

ergoselect 100/150/200

Bicycle Ergometer

Operator's Manual

201000134000 • Version 2016-09-23 / Rev 02 • English



This manual was written with the utmost care. Should you still find details that do not correspond with the system, please let us know and we will correct the issue as soon as possible.

We reserve the right to modify the design and technical features of the device and are not bound by the information and illustrations provided in this manual.

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This manual is not subject to any change order service. Please contact the manufacturer for the latest document revision.

ergoline GmbH Lindenstraße 5 72475 Bitz Germany

 Tel.:
 +49-(0) 7431 98 94 - 0

 Fax:
 +49-(0) 7431 98 94 - 128

 e-mail:
 info@ergoline.com

 http:
 www.ergoline.com

Printed in Germany



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GENERAL INFORMATION

- The product ergoselect bears the CE marking CE-0123 (Notified Body: TÜV), indicating its compliance with the provisions of the Council Directive 93/42/EEC about medical devices and fulfills the essential requirements of Annex I of this directive.
 The CE marking covers only the accessories listed in the Order Information chapter.
 The ergometer is an MDD class IIa product.
- The device fulfills the requirements of standard EN 60601-1 "Medical Electrical Equipment, Part 1: General Requirements for Safety" as well as the interference protection requirements of standard EN 60601-1-2 "Electromagnetic Compatibility – Medical Electrical Devices". The radio-interference emitted by this product is within the limits specified in EN 55011, class B.
- The symbol 🔲 means: protection class II
- This manual is an integral part of the equipment. It should be available to the equipment operator at all times. Close observance of the information given in the manual is a prerequisite for proper device performance and correct operation and ensures patient and operator safety. Please note that information pertinent to several chapters is given only once. Therefore, read the manual once carefully in its entirety.
- The symbols 🛕 🚱 mean:

Consult accompanying documents. They indicate points which are of particular importance in the operation of the device.

- Observance of the safety information protects from injuries and prevents inappropriate use of the device. All equipment users and persons responsible for assembly, maintenance, inspection and repair of the device must read and understand the content of this manual, before using or work on it. Paragraphs with special symbols are of particular importance.
- If unauthorized individuals open the control terminal, damaging the calibration sticker, any warranty claim shall become void.
- This manual reflects the equipment specifications and applicable safety standards valid at the time of printing. All rights are reserved for devices, circuits, techniques, software programs, and names appearing in this manual.
- On request ERGOLINE will provide a Service Manual.

- The ERGOLINE quality management system complies with the standards ISO 9001: 2008 and EN ISO 13485: 2003-AC2007.
- The safety information given in this manual is classified as follows:

Danger

indicates an imminent hazard. If not avoided, the hazard will result in death or serious injury.

Warning

indicates a hazard. If not avoided, the hazard may result in minor injury and/or product/property damage.

Caution

indicates a potential hazard. If not avoided, the hazard may result in minor injury and/or product/property damage.

- To ensure patient safety, the specified measuring accuracy, and interference-free operation, we recommend using only original ERGOLINE accessories. The user is responsible if non-ERGOLINE accessories are used.
- ERGOLINE is responsible for the safety, reliability, and performance of the equipment, only if
 - modifications and repair are carried out by ergoline GmbH or by an organization expressly authorized by ergoline GmbH
 - the equipment is used in accordance with the instructions given in this operator's manual.

ergoline GmbH Lindenstrasse 5 72475 Bitz Germany

Phone: +49-(0)-7431 - 9894 -0 Fax: +49-(0)-7431 - 9894 -128 e-mail: info@ergoline.com http: www.ergoline.com

SAFETY INFORMATION

Danger

• Explosion Hazard •

The device is not designed for use in areas where an explosion hazard may occur.

Explosion hazards may result from the use of flammable anesthetics, skin cleansing agents or disinfectants.

Warning Patient Hazard, Equipment Damage •

Do not expose the ergoselect to direct sunlight to prevent system components from reaching inadmissible high temperatures.

Do NOT use the ergoselect outdoors (medical device). Furthermore the device has no additional protection against the ingress of humidity. Humidity inside the device may cause equipment malfunctions and increases the risk of an electric shock.

Additionally, the device should not be operated in the vicinity of electric power plants, because they may impair equipment functions.

The ergoselect ergometer may only be used in combination with accessories approved by ergoline GmbH.

Risk to Persons

Before using the ergometer, the operator must ascertain that it is in correct working order and operating condition. The cables and connectors, in particular, must be checked for signs of damage. Damaged parts must be replaced immediately, before use.

• Equipment Malfunction •

Only the special shielded cables supplied by ERGOLINE may be used to connect the device to other pieces of equipment.

• Equipment Malfunction •

Cellular telephones may not be used in the immediate vicinity of the ergometer, because they might interfere with the proper functioning of the ergometer.

Electromagnetic interference most probably exists when the watt reading is unstable. If the displayed value changes frequently even though the speed is above 30 RPM, this may be due to electromagnetic interference.

Warning

Shock Hazard

When the ergometer is connected to other equipment or if a medical system is created, it must be ensured that the added leakage currents do not present a hazard. In case of questions, please contact your ERGOLINE dealer or the ergoline GmbH Service Department.

For use, the ergometer must always be connected to electric installations that fulfill the local requirements.

• Patient Hazard •

The German Medical Device Operator Ordinance (MPBetreibV, § 5) demands that users

- must be trained in the use of the ergometer
- must be familiar with the routines for handling and assembly of the ergometer
- must be familiar with and observe the safety rules and regulations for operation of this type of equipment
- must be informed about any other pertinent rules and regulations (e.g. safety features)
- must be informed about the potential hazards arising from the use of this type of equipment.
- make sure that no unauthorised changes are carried out.

Hint

Removing the power cord results in complete disconnection from mains (all poles).

Caution

Additional equipment connected to medical electrical equipment must comply with the respective IEC or ISO standards (e.g. IEC 60950 for data processing equipment). Furthermore all configurations shall comply with the requirements for medical electrical systems (see IEC 60601-1-1 or clause 16 of the 3Ed. of IEC 60601-1, respectively).

Anybody connecting additional equipment to medical electrical equipment config ures a medical system and is therefore responsible that the system complies with the requirements for medical electrical systems. Attention is drawn to the fact that local laws take priority over the above mentioned requirements. If in doubt, consult your local representative or the technical service department.

- IEC 60601-1+AI +A2:1995: 6.8.2.c, 19.2.b, 19.2.c,
- IEC 60601-1:2005: 7.9.2.5, 8.1, 16.2.d,
- MDD 93/42/EEC: Annex I clause 13.6.c

SAFETY INFORMATION FOR Non-Invasive Blood Pressure Measurement

Warning

• Patient Hazard •

Do not take blood pressure measurements with a cuff on patients suffering from sickle cell anemia or where skin lesions are likely to occur.

The cuff may cause hematomas in patients with severe blood coagulation disease. In these instances, the user must take a decision for or against automatic blood pressure measurements.

Caution

• Compromised Measuring Accuracy •

Arrhythmias occurring frequently during a measurement may compromise the accuracy of the measurement. In certain cases, a valid measurement will not be possible.

Electromagnetic fields are also capable of impairing the measuring accuracy.

Intended Use

The ergoselect is a computer-controlled medical ergometer. At pedal speeds between 30 and 130 RPM and loads between 6 and 999 watt, the ergometer operates independent of the pedal speed.

The speed-independent range is shown in the Appendix (Technical Specifications).

The ergoselect ergometer may only be used in exercise testing as well as for rehabilitation of cardiac and cardiovascular patients according to the instructions given in this manual. If the ergometer is used for other purposes, the manufacturer cannot be held liable for personal injuries or property damage resulting from the unintended use of the equipment.

Note - Applied Parts

• Applied parts are components that are directly in contact with the human body (e.g., blood pressure measuring devices).

Note – Stability

• Ensure the stability of the ergometer. If the maximum permitted patient weight is exceeded by 10%, the stability of the ergometer can no longer be guaranteed. It may become unstable as a result.

Note

- If the cuff pressure exceeds the maximum value of 300 mmHg during inflation, the inflation procedure will be aborted and the cuff deflated.
 As a redundant safety precaution, the cuff is immediately deflated when the cuff pressure exceeds 320 mmHg.
 You can check the proper functioning of this safety precaution by abruptly bending your arm while the cuff is being inflated, causing a brief overpressure in the cuff. The cuff must deflate immediately.
- Measurements that did not yield a valid measurement will not be repeated during the exercise test.
- If the inflation phase takes longer than 40 seconds or if an adequate pressure does not build up in the cuff within a reasonable period of time, the measurement will be aborted and the cuff deflated.
- If a valid measurement cannot be completed within 120 seconds, the measurement will be aborted and the cuff deflated.
- If the cuff pressure remains constant for some time, the measurement will also be aborted and the cuff deflated.

BIOCOMPATIBILITY

The parts of the product described in this manual, including all accessories that come in contact with the patient during the intended use, fulfill the biocompatibility requirements of the applicable standards if applied as intended.

If you have questions in this matter, please contact ERGOLINE or a representative.

Applicable Laws, Regulations and Directives

- 93/42/EEC (Medical Device Directive of the EU)
- 89/336/EEC (Electromagnetic Compatibility Directive of the EU)
- EN 1060-1 Non-invasive sphygmomanometers, Part 1: General requirements
- EN 1060-3 Non-invasive sphygmomanometers, Part 3: Supplementary requirements for electro-mechanical blood pressure measuring systems

Symbols



Symbol 'type B applied part'.

Type B applied parts have no direct contact with patients and offer the lowest protection against electric shock.



Symbol 'type BF applied part'.

Type BF applied parts are connected to the body of the patient and provide a higher degree of protection against electric shock. The applied parts are isolated.



Caution, consult accompanying documents.



Protection class II equipment.



This symbol indicates that the waste of electrical and electronic equipment must not be disposed as unsorted municipal waste and must be collected separately.

Consult operating instructions.



Catalog number.



Serial number.



Scheduled date of the next inspection (e.g., March 2011).



Toggle switch ON (voltage)



Toggle switch OFF (voltage)



CE mark per the Medical Device Directive 93/42/EEC of the European Union. Notified body: TÜV SÜD Product Service GmbH, Ridlerstr. 65, 80339 München, Germany.



Nationally Recognized Testing Laboratory NRTL label for the USA and Canada.



Do not lean against the ergometer: there is a risk of the ergometer tipping over.



Manufacturer's identification.



Date of manufacture. The number found under this symbol is the year of manufacture in the YYYY format.



PVC-free.

Latex-free.



Suitable for indicated arm circumference.



Small size.



Standard size.





Large size.



Transport and storage label: top.



Transport and storage label: keep dry.



Transport and storage label: fragile.



Transport and storage label: approved temperature range.



Transport and storage label: approved humidity, non-condensing.



Transport and storage label: approved pressure range.



Transport and storage label: do not stack.

Setup and Mains Connection

Controls and Indicators

- 1 Control terminal (model P or model K)
- 2 Blood pressure cuff connection (option)
- 3 Adjustment of the handlebar angle
- 4 Blood pressure cuff
- 5 Adjustment of the handlebar height (ergoselect 150/200 only)
- 6 Castors
- 7 Speed display (RPM) for patient
- 8 Adjustment of the saddle height (ergoselect 100/150 only)
- 9 Digital saddle height indication (ergoselect 200 only)
- 10 Power switch (green button)
- 11 Sockets for power cord and connection cables (underside of ergometer)
- 12 Levelling devices to adjust the ergometer to uneven floors



ERGOSELECT 100/150 - CONTROLS, CONNECTIONS AND INDICATORS



ERGOSELECT 200 - CONTROLS, CONNECTIONS AND INDICATORS

TRANSPORT

For short distances, the ergoselect can be lifted at the saddle and rolled away on its castors.

To cover greater distances, however, we recommend the following method:

Caution

• Equipment Damage •

Avoid strong vibrations of the ergoselect during transport.

- Disconnect the power cord from the wall outlet.
- Rotate the handlebar towards the front. Tighten the clamping lever.
- Stand in front of the ergoselect, grasp the handlebar and tilt the ergoselect towards you until it is standing on the castors only and is balanced.
- It is now possible to transport the ergoselect.
- When you have reached the new location, lower the ergoselect very carefully to avoid damage.



TRANSPORTING THE ERGOSELECT

SETUP

Place the ergoselect on a horizontal level floor.

The ergoselect must be set up in a secure and stable position - the two levelling feet at the back make for easy adjustment to uneven floor surfaces. Extend the foot concerned until the ergoselect no longer wobbles.

In case of delicate flooring, it is recommended to place a mat under the ergometer to protect the flooring from damage by the feet.



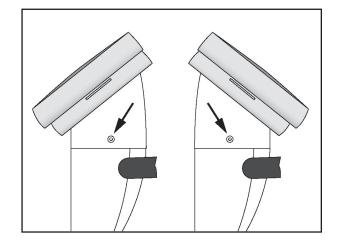


LEVELLING FEET OF THE ERGOSELECT ERGOMETER

MOUNTING THE CONTROL TERMINAL (P OR K)

The control terminal can be installed with the display either facing the patient or the operator.

It is recommended to install the terminal with the display and control keys towards the operator and the speed display towards the patient.



DIFFERENT ORIENTATIONS OF THE CONTROL TERMINAL

CONNECTING THE POWER CORD

Set the handlebar to the front upper position and secure. Tilt the ergoselect carefully towards you until it rests on the handlebar.

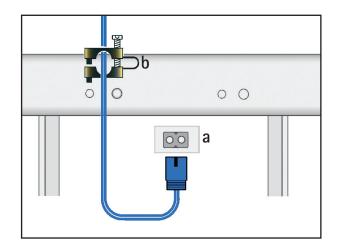


ASSEMBLY POSITION OF THE ERGOSELECT ERGOMETER



Before connecting the ergometer to the power line, check that the line voltage corresponds to the ratings on the type plate. The type plate is located on the back of the ergometer, at the bottom. The connection panel is located on the underside of the ergometer.

- Plug the power cord into socket (a) and use the supplied lock (b) to secure it against disconnection.
- Using the supplied strain relief, attach the cable to the metal frame.



CONNECTION PANEL

a Power input

b Lock



POWER CORD WITH INSTALLED STRAIN RELIEF

Caution

• Disconnection from Power Supply •

Pressing the power switch or removing the power cord disconnects the device from the power supply.

Removing the power cord results in a complete disconnection from the power supply (all poles).

Ensure that the power plug is readily accessible at all times.

CONNECTING THE ECG CABLE

ergoselect ergometers can be connected to electrocardiographs and PC-based ECG systems of most manufacturers.

Different connection cables are available to support different communication modes (digital, analog, remote start, etc.).

All ergoline ergometers are equipped with a digital interface (special adapters, which can be obtained from ergoline, are required for control of the ergometer with analog signals or for the remote start function).

The appropriate cable is plugged into the 9-pole port of the connection panel (Port 1) or into the USB port and secured at the metal frame with an additional strain relief.



EKG / PC CONNECTION

USB PORT 1 PC connection via USB (virtual COM) Digital connection (remote control from PC or ECG recorder), connection for cable adapter (analog interface + remote start)

Hint

• connecting cables •

Only use connecting cables released by ergoline.

To use the integrated USB connector, a special driver is required - contact ergoline.

CONNECTING THE BLOOD PRESSURE CUFF

- Connect the microphone at (1).
- Slip the cuff tubing onto the fitting (2) and engage. To disconnect, push back the connector's knurled sleeve.

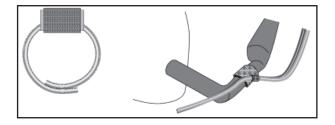
Artifacts that may be caused by patient movements during the exercise test, must be avoided if possible, while the blood pressure is being taken.

Therefore, do not forget to attach the cuff tubing to the handlebar with the supplied Velcro tape:

1

BLOOD PRESSURE CUFF CONNECTIONS

- 1 Microphone connection
- 2 Cuff tubing
- Open the large Velcro tape and wrap around handlebar.
- Secure the cuff tubing with the small Velcro tape, but do not exert pressure on the tubing.



VELCRO TAPE TO SECURE THE CUFF TUBING

PREPARING THE PATIENT

Adjusting the Saddle and the Handlebar

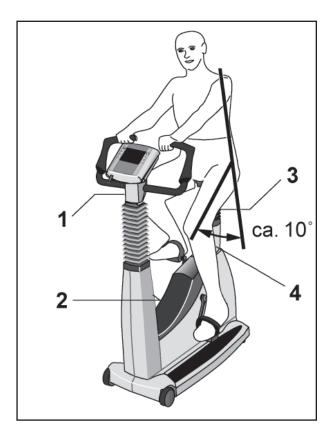
On the ergoselect 100/150, you adjust the saddle height manually with a clamping lever. On the ergoselect 200, the saddle height is electrically adjusted with the corresponding keys on the control terminal (the display below the saddle indicates the saddle height).

With the pedal in the bottom position, the angle between the axis formed by the upper body and the thigh should be approximately 10°.

Set the handlebar to a position where the patient sitting upright on the saddle can reach it easily.

To do so, open clamping lever 1 and set the handlebar to a suitable angle.

On the ergoselect 150/200, the height of the handlebar can additionally be adjusted with clamping lever 2 – the horizontal bar should be approximately at the samel level as the saddle.



Adjusting saddle and handlebar

- 1 Adjustment of the handlebar angle
- 2 Adjustment of the handlebar height (ergoselect 150/200 only)
- 3 Adjustment of the saddle height (ergoselect 100/150 only)
- 4 Saddle height display (ergoselect 200 only)

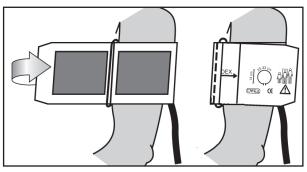
Note

- Tighten the clamping levers only as far as necessary, NOT with maximum force.
- Lubricate the clamping lever threads quarterly at minimum, using a suitable lubricant (e.g. OKS470).

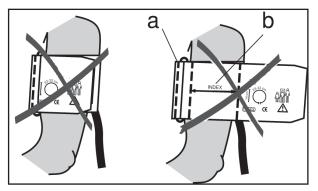
Preparing the Patient for Blood Pressure Measurements

CUFF SIZE

Always choose the cuff size suitable for the patient's arm. The maximum arm circumference is indicated on the cuff.



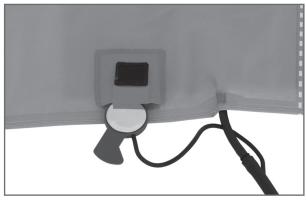
CORRECT CUFF SIZE



WRONG CUFF SIZE

MICROPHONE POSITION

Before applying the cuff, check the position of the microphone inside the red pocket (on the inside of the cuff): When the microphone is inside the pocket, its **metal side must face the arm**.



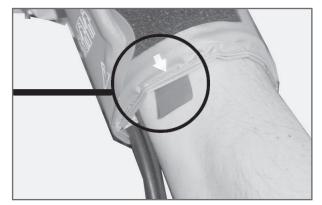
CORRECT MICROPHONE POSITION

APPLYING THE CUFF

The center of the microphone must be located exactly on the **brachial artery**. Locate the artery by palpation, if required. The **red tab** identifies the position of the microphone.

The accurate placement of the microphone is the primary condition for reliable pressure measurement during exercise tests.

The cuff must be applied directly on the skin, it may not be applied on top of clothing, paper, etc. Apply the cuff approx. **2 cm above the bend of** the elbow. The cuff should be **tight**, but it should not constrict blood vessels. The cuff **may not move** during the exercise test.



MICROPHONE PLACEMENT ON THE ARTERY

When you close the Velcro strap, check that the metal clasp (a) is inside the marked index range (b), and not outside.

The cuff tab must be located below the metal clasp (see illustration at right).

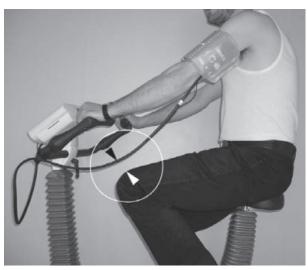
CORRECT CUFF POSITION (TAB)

CHECKING THE CUFF TUBING

Check that the cuff tubing does not knock against the patient's knee, when the patient is pedalling and the hand is on the handlebar.

Secure the cuff tubing with the Velcro tape attached to the handlebar.

Instruct your patient to move as little as possible during a blood pressure measurement and, in particular, to avoid excessive contractions of the muscles in the upper arm.



DISTANCE BETWEEN KNEE AND TUBING

Caution Patient Hazard

Apply the cuff directly on the skin. Make sure that rolled up sleeves do not impede blood circulation in the upper arm. Loose cuffs will cause erroneous measurements; overtight cuffs may constrict blood vessels or cause skin lesions and hematomas.

• Incorrect Measurements •

A loose cuff would degrade the accuracy of the measurement. Therefore, the computer aborts the measurement, if a minimum pressure is not attained within a few seconds.

Warning Patient Hazard

If, by accident, an excessive pressure builds up inside the cuff, either remove the cuff immediately from the arm or disconnect the cuff tubing from the control terminal. The same measures are recommended, if the cuff does not deflate correctly.

OPERATION

The ergometers of the ergoselect series are available with two versions of the control terminal whose functionalities differ.

The following sections describe the control and configuration of the ergometer.





Control terminal P

Control terminal K

CONTROL TERMINAL P

TURNING THE SYSTEM ON

You turn the ergometer on by pressing the power switch – the green indicator in the switch lights up. The ergometer runs a self-test. Subsequently, the main menu displays.

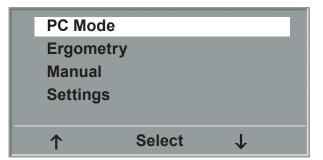
Note

- Instruct the patient not to pedal while the ergometer is being turned on and during the self-test.
- Apply the blood pressure cuff to the patient AFTER the ergometer has been turned on and the self-test completed.
- The device can be configured to default to one of the operating modes. If this option is selected, the initial screen of the selected operating mode (e.g. Ergometry) will be displayed instead of the main menu. With the Lange key, you can display the main menu.

ergoline GmbH

Selftest running

SELF-TEST SCREEN



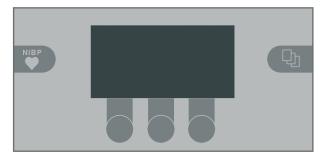
MAIN MENU

The ergometer software is controlled with 5 keys:

With this key you display the main menu or return to the previous menu level.

With this key you initiate a blood pressure measurement. A measurement in progress can be aborted with the same key.

The functions of these three softkeys change with the displayed menu - the key label describing the function is shown on the display.





OPERATING MODES WITH CONTROL TERMINAL P

An ergoselect ergometer with a control terminal P supports the following operating modes:

PC MODE

An external device (e.g. stand-alone electrocardiograph, PC-based ECG system) controls the ergometer – no intervention at all is required at the ergometer.

ERGOMETRY

The ergometer runs an automatic exercise test - some of the corresponding test protocols are user-configurable and stored in the system (see chapter "Settings").

MANUAL

The ergometer is controlled manually, i.e., the user performs all load changes via the keypad.

SETTINGS

Used to configure the ergometer.

Speed readout

At the top of the control terminal, there is a speed readout for the patient as well as three LEDs that inform the patient of the speed: too slow, too fast or correct.

The ranges for the respective speed ratings depend on the selected load (see "Technical Specifications").



SPEED READOUT

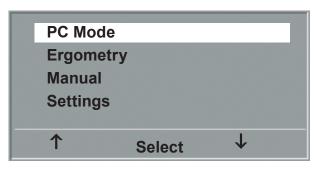
- 1 speed low (patient should pedal faster)
- 2 correct speed
- *3* speed high (= patient should pedal slower)

Note

- If, during an exercise test, the speed drops below 30 RPM, the load readout starts blinking on the display.
- To reactivate the saddle height adjustment function, press and the arrow keys will again be displayed.
- Additional blood pressure measurements an be initiated with with .

PC MODE

Use the softkeys on the right and left ($\uparrow \downarrow$) to position the bar cursor on PC MODE and confirm the selection with SELECT.



Main menu

The display changes - the ergometer is waiting for commands from the external ECG unit.

With the arrrow keys, the saddle height can be electrically adjusted on the ergoselect 200 (on the ergoselect 400, these keys adjust the height of the drive unit).

As soon as the ergometer receives commands from the controlling ECG unit or PC, the exercise test will start and

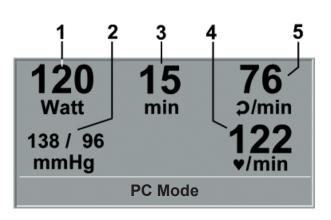
The exercise test can only be terminated with the corre-

sponding command from the controlling ECG unit.

the corresponding values will be displayed.

0-0Wattmin⊅/min--- / ---0mmHg✓/min↑Saddle

INITIAL SCREEN



DISPLAY DURING EXERCISE TEST

- 1 current load in watts
- 2 most recent BP value (systolic/diastolic values) or cuff pressure during inflation and bar graph indicating microphone signal strength (see below)
- 3 duration of exercise test (min)
- 4 heart rate at the time of the BP measurement (BPM)
- 5 pedal speed (RPM)



Note

- All functions are locked while the ergometer is operating in PC mode, except for the saddle height adjustment and the blood pressure key.
- Additional blood pressure measurements can be initiated with

Ergometry

Use the softkeys on the right and left ($\uparrow\downarrow$) to position the bar cursor on ERGOMETRY and confirm the selection with SELECT.

PC Mode		
Ergometry		
Manual		
Settings		
1	Select	\downarrow

Main menu

The stored test protocols available for selection will be displayed. There are five fixed protocols (protocols 1 to 5, see Appendix), whereas protocols 6 to 15 are user-programmable.

The protocol menu provides an overview of the test phases:

e.g.:	50 W /	2 min /	25 W
-------	--------	---------	------

means: initial (basic) load 50 watts stage time 2 minutes load increment 25 watts

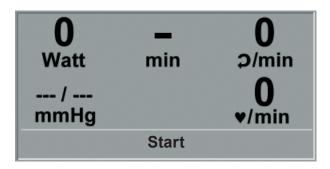
Use the softkeys on the right and left ($\uparrow \downarrow$) to position the bar cursor on one of the protocols and confirm the selection with SELECT.

The exercise test is started with the "Start" key, a blood pressure measurement at rest may precede the test (see "Settings").

When the basic load appears on the display (after approx. 15 seconds or upon termination of the blood pressure measurement) and the patient's RPM indicator blinks, the patient should start pedalling.

Pro	otocols	
1.	WHO	
2.	BAL	
3.	Hollmann	
4.	STD. France	
5.	Standard	
1	Select	\checkmark

SELECTING AN EXERCISE TEST PROTOCOL



INITIAL EXERCISE TEST SCREEN

The internal protocol will now control the entire exercise test - the display always indicates the current values.

With the +5 W and -5 W keys, the current load can be changed at any time (in increments of +/-1 W up to +/-25 W, as configured).

120 Watt 138 / 96 mmHg	15 min	76 D/min 122 Min
+ 5 W	Recovery	- 5 W

SCREEN DISPLAY DURING THE TEST

Note

- The saddle height (ergoselect 200) can be changed during an exercise test.
- To reactivate the saddle height adjustment function, press
- Additional blood pressure measurements can be initiated with NBP.

TERMINATING AN EXERCISE TEST

The exercise phase can be terminated manually at any time with the RECOVERY key.

The load will immediately be reduced to 25 watts, but a higher or lower value can be selected manually.

It is recommended that the patient continues to pedal in the recovery phase.

The END key in the middle will terminate the test.

120 Watt 138 / 96 mmHg	15 min	76 p/min 122 v/min
+ 5 W	End	- 5 W

RECOVERY PHASE

MANUAL

Use the softkeys on the right and left ($\uparrow\downarrow$) to position the bar cursor on MANUAL and confirm the selection with SELECT.

In this operating mode the user controls the entire exercise test by selecting the loads, stage times and by initiating blood pressure measurements.

PC Mod	de		
Ergom	etry		
Manua	l		
Setting	IS		
1	Select	\downarrow	

Main menu

The exercise test is started with the "Start" key, afterwards the load can be set and changed with the +5 W and -5 W keys (in increments of +/-1 W up to +/-25 W, as configured).

Blood pressure measurements an be initiated with .

O Watt	_ min	0 ⊋/min
/ mmHg		0 ▼/min
+ 5 W	Start	- 5 W

INITIAL SCREEN OF A MANUAL EXERCISE TEST

TERMINATING AN EXERCISE TEST

The exercise test can be terminated manually at any time with the END key located in the middle.

The load will immediately drop to 0 watt.

There is no recovery phase in the manual mode.

120 Watt 138 / 96 mmHg	15	76 D/min 122 V/min
+ 5 W	End	- 5 W

SCREEN DISPLAY DURING THE TEST

SETTINGS WITH CONTROL TERMINAL P

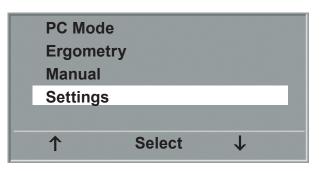
Some of the device settings are configurable to meet specific requirements. The settings will be saved and remain stored even when the ergometer is switched off.

Use the softkeys on the right and left ($\uparrow\downarrow$) to position the bar cursor on SETTINGS and confirm the selection with SELECT.

The configuration menu displays.

When all changes have been made, you can exit the configuration menu with the $[P_2]$ key.

Use the softkeys on the right and left ($\uparrow \downarrow$) to position the bar cursor on the parameter to change and confirm the selection with SELECT.



Main menu

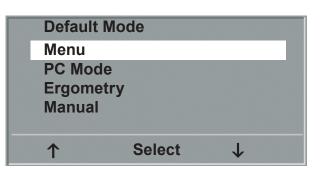
Settings	6		
Default	Mode		
Protoco	ls		
Contras	t		
Load Ch	nange		
Langua	ge		
1	Select	\downarrow	

CONFIGURATION MENU

DEFAULT MODE

In this menu you choose the default mode activated when the ergometer is turned on. When first turned on, the ergometer will display this menu.

Use the softkeys on the right and left ($\uparrow \downarrow$) to position the bar cursor on your preferred default mode and save the selection with SELECT.



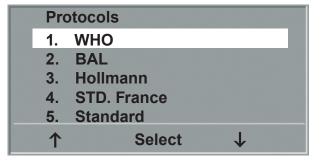
SELECTING THE DEFAULT MODE

PROTOCOLS

Protocols 6 - 15 are user-programmable (protocols 1 - 5 are fixed, see Appendix for protocol parameter details). Standard values for the following parameters can be entered:

- protocol type (step or ramp)
- initial load
- stage time
- load increment (load increase with each stage)

Use the softkeys on the right and left ($\uparrow \downarrow$) to position the bar cursor on the protocol to change (No. 6 - 15) and confirm the selection with SELECT.



SELECTING THE EXERCISE TEST PROTOCOL TO EDIT

Use the softkeys $\uparrow \downarrow$ to select the parameter to edit.

At Select, for example, you can choose the protocol type:

- Step (load increase in steps) or

- Ramp (continuous load increase).

Press SELECT to save the selected protocol type.

To cancel the selection, press the Q_1 key.

Proto	col	6.
Selec	t	Step
Basic	Load	25 W
Stage	Time	2 min
Load	Stage	25 W
1	Select	\checkmark

SELECTING THE PARAMETER TO EDIT

All other parameters are edited in the same way.

Using the arrow keys ($\uparrow \downarrow$), highlight a parameter and confirm the selection with SELECT: the corresponding value appears in reverse video and can be changed with the arrow keys $\uparrow \downarrow$.

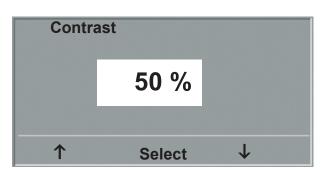
Pressing SELECT will save the new value. You exit the configuration with \bigcirc .

Protocol		6.	
Select		Step	
Basic Lo	ad	25 W	
Stage Tir	ne	2 min	
Load Sta	ge	25 W	
↑	Select	\checkmark	

EDITING THE PARAMETER VALUE

CONTRAST

The display contrast is adjustable in the range from 0 to 100%.



Adjusting the display contrast

LOAD CHANGE

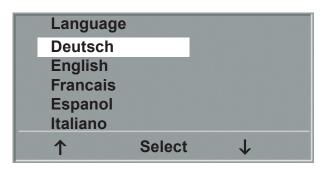
Here you determine the increments for each load change. Depending on your choice, each key press will change the load by +/-1, 5, 10 und 25 Watts.

Load C	Change	
+/	1 Watt	
+/	5 Watt	
+/	10 Watt	
+/	25 Watt	
-		
1	Select	\checkmark

SELECTING THE INCREMENT FOR MANUAL LOAD CHANGES

LANGUAGE

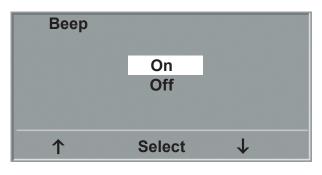
The texts can be displayed in different languages.



LANGUAGE MENU

BEEP

The audio signal emitted during blood pressure measurements can be turned on and off.



BEEP DURING BP MEASUREMENTS

SOFTWARE VERSION

Select this option to view the installed software version.

DATE/TIME

To begin with, you select DATE or TIME and confirm the selection. Then the value displayed in reverse video can be edited with the $\uparrow \downarrow$ keys and saved with SELECT.

The time is adjusted in the same way. You exit the configuration with

Date		
	22. 08. 2016	
Time	17:33:05	
	17:33:05	
1	Select	\checkmark

SETTING THE DATE

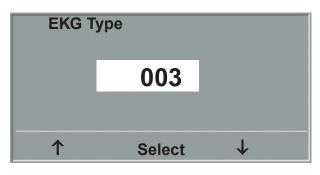
Date	22. 08. 2016	
Time	17:33:05	
1	Select	\downarrow

Setting the day

EKG TYPE

The selected EKG Type determines the communication method with the ECG recorder, PC-based ECG system, etc.

To prevent an accidental change of this setting, the menu is protected with a password. Using the arrow keys, enter 003 and confirm the entry with SELECT.





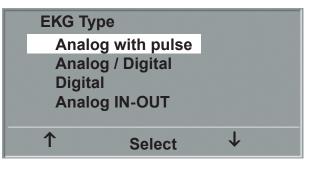
All ergoselect ergometers support the following communication modes:

- Analog with pulse Remote start mode; prior the each load change, the ergometer generates a control pulse and sends the corresponding data via the interface.
- Analog / Digital
 An analog voltage controls the load blood pressure measurements can be initiated with digital commands.
- Digital (default) The communication with the ergometer is entirely controlled with digital commands.
- Analog IN-OUT The entire communication (load control and BP measurements) is controlled with analog signals. No digital data will be sent.

Select the communication mode and confirm with SELECT.

Note

- The EKG Type needs to be selected only when the ergometer is connected to an ECG unit. The selection is part of the installation procedure.
- The "Analog/Digital" and "Digital" communication is only possible when PC Mode is selected from the main menu or when this is the default mode.



Selecting the ergometer communication mode

RPM

Here you determine the RPM limits. When these limits are exceeded, the LEDs for high or low speed (RPM) will illuminate.

Select the value to change (Min. or Max.) and confirm with SELECT.

Using the arrow keys, change the value and save the new value with SELECT.

RPM		
Min ↑	0 70	
	54	
Max↓	50 130	
	64	
1	Select	\downarrow

SETTING THE RPM LIMIT VALUES

Note

• The limits selected in this menu only apply to the load range between 6 and 150 watts. At higher loads the RPM limits automatically adapt to the respective loads:

Load (watts)	Green RPM range (1/min)
6 - 150	54 – 64 (adjustable)
151 - 250	58 - 65
251 - 350	68 - 75
351 - 450	78 - 85
451 - 550	88 - 95
551 - 650	98 - 105
651 - 750	108 - 115
751 - 850	118 - 125
851 - 950	> 125
951 - 999	> 130

PULSE DISPLAY

The pulse readout on the display can be turned off.

CONTROL TERMINAL K

TURNING THE SYSTEM ON

You turn the ergometer on by pressing the power switch the green indicator in the switch lights up. The ergometer runs a self-test. Subsequently, the main menu displays.

ergoline GmbH

Selftest running

SELF-TEST SCREEN

Note

- Instruct the patient not to pedal while the ergometer is being turned on and during the self-test.
- Apply the blood pressure cuff to the patient AFTER the ergometer has been turned on and the self-test completed.
- The device can be configured to default to one of the operating modes. If this option is selected, the initial screen of the selected

operating mode (e.g. Ergometry) will be displayed instead of the main menu. With the the key, you can display the main menu.

Exercise Test	PC Mode
Training	Manual
Test	Settings

MAIN MENU

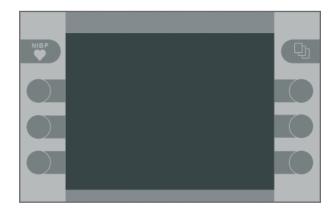
The ergometer software is controlled with 8 keys:

With this key you display the main menu or return to the previous menu level.

With this key you initiate a blood pressure measurement. A measurement in progress can be aborted with the same key.

\bigcirc	
\bigcirc	
\bigcirc	

The functions of these six softkeys change with the displayed menu - the key label describing the function is shown on the display.



Keypad K

OPERATING MODES WITH CONTROL TERMINAL K

An ergoselect ergometer with a control terminal K supports the following operating modes:

PC MODE

An external device (e.g. stand-alone electrocardiograph, PC-based ECG system) controls the ergometer – no intervention at all is required at the ergometer.

ERGOMETRY

The ergometer runs an automatic exercise test - some of the corresponding test protocols are user-configurable and stored in the system (see chapter "Settings").

TRAINING

Ten different training protocols with warm-up, exercise and recovery phases can be custom-configured (see chapter "Settings").

A POLAR receiver is integrated in the ergometer and provides the relevant data for heart-rate controlled training sessions.

TEST

Integrated test protocols (steep ramping test, PWC tests) allow an assessment of the physical fitness.

MANUAL

The ergometer is controlled manually, i.e., the user performs all load changes via the keypad.

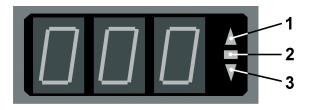
SETTINGS

Used to configure the ergometer.

Speed readout

At the top of the control terminal, there is a speed readout for the patient as well as three LEDs that inform the patient of the speed: too slow, too fast or correct.

The ranges for the respective speed ratings depend on the selected load (see "Technical Specifications").



SPEED READOUT

- 1 speed low (patient should pedal faster)
- 2 correct speed
- 3 speed high (= patient should pedal slower)

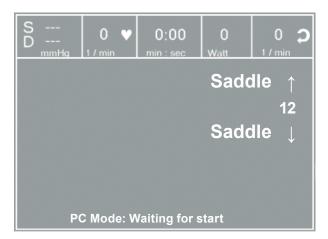
Note

- If, during an exercise test, the speed drops below 30 RPM, the load readout starts blinking on the display.
- Additional blood pressure measurements can be initiated with

PC MODE

When the PC Mode key has been pressed, the screen appears as shown at right. The ergometer is waiting for commands from the external ECG unit.

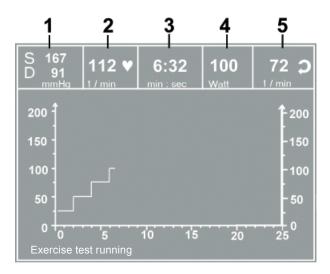
With the arrrow keys, the saddle height can be electrically adjusted on the ergoselect 200 (on the ergoselect 400, these key adjust the height of the drive unit).





As soon as the ergometer receives commands from the controlling ECG unit or PC, the exercise test will start and the corresponding values will be displayed.

The exercise test can only be terminated with the corresponding command from the controlling ECG unit.



DISPLAY DURING EXERCISE TEST

- 1 most recent BP value (systolic/diastolic pressures) or cuff pressure during inflation and bar graph indicating microphone signal strength (see below)
- 2 Sp02 (%), heart rate (BPM)
- *3 duration of exercise test (minutes:seconds)*
- 4 current load in watts
- 5 pedal speed (RPM)



Note

- All functions are locked while the ergometer is operating in PC mode, except for the saddle height adjustment and the blood pressure key.
- Additional blood pressure measurements can be initiated with with .

Ergometry

The ergometer is controlled by an internally stored protocol.

Pressing the "Ergometry" key will display the test protocol used last.

Press the "Start" key to re-start the protocol, or press the "Select" key to display the protocol parameters or to switch to another test protocol.

There are five fixed protocols (protocols 1 – 5, see Appendix), whereas protocols 6 – 15 are user-programmable.

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INITIAL SCREEN OF AN EXERCISE TEST

With the arrow keys you can display the test protocol. With "Select" you confirm the selection.

The selected exercise test is started with the "Start" key, a blood pressure measurement at rest may precede the test (see "Settings").

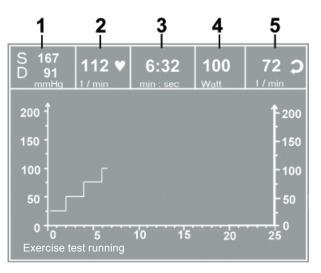
The display changes to the exercise test screen, where load and heart rate are represented both by numeric values and waveforms.

When the basic load appears on the display (after approx. 15 seconds or upon termination of the blood pressure measurement) and the patient's RPM indicator blinks, the patient should start pedalling.

The internal protocol will now control the entire exercise test - the display always indicates the current values.

S D mmHg 1 / min	• 0:00 min : sec	0 Watt	0 2
Protocol WHC)		*
Basic Load	25 Watt		
Stage Time	2 min		
Load Stage	25 Watt		
Recovery Load	25 Watt		*
Recovery Time	10 min		
NIBP Lead Time	60 sec		
		્	Select
S	elect protoco	1	





DISPLAY DURING EXERCISE TEST

- 1 most recent BP value (systolic/diastolic pressures) or cuff pressure during inflation
- 2 heart rate (BPM)
- 3 duration of exercise test (minutes:seconds)
- 4 current load in watts
- 5 pedal speed (RPM)

Adjustments During the Exercise Test

Press the D key to display the configuration menu. This is what you can do during the test

- increase or decrease the current load in increments (adjustable between 1 watt and 25 watts)
- hold the current load
- end the exercise phase and advance to the recovery phase

Pressing 😳 again displays another menu where you can

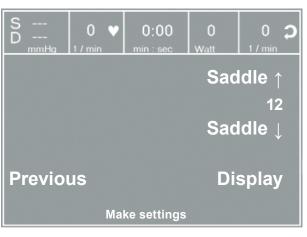
change the saddle height and the display mode (see "PC

• terminate the test.

Mode").

S D mmHg 1 / min	0:00 min : sec	0 Watt	0 🎝 1 / min
Load +	Reco	very p	hase
Load –			End
Load const.		Pre	vious
Make settings			

CONFIGURATION MENU I



CONFIGURATION MENU II

TERMINATING THE TEST

Once the full protocol has been completed, the test will be terminated.

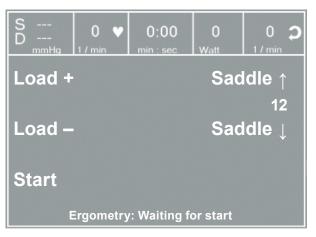
However, it is possible at any time to manually terminate the test or switch to the recovery phase (see above).

MANUAL

In this operating mode the user controls the entire exercise test by selecting the loads, stage times and by initiating blood pressure measurements.

The exercise test is started with the "Start" key, afterwards the load can be set and changed with the [Load +] and [Load -] keys (in increments of 1 W up to 25 W, as configured).

Blood pressure measurements can be initiated with **PP**.



SCREEN DISPLAY IN MANUAL MODE

TERMINATING AN EXERCISE TEST

The exercise test can be terminated manually at any time with the END key located in the middle.

The load will immediately drop to 0 watt.

There is no recovery phase in the manual mode.

TRAINING

Cardiologic training sessions can be performed with ergoselect ergometers equipped with control terminal K. For a detailed description of the protocols, please refer to the Appendix.

Pressing the "Training" key will display the training protocol used last.

Press the "Start" key to re-start the protocol, or press the "Select" key to display the protocol parameters or to switch to another training protocol.

All training protocols 1 - 10 are user-configurable (see "Settings for Control Terminals K").

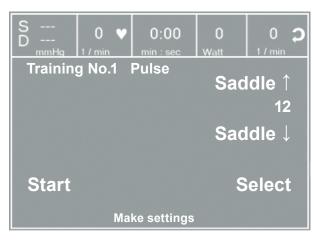
Use the arrow keys to display the protocol to use and the corresponding parameters. Confirm the selection with the "Select" key.

You initiate the training session with the "Start" key.

The display changes to the training session screen, where load and heart rate are represented both by numeric values and by waveforms.

When the basic load appears on the display (after approx. 15 seconds or upon termination of the blood pressure measurement) and the patient's RPM indicator blinks, the patient should start pedalling.

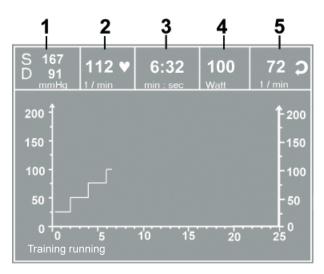
The internal protocol will now control the entire training session - the display always indicates the current values.



INITIAL SCREEN OF THE TRAINING SESSION

S 0 D 0 1 / min	♥ 0:00 min : sec	0 Watt	0 🎝
Training No.	1 Pulse	mau	
Basic Load	25 Watt		\uparrow
Warmup	2 min		
Training time	20 min		
Recovery Load	20 Watt		
Recovery Time	3 min		*
Load increment	8 Watt/min		
Training pulse	100 P/min		
Maximum load	80 Watt	Se	lect
	Select protoco		





DISPLAY DURING EXERCISE TEST

- 1 most recent BP value (systolic/diastolic pressures) or cuff pressure during inflation
- 2 heart rate (BPM)
- 3 duration of exercise test (minutes:seconds)
- 4 current load in watts
- 5 pedal speed (RPM)

Adjustments During the Training Session

Press the 🕒 key to display the configuration menu. This is what you can do during the training session

- end the training session and advance to the recovery phase,
- directly terminate the training session,
- change the display mode (see "PC Mode").



CONFIGURATION MENU

TRAINING WITH CHIP CARD

As an alternative to the training protocols saved in the ergometer, it is possible to load training protocols from the chip card.

The training protocols are saved to the chip card by means of a PC program ("ergoline opticare professional" or "ergoline opticare basic").

Upon completion of the training session, the entire procedure (incl. load and heart rate waveforms) is saved to the chip card and can be reviewed and analyzed at the PC.

STARTING THE CHIP CARD TRAINING SESSION

Select the "Training" mode and insert the chip card into the card reader (on the side of the control terminal).

The ergometer switches to the chip card mode and reads the data stored on the card.



READING THE CHIP CARD DATA

The name and the weight stored on the card are displayed.

You can use the arrow keys to enter the current weight.

Press the "Next" key and the initial screen will display. You can initiate the displayed training protocol or select another protocol from the chip card.

The chip card training session proceeds in the same way as the exercise tests stored in the ergometer.

ENTERING THE WEIGHT

TERMINATING THE TRAINING SESSION

After termination of the training session (automatic termination when the programmed recovery phase has been completed, or manual termination) the test subject can state how the test was perceived (BORG scale).

Training Chipcard	
Stress very very heavy	^
very heavy	
heavy	
a little stronger	\checkmark
easy very easy	
very very easy	ОК

ENTERING THE BORG VALUE

Subsequently all training data are written to the chip card and are then available for analysis with a special program (e.g. opticare basic).



WRITING TO THE CHIP CARD

SETTINGS FOR CONTROL TERMINALS K

Some of the device settings are configurable to meet specific requirements. The settings will be saved and remain stored even when the ergometer is switched off.

Select SETTINGS to display the configuration menu.

When all changes have been made, you can exit the configuration menu with the the key.

Use the softkeys ($\uparrow \downarrow$) to position the bar cursor on the parameter to change and confirm the selection with SELECT.

Settings	
Default Mode	
Protocols	↑
Contrast	
Load Change	
Language	
Веер	¥
Software Version	
Date/Time	
Training	Select
EKG Type	
Choose fu	inction

CONFIGURATION MENU

DEFAULT MODE

In this menu you choose the default mode activated when the ergometer is turned on. When first turned on, the ergometer will display this menu.

Use the softkeys ($\uparrow \downarrow$) to position the bar cursor on your preferred default mode and save the selection with SELECT.

PROTOCOLS

Protocols 6 - 15 are user-programmable (protocols 1 - 5 are fixed, see Appendix for protocol parameter details). Standard values for the following parameters can be entered:

- test protocol type (step/ramp)
- initial load
- stage time
- load increment (load increase with each stage)
- NIBP lead time (blood pressure measurement)
- recovery load
- recovery time

Use the softkeys (\uparrow \downarrow) to position the bar cursor on the protocol to change (No. 6 – 15) and confirm the selection with SELECT.

Pro	tocols	
1.	WHO	
2.	BAL	
3.	Hollmann	\uparrow
4.	STD. France	
5.	Standard	
6.	25W / 2min / 25W	Ļ
7.	25W / 2min / 25W	
8.	25W / 2min / 25W	
9.	25W / 2min / 25W	Select
10.	25W / 2min / 25W	
	Choose function	

SELECTING THE EXERCISE TEST PROTOCOL TO EDIT

Use the softkeys $\uparrow \downarrow$ to select the parameter to edit.

At Select, for example, you can choose the protocol type:

- Step (load increase in steps) or

- Ramp (continuous load increase).

Press SELECT to save the selected protocol type.

To cancel the selection, press the 🕒 key.

Protocol Select		6 Step	
Basic Load	25	Watt	
Stage Time	2	min	\uparrow
Load Stage	25	Watt	
NIBP Lead Time	60	sec	
Recovery Load	25	Watt	•
Recovery Time	2	min	Select
Choc	ose fu	Inction	

SELECTING THE PARAMETER TO EDIT

All other parameters are edited in the same way.

Using the arrow keys ($\uparrow \downarrow$), highlight a parameter and confirm the selection with SELECT: the corresponding value appears in reverse video and can be changed with the arrow keys $\uparrow \downarrow$.

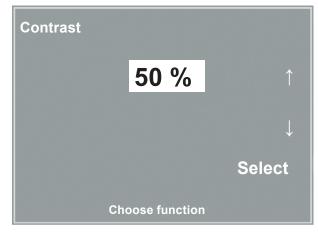
Pressing SELECT will save the new value. You exit the configuration with \Box_2 .

Protocol		E	5.
Select		Step	
Basic Load	25	Watt	
Stage Time	2	min	\uparrow
Load Stage	25	Watt	
NIBP Lead Time	60	sec	
Recovery Load	25	Watt	
Recovery Time	2	min	Select
Choo	se fu	Inction	

EDITING THE PARAMETER VALUE

CONTRAST

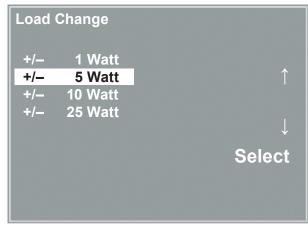
The display contrast is adjustable in the range from 0 to 100%.



EDITING THE PARAMETER VALUE

LOAD CHANGE

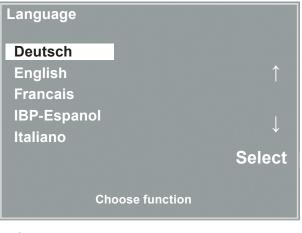
Here you determine the increments for each load change. Depending on your choice, each key press will change the load by +/-1, 5, 10 und 25 watts.



SELECTING THE INCREMENT FOR MANUAL LOAD CHANGES

LANGUAGE

The texts can be displayed in different languages.



LANGUAGE MENU

BEEP

The audio signal emitted during blood pressure measurements can be turned on and off.

SOFTWARE VERSION

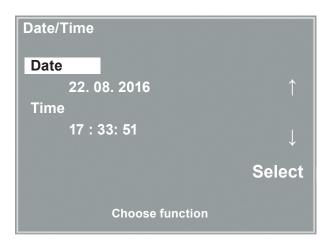
Select this option to view the ergometer's installed software version.

DATE/TIME

To begin with, you select DATE or TIME and confirm the selection.

Then the value displayed in reverse video can be edited with the $\uparrow\downarrow$ keys and saved with SELECT.

The time is set in the same way. You exit the configuration with the same way.





Date/Time	
Date	
22. 08. 2016 Time	1
17 : 33: 51	
	\downarrow
	Select
Choose function	

CHANGING THE DATE

TRAINING

Ten training protocols consisting of warmup, training and recovery phase are user-configurable. Depending on the selected training mode (pulse, constant, interval), there will be different parameters to define for the training phase: First of all you select and confirm the protocol you wish to configure.

Then you select the parameters with the arrow keys ($\uparrow\downarrow$) as usual and edit them.

Training	
1. Pulse	
2. Constant	
3. Interval	1
4. Interval	
5. Pulse	
6. Pulse	\downarrow
7. Pulse	
8. Pulse	
9. Pulse	Select
10. Pulse	
Cł	oose function

SELECTING THE EXERCISE TEST PROTOCOL TO EDIT

For all training modes (pulse, constant load and interval), the warmup phase, the duration of the training session and the recovery phase are defined first.

Depending on the selected training mode, you can edit the corresponding parameters afterwards:

10 - 300

sec

•	Pulse-controlled t Training pulse: Maximum load:	raining: 40 - 250 1 - 999	1/min Watt
•	Constant load: Training load:	1 - 999	Watt
•	Interval training: Load Stage 1: Stage Time 1: Load Stage 2:	1 - 999 10 - 300 1 - 999	Watt sec Watt

Stage Time 2:

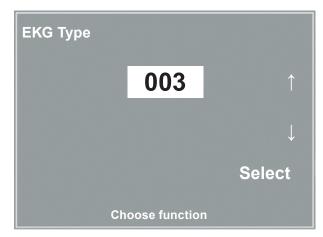
Training Select	Puls	se	
Basic Load	20	Watt	
Warmup	2	min	↑
Training time	20	min	
Recovery Load	20	Watt	
Recovery Time	3	min	\downarrow
Load increment	8	W/min	
Training pulse	100	1/min	
Maximum load	50	Watt	Select
Cho	ose fui	nction	

EDITING THE TRAINING PROTOCOL

EKG TYPE

The selected EKG Type determines the communication method with the ECG recorder, PC-based ECG system, etc.

To prevent an accidental change of this setting, the menu is protected with a password. Using the arrow keys, enter 003 and confirm the entry with SELECT.

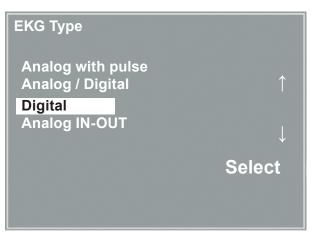


ENTERING THE EKG TYPE PASSWORD

All ergoselect ergometers support the following communication modes:

- Analog with pulse Remote start mode; prior the each load change, the ergometer generates a control pulse and sends the corresponding data via the interface.
- Analog / Digital An analog voltage controls the load - blood pressure measurements can be initiated with digital commands.
- Digital (default) The communication with the ergometer is entirely controlled with digital commands.
- Analog IN-OUT The entire communication (load control and BP measurements) is controlled with analog signals. No digital data will be sent.

Select the communication mode and confirm with SELECT.



SELECTING THE ERGOMETER COMMUNICATION MODE

Note

- The EKG Type needs to be selected only when the ergometer is connected to an ECG unit. The selection is part of the installation procedure.
- The "Analog/Digital" and "Digital" communication is only possible when PC Mode is selected from the main menu or when this is the default mode.

RPM

Here you determine the RPM limits. When these limits are exceeded, the LEDs for high or low speed (RPM) will illuminate.

Select the value to change (Min. or Max.) and confirm with SELECT.

Using the arrow keys, change the corresponding value and save the new value with SELECT.

RPM		
Min ↑	0 70	54 ↑
Max↓	50 130	64 ↓
		Select
	Choose function	

SETTING THE RPM LIMIT VALUES

•	• The limits selected in this menu only apply to the load range between 6 and 150 watts. At higher loads the RF limits automatically adapt to the respective loads:	
	Load (watts)	Green RPM range (1/min)
	6 - 150	54 – 64 (adjustable)
	151 - 250	58 - 65
	251 - 350	68 - 75

78 - 85 88 - 95

98 - 105

108 - 115

118 - 125

> 125

> 130

Note

Pulse Display

351 - 450

451 - 550 551 - 650

651 - 750

751 - 850

851 - 950

951 - 999

The pulse readout on the display can be turned off.

CLEANING, MAINTENANCE, DISPOSAL

GENERAL CLEANING

Wipe the device surface down with a cloth moistened with soap water or a disinfectant.

The cloth should not be dripping wet; do not allow liquids to enter the device.

CLEANING THE SADDLE

Clean the saddle with a soft and dry or moist cloth (**Disin-fectants used should not contain any alcohol**).

CLEANING THE UPHOLSTERY (E.G. COUCH ERGOMETER)

Wipe the upholstery down with a soft cloth moistened with soap water.

The cloth should only be moist and not dripping wet. If the cleaning agents and disinfectants used are caustic or contain alcohol, they may damage and/or discolor the upholstery.

DISINFECTION

Only the following disinfectants are approved for disinfection:

Schülke & Mayr GmbH:

- Antifect® AF, FF, FD 10
- Terralin ® (0,5 %)
- Quartamon Med®

B. Braun Melsungen AG:

- Hexaquart plus[®] (0,5 % / 5,0 %)
- Hexaguart S[®] (1,5 % / 5,0 %)
- Meliseptol[®]
- Melsept SF[®] (0,5 % / 5,0 %)

ECOLAB:

• Incidin Foam®

Warning

• Shock Hazard •

Disconnect the device from the power line before cleaning.

• Equipment Damage •

- Do not allow liquids to enter the equipment. Devices into which liquids have entered must be immediately cleaned and checked by a service technician, before they can be reused.
- Do not use acids, alkaline solutions (household cleaners) or caustic disinfectants.

Note

• The use of cleaning agents and disinfectants containing alcohol is not permitted!

Hint

• Strictly observe the manufacturer's instructions for use.

CLEANING THE BLOOD PRESSURE CUFF

REMOVING THE MICROPHONE

Pull the end of the cuff through the metal clasp and fold out the cuff.

Pull on the short Velcro tab to open the microphone pocket and carefully remove the microphone.

CLEANING

Clean the cuff and tubing with a moist cloth. You can use a dishwashing liquid or mild soap water (no cleaning agents containing alcohol).

Clean the microphone with a cloth moistened with alcohol or soap water.

Allow the microphone to dry before reinserting it in its pocket.

DISINFECTION

For disinfection, spray a disinfectant sparingly on the cuff, the tubing and the microphone.

After the contact time indicated by the manufacturer, wipe all components dry.

Only the following disinfectants are approved for disinfection:

Schülke & Mayr GmbH:

- Antifect[®] AF, FF, FD 10
- Terralin[®] (0,5 %)
- Quartamon Med®

B. Braun Melsungen AG:

- Hexaquart plus[®] (0,5 % / 5,0 %)
- Hexaquart S[®] (1,5 % / 5,0 %)
- Meliseptol[®]
- Melsept SF[®] (0,5 % / 5,0 %)

ECOLAB:

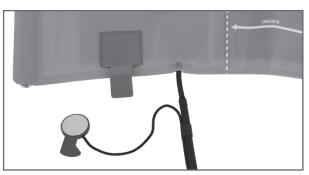
Incidin Foam[®]

INSERTING THE MICROPHONE

Slip the microphone into the pocket, the metal side facing the arm.

Guide the microphone cable out of the pocket and to the right of the Velcro tab. Then close the tab.

Fold the end of the cuff over and introduce it into the metal clasp.



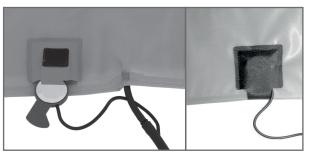
REMOVING THE MICROPHONE

WarningEquipment Damage

- Cuff, microphone and tubing may not under any circumstances:
 - be immersed in liquids
 - be cleaned in a water bath or in running water.

Hint

• Strictly observe the manufacturer's instructions for use.



INSERTING THE MICROPHONE

MAINTENANCE

CHECKS BEFORE EACH USE

Before each use, visually inspect the device for signs of damage.

If you detect damages or impaired functions which may result in a hazard to the patient or the operator, the device must be repaired before it can be used again.

TECHNICAL SAFETY INSPECTIONS AND TECHNICAL INSPECTIONS OF THE MEASURING SYSTEM

The technical safety inspections and the inspections of the measuring system must be completed every two years according to the rules of the art by a Service Engineer authorized by ergoline.

Similarly, the automatic sphygmomanometer in the control terminal must be checked and calibrated by an authorized specialist every two years to fulfill legal requirements. The date of the next inspection is indicated on the inspection sticker attached next to the type plate on the ergometer.

DISPOSAL

Do not dispose the product described in this Operator Manual as unsorted municipal waste. It must be collected separately.

Please contact the authorized manufacturer ergoline GmbH to obtain information concerning the decommissioning of your equipment. There is no proper waste management, proper disposal is documented by ergoline GmbH. Consult operating instructions.



TECHNICAL SPECIFICATIONS

Ergometer

Model	modular ergometer system ergoselect models ergoselect 100 P / K, ergoselect 150 P / K, ergoselect 200 P / K
Operating Mode	continuous operation
Power	100 - 240 V / 50 - 60 Hz (100 VA max.)
	specification power cord US: SPT 2x18AWG 125 V / 10 A "hospital" or "hospital grade"
	specification internal backup battery: IEC: CR 2032 /3V 230 mAh
Braking Principle	computer-controlled eddy current brake with torque measurement; speed independent to DIN VDE 0750-0238
Load Range	6 – 999 watts, speed independent (see diagrams)
Speed Range	30 to 130 RPM
Deviation of Measured Load	to DIN VDE 0750-0238
Load Increments	user programmable
Internal Protocols	 Control Terminal P: 5 fixed incremental exercise test protocols (e.g. WHO) 10 user-programmable protocols manual load control
Internal Protocols	5 fixed incremental exercise test protocols (e.g. WHO)10 user-programmable protocols
Internal Protocols Permitted Patient Weight	 5 fixed incremental exercise test protocols (e.g. WHO) 10 user-programmable protocols manual load control Control Terminal K: 5 fixed incremental exercise test protocols (e.g. WHO) 10 user-programmable protocols manual load control 4 fixed test protocols (e.g. PWC)
	 5 fixed incremental exercise test protocols (e.g. WHO) 10 user-programmable protocols manual load control Control Terminal K: 5 fixed incremental exercise test protocols (e.g. WHO) 10 user-programmable protocols manual load control 4 fixed test protocols (e.g. PWC) 10 user-programmable training protocols max. 160 kg (ergoselect 100 / ergoselect 150 / ergoselect 200)
Permitted Patient Weight	 5 fixed incremental exercise test protocols (e.g. WHO) 10 user-programmable protocols manual load control Control Terminal K: 5 fixed incremental exercise test protocols (e.g. WHO) 10 user-programmable protocols manual load control 4 fixed test protocols (e.g. PWC) 10 user-programmable training protocols max. 160 kg (ergoselect 100 / ergoselect 150 / ergoselect 200) max. 200 kg (ergoselect 200 with stabilizer plate) continuous, for patients between 120 cm and 210 cm ergoselect 100/150: manual adjustment of saddle height ergoselect 200: electrical adjustment of saddle height with

Displays	LCD: 68 x 34 mm, 128 x 64 pixels (Control Terminal P) 115 x 88 mm, 320 x 240 pixels (Control Terminal K) additional LED display for speed (RPM)
Interfaces	PORT 1 (DSUB-9-pole): digital remote control RS232 by PC or ECG recorder, remote start of ECG recorder (optional) USB: digital remote control by PC (driver required)
Dimensions, Weight	ergoselect 100/150: length: 900 mm width: 420 mm (width of handlebar: approx. 535 mm) height: 900 mm - 1350 mm weight: approx. 56 kg
	ergoselect 200: length: 900 mm width: 460 mm (width of handlebar: approx. 535 mm) height: 900 mm - 1350 mm weight: approx. 67 kg
Safety Standards	DIN EN 60601-1, DIN EN 60601-1-2, DIN VDE 0750-238
Protection Class / Degree of Protection	II D / B (ergometer) BF (blood pressure module)
MDD Classification	class Ila to 93/42 EEC
RF Emission	class B to DIN EN 55011 / 5.0 DIN EN 60601-1-2
Environment	operation:temperature:+10 to +40 °C (50 to 104 °F)rel. humidity:30 to 75%, no condensationatmospheric pressure:800 to 1060 hPatransport and storage:-20 to +70 °C (-40 to +158 °F)rel. humidity:10 to 95%, no condensationatmospheric pressure:500 to 1060 hPa
Blood Pressure Module	
Measuring Method:	auscultatory method, oscillometric; for resting BP, the results from both measurements are compared for plausibility
Measuring Range	systolic pressure:40 to 280 mmHgdiastolic pressure:40 to 280 mmHgpulse rate:35 to 230 P/min
Measurement Error, systematic	systole: +/- 3 mmHg diastole: +/- 3 mmHg (temperature: +15 +25 °C)
Standard deviation (clinical trial)	systole / diastole: 7 mmHg (max.)

Inflation Pressure	300 mmHg max.; during inflation the inflation pressure automatically adapts to patient's BP
Inflation Rate	between approx. 6 seconds (to 140 mmHg) and approx. 18 seconds (to 300 mmHg)
Max. Cuff Pressure	300 mmHg
Cuff Deflation Method	pulse-dependent deflation rate approx. 3 mmHg/beat or approx. 3 mmHg/s
Calibration	calibration with external pressure meter
Artifact Rejection	automatic artifact rejection and comparison of the resting BP readings from both methods for plausibility

EXERCISE TEST PROTOCOLS

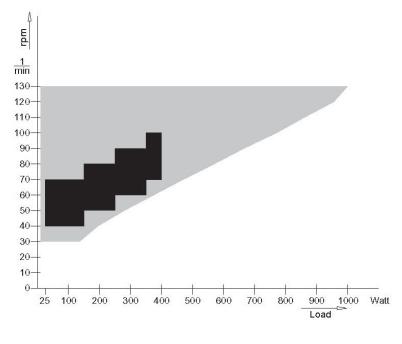
Protocol	initial load [W]	time in stage [min]	load increment [W]	recovery load [W]	recovery time [min]
1. WHO	25	2	25	25	99
2. BAL	50	3	50	25	99
3. Hollmann	30	3	40	25	99
4. STD France	30	3	30	25	99
5. Standard	20	1	25	25	99
6 15. (user-programmable)	25	2	25	25	99
Adjustment Range	20 - 100	1 - 30	1 - 400	20 - 100 (*)	1 - 99

(*) With Control Terminal P, the recovery load is fixed at 25 W.

TEST PROTOCOLS (CONTROL TERMINAL K ONLY)

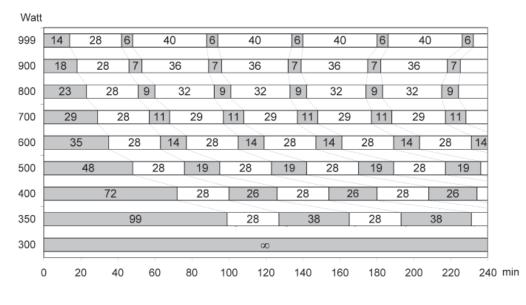
Protocol	initial load [W]	duration [sec]	load increment [W]	time in stage [sec]	recovery load [W]	recovery time [min]
ramping protocol	0	120	25	10	25	99
PWC-130 (*)	25	0	25	120	25	99
PWC-150 (*)	50	0	25	120	25	99
PWC-170 (*)	50	0	50	120	25	99

(*) The program advances to the recovery phase as soon as the target heart rate (130/150/170) is reached.



FAMILY OF CHARACTERISTICS OF THE BRAKING TORQUE CONTROL RANGE

black: speed-independent range to DIN VDE 0750-0238 **black + grey:** speed-independent range of the ergoselect ergometer



FAMILY OF CHARACTERISTICS OF THE LOAD PERIODS ACCORDING TO IEC 60601-1

Under permanent load, the load periods and pauses (white) shall be observed.

Electromagnetic Compatibility EN 60601-1-2

Changes or modifications to this system not expressly approved by ergoline could cause EMC issues with this or other equipment.

This system is designed to comply with applicable regulations regarding EMC.

Its compliance with these requirements has been verified. It needs to be installed and put into service according to the EMC information stated as follows.

Warning RF INTERFERENCE

• Use of portable telephones or other radio frequency (RF) emitting equipment near the system may cause unexpected or adverse operation.

Caution

• Equipment Malfunction •

• The equipment or system should not be used adjacent to, or stacked with, other equipment. If adjacent or stacked use is necessary, the equipment or system should be tested to verify normal operation in the configuration in which it is being used.

GUIDANCE AND MANUFACTURER'S DECLARATION - ELECTROMAGNETIC EMISSIONS

The ergoselect ergometer is intended for use in the electromagnetic environment specified below. It is the responsibility of the customer or user to ensure that the ergoselect ergometer is used in such an environment.

Emissions Test	Compliance	Electromagnetic Environment – Guidance
RF emissions to EN 55011	Group 1	The ergoselect ergometer uses RF energy only for its in- ternal function. Therefore, its RF emissions are very low and are not likely to cause any interference in nearby electronic equipment.
RF emissions to EN 55011	Class B	The ergoselect ergometer is suitable for use in all estab-
Harmonic emissions to EN 61000-3-2	Class A	lishments, including domestic and those directly con- nected to the public low-voltage power supply network
Voltage fluctuations/flicker emissions to EN 61000-3-3	Complies	that supplies buildings used for domestic purposes.

GUIDANCE AND MANUFACTURER'S DECLARATION - ELECTROMAGNETIC IMMUNITY

The ergoselect ergometer is intended for use in the electromagnetic environment specified below. It is the responsibility of the customer or user to ensure that the ergoselect ergometer is used in such an environment.

ImmunityTest	IEC 60601 Test Level	Compliance Level	Electromagnetic Environment – Guid- ance
Electrostatic discharge (ESD) to EN 61000-4-2	± 6 kV contact ± 8 kV air	± 6 kV ± 8 kV	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 to EN 61000-4-4	\pm 2 kV for power supply lines \pm 1 kV for input and output lines	± 2 kV passed	Mains power should be that of a typical commer- cial or hospital environment.
Surge to EN 61000-4-5	± 1 kV differential mode ± 2 kV common mode	± 1 kV N/A	Mains power should be that of a typical commer- cial or hospital environment.
Voltage dips, short interrup- tions and voltage variations on power supply input lines to EN 61000-4-11	< 5 % UT (> 95 % dip in UT) for 0.5 cycles 40 % UT (60 % dip in UT) for 5 cycles	< 5 % UT 40 % UT	Mains power should be that of a typical commer- cial or hospital environment. If the user of the ergoselect ergometer requires continued operation during power mains interruptions, it is recommend- ed that the ergoselect ergometer be powered from an uninterruptible power supply or a battery.
	70 % UT (30 % dip in UT) for 25 cycles	70 % UT	
	< 5 % UT (> 95 % dip in UT) for 5 s	< 5 % UT	
Power frequency (50/60 Hz) magnetic field to EN 61000-4-8	3 A/m	passed	Power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercial or hospital environment. The ergoselect ergometer has no components susceptible to magnetic fields.

GUIDANCE AND MANUFACTURER'S DECLARATION - ELECTROMAGNETIC IMMUNITY

The ergoselect ergometer is intended for use in the electromagnetic environment specified below. It is the responsibility of the customer or user to ensure that the ergoselect ergometer is used in such an environment.

ImmunityTest	IEC 60601 Test Level	Compliance Level	Electromagnetic Environment – Guid- ance
			Portable and mobile RF communications equip- ment should be used no closer to any part of the ergoselect ergometer, including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter.
			Recommended separation distance: d = $1.2 \sqrt{P}$
			d = 1.2 √P for 80 MHz to 800 MHz
Conducted RF to EN 61000-4-6 Radiated RF to EN 61000-4-3	3 Vrms 150 kHz to 80 MHz 3 V/m 80 MHz to 2.5 GHz	3 V 3 V/m	 d = 2.3 √P for 800 MHz to 2.5 GHz where P is the rated output power rating of the transmitter in watts (W) according to the transmitter manufacturer and d is the recommended separation distance in meters (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 (b). Interference may occur in the vicinity of equipment marked with the following symbol
NOTE 2: These guidelines objects, and peo (a) Field strengths from fix broadcast and TV broad	ople. red transmitters, such as base station deast cannot be predicted theoretical	s for radio (cellular/cordless) t	fected by absorption and reflection from structures, telephones and land mobile radio, AM and FM radio e electromagnetic environment due to fixed RF transmit- in the location in which the ergoselect ergometer is used
	RF compliance level above, the ergos ce is observed, additional measures n	-	served to verify normal operation. orienting or relocating the ergoselect ergometer.
(b) Over the frequency ran	ge from 150 kHz to 80 MHz, field stre	enaths should be less than 3 \	//m

RECOMMENDED SEPARATION DISTANCES BETWEEN PORTABLE AND

MOBILE RF communications equipment and the ergoselect ergometer

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

Rated Maximum Output Power of	Separation Distance According to Frequency of Transmitter [m]			
Transmitter [W]	150 kHz to 80 MHz d = 1.2 √P	80 MHz to 800 MHz d = 1.2 √P	800 MHz to 2.5 GHz d = 2.3 √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	3.7	3.7	7.37	
100	11.7	11.7	23.3	

For transmitters rated at a maximum output power not listed above, the recommended separation distance d in meters (m) can be estimated 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.



ergoline GmbH Lindenstraße 5 72475 Bitz Germany

 Tel.:
 +49-(0) 7431 98 94 - 0

 Fax:
 +49-(0) 7431 98 94 - 128

 e-mail:
 info@ergoline.com

 http:
 www.ergoline.com