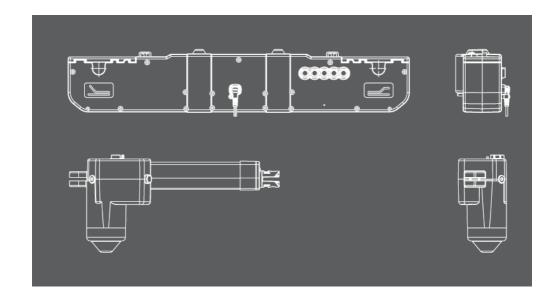
Hermann Bock calls its electrically operated nursing and therapy beds health beds, because they considerably facilitate the care recipient's recovery process in both physical and mental aspects while relieving pain at the same time thanks to their versatile functions. When applied as medical product, electrically operated beds require particular consideration with respect to the continuous safety inspections. These include the safe and professional handling of the bed, the daily check of the electrical equipment, and the proper maintenance and cleaning.

In order to avoid damages to the cables, the cable installation should be places off-side potential damage areas. Also avoid contact with square-edged components. Notes for an appropriate cable installation can be found in chapter 7.7. All potential risks of exposure to too high contact voltages should be excluded, as this helps to prevent injuries caused by any electrical shock. This may especially occur when the mains connection has been damaged, the leak currents are unacceptable or too high, or liquids have penetrated the motor housing, e.g. caused by improper cleaning.

Bock Hazard Note

The simultaneous use of electrical devices my cause, especially in the direct environment of the ready-to operate bed, low electromagnetic interactions between the electrical devices, such as radio noises.

When such a rare case occurs, you should extend the distance between the devices. Do not use the same wall socket or switch off the noisy device temporarily. If the bed is not operated in line with its purpose, thus simultaneously with electrical, medical devices, you should deactivate the functions of the bed for the time being. The deactivation can be done by the integrated locking device on the back side of the hand control.



4. The drives

4.1 The 24 Volt drives

Hermann Bock provides their health care beds with various drive systems.

4.1.1 The drives system

The dual drive and actuator consists of 4 main components.

- Housing
- Drive
- Gearbox
- Spindle with nut

The housing principle and its double drive and the single drive guarantees the permanent function of all drive components. The special construction design is based on two load absorbing housing cases. Owing to a detailed internal engineering, the patented design of the inner housing constitutes an essential requirement for the precisely fitting intake of the drive technology. We do not use pre-assembled components. The totally easy assembly/disassembly and the spacious installation compartment for battery and electronics positioned above the robust hinged cover make the housing of the double drive stand out. The drive comes with a mains isolation in the mains plug and has an emergency lowering function.

4.2 The external switch-mode power supply SMPS

The SMPS wall power supply (switch-mode power supply) is an electronic transformer, which has an integrated performance control. A constant voltage until the maximum load (without lost in velocity) and a safety against overload are given. The external transformer offers safety beginning at the wall socket, as there the line voltage is directly converted into 29V low voltage, with which the bed is operated. The transformer is connected to the motor cable and can be changed separately in case of damages.

The wall power supply already corresponds to the upcoming new European standards for electronic domestic appliances. In the standby-mode it has an energy consumption of max. 0.5 W. Due to its variable input voltage of 100 V-264 V it is applicable worldwide. With the SMPS power supply alternating electrical magnetic fields are not measurable and in use comparing to motors with mains isolation even lower (due to the concurrent flow).

Bock Hazard Note

Never open any drive components! Both repair and exchange ofn components are only allowed to be done by especially authorised experts.

Bock Top Advice

Once a year the 9 Volt battery of the motor should be tested and if necessary exchanged. Furthermore regular visual inspections should be done.



5. Assembly and operation

5.1 Technical data

Technical o	data		Vareflex
Lying surface dim	ension: cm		90 x 200
External dimension	n: cm		103 x 220
Max. safe capacit	y: kg		170
Max. person weig	ht: kg		135
Height adjustmen	t: cm		30 - 80
max. indicated a	ngel toward	ls horizontal:	_
- Back bar			70°
- Foot bar			20°
- Trendelenburg-	Position (opt	ional)	Not possible
Side rail height wi	th wooden s	lats: cm	39
Possible side rai	l solutions:		
- Continuous woo	oden/steel si	de rails	•
Lifter space: cm			>15
Sound leval: dB(A	۸)		< 65
Weights:			
Total weight incl.	wooden side	rails: kg	83,5
Lying surface: kg			23 + 17,8
End panels: kg			2x 24
Continuous wood	en side rails:	: kg /set	12,5
Continuous wood	en side rails:	: kg /set	15,8
Electrical data			
Motor		Input voltage: V	100-240
		Frequency: Hz	50/60
		max. preformance: W	90



5.2 Vareflex

Vareflex offers a high lying offers a high lying comfort for people with the need of care, fragile or handicapped people. Due to its easy operation it supports the care ideally.

- > Vareflex is not suitable for the use in hospitals.
- > Vareflex is not suitable for the transport of the patient. The beds are simply defined for the moving inside the patient's room, e.g. for cleaning or access to the patient.
- > Vareflex is suitable for persons up from twelve years and a body height of 150 cm.
- > Under certain circumstances, Vareflex can (if required) be used in combination with medical purposes and other electrical medical devices (e.g. draining devices, ultrasound nebulizers, nutrition systems, anti-decubitus systems, oxygen concentrators, etc.). In this case, it would be necessary, to deactivate all bed functions by means of the integrated locking device, until the treatment is completed.

Important: The beds are not equipped with any particular connections that provide for a potential equalization. Electrical medical devices connected to the patient's intravascular or intracardiac system must not be used. The operator of the medical device shall be responsible for the conformity of the combination of the devices with the requirements of DIN EN 60601-1:2006.

Particular features

Vareflex offers modern and reliable technique combined with easy operations of different functions.

Vareflex is optionally available with a lying surface which provides both 4 or 5 different adjustment functions. The electrical adjustment of the lying surface's back rest can be done by means of the hand control. In case of the 4- resp. 5-sectioned lying surface, the electrical control of the back- and leg part takes place by means of the hand control including its automatic tripple function.

Getting Vareflex ready for use

Please remove all packaging leftovers from the health care bed before carrying out the assembly.



Remove the two lying surface parts from the transport system and put them together. Fasten them with the before removed screws and the provided Allen key.



Put the lying surface on the floor. Now lift the lying surface on one side and put it into the triangular connecting links of the end panels. Do not click it into place correctly, but leave it as in the picture on the right side. Repeat the same with the other side.



This is how the end panels and the lying surface should be connected.



Assemble the side rails (wooden or steel) on one end panel. Therefore you need to put the side rails on the side rail fittings. (The labelling up and down has to be followed).



Afterwards put the side rails on the other side on the side rail fitting. Thereby correct the cap between side rails and the end panel.



Now let the lying click into place completely with the end panels.



Now lock the lying surface on all 4 corners.



Connect the single motors.

- 1. End panel foot part
- 2. End panel head part
- 3. Lying surface foot part
- 4. Lying surface head part
- 5. Hand control



Screw the cover cap onto the control box.

The net cable has to be fastened to the lying surface with the provided pull relief. Connect the mains plug.

After the assembly process resp. prior to the initial operation of bed, it will be necessary to run the adjustment area of the lying surface using the hand control, in order to check for the ideal positioning of the cables. The adjustment area must be accessible without any obstacles.

6. Accessories

As it is our goal to satisfy every need of our customers, Hermann Bock offers a wide range of practical and mobility-promoting accessories, so that each health bed can be exactly customized to the individual needs of the care recipient. The assembly is done in a quick and easy manner using the fixing points on the beds that have already been prepared for this purpose. It goes without saying, that every element of our additional accessories meets the special quality and safety standards of Bock. The bed extensions available for lengths of up to 220 cm makes it also possible for tall people to benefit from the high lying comfort with equal functionality. In addition to the standard equipment included in the delivery as basic equipment, you can also choose from our variety of accessories, which is available for each bed model. This optional accessory varies depending on the bed model and is fitted to its special functions and place of use. The range stretches from technical elements over mattresses up to the occasional extra bed. A wide offer of wooden colours and a variety of colours allow for the harmonious integration of each health bed with any kind of furniture.

6.1 Special dimensions

Special dimensions are an essential part of the manufacture at Hermann Bock.

Ideal lying comforts for persons in need of care who have a particular physique can only be achieved by means of custom-built models. With its customized models, Hermann Bock enables customers to have their health bed tailored to fit the individually physical constitution of the person in need of care. For body heights up from 190 cm, Hermann Bock recommends the employment of a bed extension that allows an extension of the lying surface to a length of up to 220 cm. That way, the high lying comfort can also be ensured for tall persons, and, of course the functionality remains the same.

6.2 Assembly - Bed extensions (therapy beds)

The scope of delivery consists of:

- 2 Adapter units for left and right foot part
- 1 wire bracket for the foot part
- 1 set of side rails
- Fixation screws

How to carry out the easy assembly

- 1. Take off the mattress from lying surface.
- 2. Remove foot end panel.
- 3. Plug adapter units into the frame of the lying surface at the foot end and fasten with screws.
- 4. Put wire bracket into the frame of the lying surface, drill holes (d = 4.2 mm) and fasten with screws.
- 5. Do not slide the foot end panel further than shortly before the release button.
- 6. IMPORTANT: Make sure to read the labels attached on the top and bottom of the side rails' end caps, as these must not be confounded with each other.
- 7. Than hook the side rails into the pre-assembled metal guides.
- 8. Fasten the end panels correctly.



Bock Hazard Note

The bedside use of accessories or medically necessary appliances, e.g. I.V.poles requires the nursing person's careful attention with regard to the avoidance of crush and shear zones to the care recipient when adjusting the back or leg rests.

6.3 Assembly- Accessories

The following standard equipment can be combined with the bed models:



Side rail attachment

Scope of delivery: side rail attachment, completely mounted

 Open the plastic cap, plug in the side rail attachment, position it into the middle and close the cap. Please make sure, that the release button of the side rail extension faces outwards.

Important note:

The bock side rail extension has been designed for the use on all bock wooden side rail models. Company Bock assumes no liability for damages arising from the use in combination with third-party products!

Lifting pole with triangle grip, 6,5 kg (Fig. 2)

The safe capacity of the lifting pole amount to max. 75 kg.

Scope of delivery: 1 lifting pole 1 triangle grip

- Insert the lifting pole into the provided connector on the head end of the lying surface.
- The height adjustable area of the triangle grip is not allowed to fall below <= 550 mm up >= 700 mm, measured from top of the mattress (mattress height 100mm and 120 mm) to the button line of the horizontal grip.

ATTENTION: Do not swing the lifting pole outside of the lying surface.

Under normal use the triangle grip has a durability of at least five years. We refer to the safety, technical tests.

Side rail bumpers, 1.4 kg

Scope of delivery: 1 cover 1 bumper

 Open the zipper of the bumper and slip it over the side rail and close the zipper or velcro again



Tray, 4.0 kg (Fig. 4) Scope of delivery:

 The tray is to be placed on the side rails and is secured against slipping by two distance holders



Universal clamp, (0,6 kg)

Scope of delivery: 1 Clamp, 1 fastening ring

The universal clamp is a special holding appliance that provides more flexibility as basis element and allows for the flexible positioning of the modular functional equipment. It is optionally possible to attach quivers, fixtures for urine bottles, infusion systems or a lamp individually or together. Furthermore, the universal clamp can be shifted along the side rails according to preference or requirement.

Universal clamp with: drainage holder, urine bottle holder, with

flexible tube and hand control holder (from left to right)

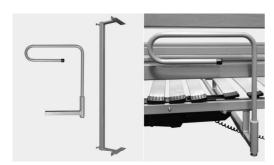
 The universal clamp is assembled to the top side rail and fastened with the mounting ring.







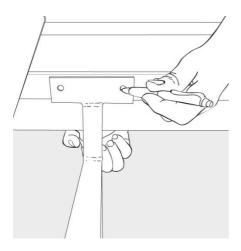
Assist handle for beds with the actuator in the end panels 3,0 kg Scope of delivery: 1 Getting-up aid incl. 1 cross bar support, 4 bolts 4 mm



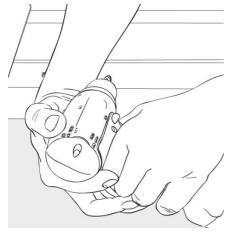
Left side: Scope of delivery, right side: mounted assist handle

Bock Hazard Note

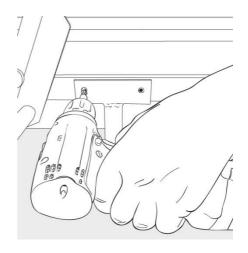
For safety reasons only use original Hermann Bock accessories for the corresponding bed models. A detailed overview can be found on a separate data sheet. Hermann Bock does not assume any reliability for accidents, damages or risk caused by a third party accessory!



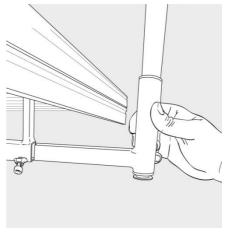
Put the cross-bar support to frame of the lying surface from the bottom up and mark the drill holes on the frame.



Drill holes into the previously marked spots (3.5 mm) of the lying surface frame.



Use the provided bolts to fasten the cross-bar with the lying surface frame.



Push the assist handle into the cross-bar bracket adjust it to the desired position and fasten it.

6.4 Mattresses

There are, in general foam and latex mattresses suitable for the Hermann Bock health beds. However, a volumetric weight of at least 35 kg/m3 is required along with the dimensions of 90×190 cm, 100×190 cm, 90×200 cm and 100×200 cm. The height of the mattress which is allowed to be used:

with steel-, aluminium bars, wooden slats or ripoplan lying surface 15 cm (for interior furnishing beds 16 cm)

lying surfaces with a spring system 12 cm

If the height limit is exceeded, an additional side rail attachment is required which is available as additional equipment. When using foam mattresses, it is recommended to use a cut foam mattress pad to allow a better fitting with the lying surface.



7. Cleaning, maintenance and disinfection

The several bed elements consists of hight-quality materials. The surface of the steel tubes is covered with a durable PES powder coating. All surfaces of the wooden parts are sealed with an ecologically compatible overlay. All bed elements are easy to clean and cared for using wipe and spray disinfection means according to the applicable cleaning requirements with respect to the various areas of application. Observing the following care instructions will retain the usability and visual appearance of your nursing bed for a long time.

7.1 Cleaning and care

Steel tubes and vanished metal parts:

Please use a wet wiper and a standard, mild household detergent for the cleaning and care of these surfaces.

Wooden-, decorative-, and plastic elements:

All standard furniture cleaners and cleaning detergents can be used. The cleaning of the plastic elements using a wet wiper without detergent additives should generally be sufficient. For the care of the plastic surfaces you should use a product, which is specifically suitable for plastics.

Drive:

In order to prevent the intrusion of moisture into the drive, it is recommended to use a slightly moist wiper to clean the housing of the drive.

Lying surface systems ripolux, ripoplan and ripolux neo:

Use a moist wiper without adding any detergents or, if deemed necessary, a detergent which is exclusively suitable for plastics to clean the supporting- and spring elements as well as the plastic surfaces. In case of heavy contamination, just remove the spring elements from the supporting elements . The dismounted plastics elements can be rinsed or spray-washed with hot water to get them clean. As regards the disinfection, the components should be sprayed with a detergent suitable for plastics. Most of the moisture drips off the plastic surface by slightly shaking it, while the rest will dry on its own within a short time. Remount the elements after they have completely dried. If required you can also remove each of the individual lying surface elements from the frame to clean them. Remount the elements after they have completely dried. If required you can also remove each of the lying surface elements from the frame to clean them.

7.2 Disinfection

All methods in accordance to the standard EN 12720 can be used for the wipe disinfection. However, you should apply only mild and gentle methods so as to retain the material resistance of the plastic elements such as the drive housing, decorative elements, ripolux and ripolan. Concentrated acids, aromatic and chlorinated hydrocarbons, as well as detergents containing highly concentrated alcohol, ether, ester and ketone may damage the material and should therefore be avoided.

7.3 Hazard avoidance

Please make sure to consider the following guidelines with respect to the electrical component parts of your nursing beds as it is crucial to avoid hazards related to cleaning and disinfection. The non-observance of these guidelines may result in considerable damage of the electrical lines and the drive.

- 1. Disconnect the mains supply and position it in such a way that contact with excessive amounts of water or detergents can be excluded.
- 2. Check all plug-connections for correct position according to the instructions.
- Check the wires and electrical component parts for damages. Should you detect any damages, do not perform any cleaning operations, but first have the defects repaired by the manufacturer or authorized staff.
- 4. Check the mains supply for residual moisture before starting the operation and dry or blow out the device, according to need.
- 5. On any suspicion of the intrusion of moisture into the electrical components, disconnect the mains supply immediately and do not re-establish the connection. Put the bed out of operation immediately, attach an appropriate visible labelling and get in contact with the manufacturer/supplier.

Bock Hazard Note

It is absolutely not recommended to use abrasive cleansers resp. detergents containing grinding particles, cleaning pads or stainless steel cleaners for the cleaning. Do neither use organic solvents such as alkyl/aromatic haloids and ketones nor detergents containing acid or alkaline. Never clean the bed using a water hose or high-pressure cleaner, as this might lead to the intrusion of fluid into the electrical components which causes malfunctions and hazards.

Bock Top Advice

Scrapes and varnish chippings that go through the entire varnish coating should be preventively sealed with appropriate repair means against the infiltration of moisture.

Guidance and manufacturer's declaration

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

Emissions test	Compliance	Electromagnetic environment - guidance
RF emissions CISPR 11 (partly)	Group 1	The medical used bed 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 equipment.
RF emissions CISPR 11 (partly)	Class B	The <i>medizinisches Bett</i> is suitable for use in all establishments other than domestic and those directly connected to the public-voltage power supply network that supplies buildings used for domestic purposes.
Harmonic emissions IEC 61000-3-2	Class A	
Voltage fluctuations/flicker emissions IEC 61000-3-3	Complies	

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

Immunity test	IEC 60601 test level	Compliance level	Electromagnetic environment – guidance
Electrostatic discharge (ESD) IEC 61000-4-2	± 6 kV contact ± 8 kV air	± 6 kV contact ± 8 kV air	Floors should be wood, concrete or ceramic tile. If floors are covered with synthetic material, the relative humidity should be at least 30 %.
Electrostatic transient/ burst IEC 61000-4-4	± 2 kV for power supply lines ± 1 kV for input/output lines	± 2 kV for power supply ines ± 1 kV for input/output lines	Mains power quality should be that of a typical commercial or hospital environment.
Surge IEC 61000-4-5	± 1 kV differential mode ± 2 kV common mode	± 1 kV differential mode ± 2 kV common mode	Mains power quality should be that of a typical commercial or hospital environment.
Voltage dips, short interruptions and voltage variations on power supply input lines IEC 61000-4-11	< 5 % U, (>95 % dip in U,) for 0,5 cycle 40 % U, 160 % dip in U,) for 5 cycles 70 % U, 30 % dip in U,) for 25 cycles < 5 % UT (>95 % dip in U,) for 5 sec	< 5 % U ₁ (>95 % dip in U ₁) for 0,5 cycle 40 % U ₁ (60 % dip in U ₁) for 5 cycles 70 % U ₁ (30 % dip in U ₁) for 25 cycles < 5 % U ₇ (>95 % dip in U ₁) for 5 sec	Mains power quality should be that of a typical commercial or hospital environment. If the user of the medizinisches Bett requires continued operation during power mains interruptions, it is recommended that the medizinisches Bett be powered from an uninterruptible power supply or a battery.
Power frequency (50/60 Hz) magnetic field IEC 61000-4-8	3 A/m	3 A/m	Power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercial or hospital environment.

NOTE: U, is the a. c. mains voltage prior to application of the test level.

Electromagnetic immunity

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

Immunity test	IEC 60601 test level	Compliance level	Electromagnetic environment – guidance
Conducted RF	зV	з۷	Portable and mobile RF communications equipment should be used no closer to any part of the EQUIPMENT medizinisches Bett, including cables, than the
IEC 61000-4-6			recommended separation distance calculated from the equation applicable to the frequency of the transmitter.
Radiated RF	3 V/m	3 V/m	_
IEC 61000-4-3			Recommended separation distance: $d = \left(\frac{7}{E_1}\right)\sqrt{P}$
			$d = [\frac{3.5}{E}] \sqrt{P}$ 80 MHz to 800 MHz $d = [\frac{7}{E}] \sqrt{P}$ 800 MHz to 2,5 GHz where p is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer and d is the recommended separation distance in metres (m).* Field strengths from fixed RF transmitters, as determined by an electromagnetic
			site survey, a should be less than the compliance level in each frequency range. Interference may occur in the vicinity of equipment marked with the following
			symbol:

- 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.
- 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 Breakington and the predicted theoretically with accuracy. To assess the electromagnetic environment due to fixed Br transmitters, an electromagnetic site survey should be considered. If the measured field strength in the location in which the medizinisches Bett is useed exceeds the applicable RF compliance level above, the medizinisches Bett should be observed to verify normal operation. If shormal performance is observed, additional measures may be necessary, such as reorienting or relocating the medizinisches Bett.
- b Over the frequency range 150 kHz to 80 MHz, field strengths should be less than [V,] V/m.

Recommended separation distances between portable

and mobile RF communications equipment and the medizinisches Bett.

The medizinisches Bett is intended for use in an electromagnetic environment in which radiated RF disturbances are controlled.

The customer or the user of the medizinisches Bett can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and the medizinisches Bett as recommended below, according to the maximum output power of the communications equipment

	Separation distance according	rding to frequency of transr	nitter
Rated maximum output of transmitter W	150 kHz to 80 MHz $d = \left[\frac{3, 5}{V_{1}}\right] \sqrt{P}$	80 MHz to 800 MHz $d = \left[\frac{3, 5}{E_1}\right] \sqrt{P}$	B00 MHz to 2,5 GHz $d = \left[\frac{7}{E_1}\right] \sqrt{P}$
0,01	0.12	0.12	0.23
0,1	0.37	0.37	0.74
1	1.17	1.17	2.33
10	3.69	3.69	7.38
100	11.67	11.67	23.33

For transmitters rated at a maximum output power not listed above the recommended separation distance d in metres (m) can be estimated using the equation applicable to the frequency of the transmitter, where P is the NOTE 1

At 80 MHz and 800 MHz, the separation distance for 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.

8. Declaration of conformity

Manufacturer: Product description/ model

Hermann Bock GmbH Nickelstraße 12 33415 Verl Medical used bed in general

<u>Classification:</u> <u>Choosed conformity evaluation process:</u>

Medical products class I, norm 1 and 12 referring to appendix IX of MDD

Appendix VII of MDD

Hereby we declare that, the above specified products fulfill the precautions of the guideline 93/42/EWG of the advice for medical products. The entire associated documentation is kept in the premises of the manufacturer. This declaration of conformity is valid from 1.4.2013.

Applied standards: Harmonized standards for which the proof of concordance

can be supplied:

DIN EN 60601-1:2007 Medical electronic devices- Part 1:

General definitions for the safety including the essential

characteristics

DIN EN 60601-1-2:2007 <u>Medical electronic devices- Part 1-2:</u>

General definitions for the safety including the essential characteristics – complement standard: electromagnetic

tolerance - requirements and testing

DIN EN 60601-2-52:2010 Medical electronic devices- Part 2-52:

Special definitions for the safety including the essential

characteristics for medical beds

DIN EN ISO 14971:2013 Application of the risk management for medical products

Verl, 26. February 2014

Klaus Bock (General Manager) Dr. Stefan Kettelhoit (General Manager)

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9. Continuous functionality check including service

The safety standards of an electrically operated nursing bed are subject to the compliance with the specified European standards. This includes the manufacturer's strict adherence to the specifications as well as official standards defined by the government which are in accordance with the safety recommendations of the BfArM (Federal institution for drugs and medical devices) for the enforcement of the Medical Products Act. Regularly conducted inspections ensure the maintenance of high safety standards and in order to avoid hazards from occurring, the continuous and strict adherence to the regular inspection of the proper functionality is mandatory. The manufacturer may have no influence on the operator's adherence with respect to the observance of these instructions concerning the beds. However, Bock facilitates the observance of the necessary precautionary measures to be taken by means of their time-saving services.

The execution of the inspection, assessment, and documentation must be performed only by or under supervision of professional persons such as electricians or electro-technically instructed persons who have a thorough knowledge of the relevant provisions and are able to recognize possible impacts and hazards. In case that there is no suitable person on part of the operator in order to perform the Safety-technical control, Bock's service offers you to carry out the Safety-technical control including check and observance of the respective inspection terms for a charge. It is stipulated by the company Hermann Bock GmbH to execute an Safety-technical control for at least once a year and before and after each re-use of the bed. In order to facilitate the execution of all necessary safety inspections, the company Hermann Bock GmbH provides you with the Safety-technical control-checklist which can be found in the assembly- and operation manual. Please make a copy of the checklist as a form for your safety-technical inspection. The Safety-technical control-checklist serves as evidence report of the performed inspection and needs to be kept on file. The Safety technical control-checklist is also available as download from our website: www.bock.net.

Bei nicht autorisierten technischen Änderungen am Produkt erlöschen alle Garantieansprüche.

Bock Hazard Note

The bed has to be cleaned and disinfected prior to every re-use. This provision is accompanied by the requirement of a visual inspection which needs to be carried out in order to prevent mechanical damages.

Bock Top Advice

The Bock-Safety-technical control-training takes place either on your site or at ours and trains your technical staff in the performance of the inspection of the Safety-technical control on Bock health beds, so that they will be in the position to carry out safety-technical inspections in an appropriate way.

Constant functionality checks In accordance with MPBbetriebV, BGV A3 and DIN EN 62353 (electrical measurements) Test specimen: O Bed O Insert frame O Controller/ Main drive Model name: Serial/inventory number: Location Person in charge: Visual, mechanical and electrical examination Result Is the overall condition of the bed alright? Yes No Description of defects: 2. Are the address/ type labeling on the bed and on the motors visible? Yes No Description of defects: Manufacturer's details such as safety guidelines and assembly or operating No 3. Yes instructions present? Description of defects: 4. Mechanical construction defect free and without torn welding seams? No Yes Description of defects: 5. Yes No Firm fit and completeness of all plastic end caps and mechanical connecting elements (screws etc.)? Description of defects: Wooden slats, carrier plates and dowels for ripolux/ripoplan without cracks or breakages? Yes No Description of defects: Tight fit in correct position of all sprung slats and carrier plates? No Yes Description of defects: 8. Tight fit and straight alignment of all spring elements? Yes No Description of defects: 9 Preassure load of the spring elements? Yes No Description of defects: Tight fit and no cracks or breakages of head and foot end panels? 10. Yes No Description of defects: Adjusting space of lying surface and room for lifting height sufficient without obstructions at 11. Yes No current location? Description of defects: 12. Safe grid mechanism of lower leg section in every step even under charge? No Description of defects: No 13. Side rail bars without cracks, breakages or damages? Yes Description of defects: 14. Adequate fastening and respectively secure fit of the side rail bars? Yes No Description of defects: 15. Load test of the side rails without distortion? No Yes Description of defects: 16. Easy run of side rail bars within the tracks and easy locking? Yes No Description of defects:

Bed-accessories (lifting pole, triangle grab handle, belts, control box etc.) without damages

No

No

No

No

Yes

Yes

Yes

Yes

Correct functions of side rails?

Distance between side rail bars max. 12 cm?

Height of side rails above mattress at least 22 cm?

Description of defects:

Description of defects:

Description of defects:

and with secure fixing?

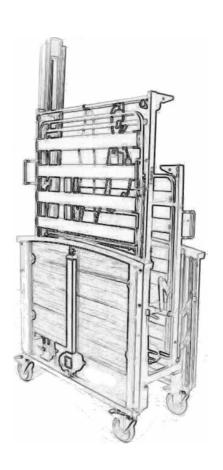
17.

18.

19.

20.

	Description of defects:				
21.	Safe breaks, arresting and free running of wheels?		Yes		No
	Description of defects:				
22.	Mains cable, connecting cables and plugs without scratches, dents, kinks, porous		Yes		No
	parts or bare wires?				
	Description of defects:				
23.	Pull relief fastened and efficient?		Yes		No
	Description of defects:				
24.	Internal plugs fully inserted and connected with strain relief?		Yes		No
	Description of defects:				
25.	Mains cable and plug without damage?		Yes		No
	Description of defects:				
26.	Correct and secure cable leading and cable connections?		Yes		No
	Description of defects:				
27.	Housings of motors and hand controls sealed and without damages?		Yes		No
	Description of defects:				
28.	Leak-prevention of motor for models older than 2001 present?		Yes		No
	Description of defects:				
29.	Motor lifting poles without damages?		Yes		No
	Description of defects:				
30.	Testing of hand controls: all buttons fully usable?		Yes		No
	Description of defects:				
31.	Testing of disabler on hand control: everything correct?		Yes		No
	Description of defects:				
32.	Testing of battery: faultless function?		Yes		No
	Description of defects:				
33.	Resistance of protective conductor: not applicable, because no protective conductor		Yes		No
	present (security class II)				
	Description of defects:				
34.	Resistance of isolator (for old appliances)				
	(initiate proof voltage and measure resistance; measured value must be more than $7~\text{M}\Omega$):				
	Description of defects:				
35.	Alternative leakage current, maximum value		Yes	1	No
33.	(device over 200 V, security class II, type B, threshold value = 0,1 mA):	Ш	163	_	NO
36.	Description of defects: Exceeds the patient-, mattress and accessory weight the assigned safe capacity (see		Yes	1	No
30.	technical data)?		res	<u></u>	INO
	Description of defects				
	Description of defects:				
Oue!	I condition of the hade exempting foultless?		V	_	No
overal	I condition of the bed: everything faultless?		Yes	<u> </u>	No
\lat					
Notes:					
Place a	and date:				
Signati	ure of examinant:				
Next e	xamination:				





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Subject to technical modification

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Our SALES PARTNERS:

Our business partners pursue the same strategy as we do, thus quality, innovation and outstanding standards, which are approved worldwide. We can rely on our business partners as you can rely on us.

Please note that repairs, spare part deliveries, trainings, safety controls and services can only be carried out by authorized personnel and our sales partners.

Otherwise all warranty claims will extinguish.

Here a small extract:

Belgium AXAMED nv-sa | www.axamed.be
Danmark Medema Danmark AS | www.medema.dk

Dubai Bridgeway Medical S. | www.bridgewayhealthcare.com

Great Britain Carebase | www.carebase.net
Estonia ITAK Ltd. | www.itak.ee

Finland RESPECTA OY | www.respecta.fi

Greece Wheel Rehabilitation Products | www.wheel.gr Israel Ugality of Life Center | www.iqlc.com

Italy Prontomed | www.prontomed.it
Croatia BEZ LIMITA d.o.o. | www.bezlimita.hr

Lebanon ALBERT MASSAAD s.a.r.l. | www.albertmassaad.com

Luxembourg Stoll | www.matelas.lu

New Zealand Cubro Ltd. | www.cubro.co.nz Neherlandsd Eureva B.V. | www.eureva.nl

Norway Medema Norge AS | www.medema.no Austria Reha Service GesmbH | www.rehaservice.at

Poland Timago International Group Sp. z o.o. | www.timago.pl

Portugal MACHADOS | www.machadosmadeira.com

Romania Donis srl | www.donis.ro

Russia Lazerlink

Swiss Sodimed | www.sodimed.ch

Serbia Proxi-Med d.o.o | www.proxi-med.co.rs
Slovenia Medimaj d.o.o. | www.medimaj.com
Slovakia Servis Invo | www.servisinvo.sk
Bock Spain Ferran Asensio Jou | www.bock.net/es/

Czechia Ortoservis I www.ortoservis.cz

Ukraine ADS Ukraine