Gebrauchsanweisung

Aneroid Blutdruckmessgerät und Manschette

Instructions

Aneroid Sphygmomanometers and Cuffs

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English Information on the various models of Riester aneroid sphygmomano meters

You have acquired a high-quality **Riester** precision aneroid sphygmomanometer, which has been manufactured in accordance with the ISO 81060-1: 2007 standard [DIN EN ISO 81060-1: 2013] and is subject to the strictest quality controls. The excellent quality will guarantee years of reliable measurements.

Intended purpose: The aneroid sphygmomanometers from Riester are used by physicians and persons trained in auscultatory blood pressure measurement to determine the systolic and diastolic blood pressure in hum-ans (adults, children, babies and neonates). Riester aneroid sphygmomanometers are intended exclusively for blood pressure measurement on healthly skin on the upper arm or thigh. The product is normally used professionally in a medical practice or a hospital. The sphygmomanometer/aneroid manometer is intended for use as a diagnostic aid.

Product description / indication: All blood pressure measuring devices / aneroid manometers distributed by company Rudolf Riester GmbH are of the same basic structure. Blood Pressure measuring devices / aneroid manometers consists of measuring system, cuff and inflation system (consisting of the pump [ball with non-return valve], and the screw valve lair lock or air drain]. All devices work according to the same measuring principle: Indirect blood pressure measurement according to the Korotkoff method.

.... ttention to the paragraph "blood pressure measurement" in the instructions for use, the blood pressure urement can only done from doctors and persons trained in auscultatory blood pressure measurement

information on the label.

-Check the pointer zero position as described in "Test of precision".

-For Germany, the Medical Devices Operator Ordinance applies. See also point "metrological control".

-A description of where the LOT No. is located can be found in the cuff table at the end of the instructions for use.

1. Model exacta® and sphygmotensiophone
Thou are equinned with 2-tube cuffs because the pressure gauge is not directly connected to the inflation

system.

2. Model: e-mega, R1 shock-proof®, minimus® II, precisa® N, precisa® N shock-proof, babyphon® and ri-san® 1-tube.

and ri-san® 1-tube.
They are equipped with 1-tube cuffs. Here the inflation system is directly connected to the manometer.

3. Model e-mega, minimus® III, precisa® N double-tube
They are equipped with double tube cuffs. Here the inflation system is directly connected to the manometer.

4. Model sanaphon® and ri-san®
These devices are devices for self-measurement. The manometer and the inflation system form a unit. In the 1-tube cuff, a chestpiece is integrated, which absorbs the Korotkoff sounds and forwards them to the ear via the enclosed stethoscope, which must be screwed into the thread on the outside of the cuff.

5. Models big ben Round / Square desk, wall, mobile and anesthesia and ri-former® big ben

Installation instructions for the different models (with the exception of the desk model): 5.1. Wall modell / ri-former® big ben
Remove the wing nut underneath the cuff basket and take off the wall bracket. Hold the bracket against the
wall at the desired location and mark the positions of the mounting holes for drilling. Drill the holes and insert
screw anchors in them. Now you can fasten down the wall bracket with screws. Place the unit on the wall
bracket so that the top part of the wall bracket engages the edge of the cuff basket and the bottom part fits
over the screw that protrudes from the underside of the cuff basket. Now replace and tighten the wing nut on

5.2. mobile stand version / bigben

Adjustment:
By opening the retaining screw the desired height can be adjusted. After adjusting the height the mobile stand must be fastened by the retaining screw again.

5.3 anesthesia model / bigben
Remove the wing nut underneath the cuff basket and take off the wall bracket. Affix the wall bracket to the back of universal clamp No. 10384 using the screws included. Place the unit on the wall bracket so that the opp art of the wall bracket engages the edge of the cuff basket and the bottom part fits over the screw that protrudes from the underside of the cuff basket. Now replace and tighten the wing nut on the protruding screw.

Selection of suitable cuff sizes

A. Nylon-velcro cuffs, Disinfectable one piece cuff
Our nylon-velcro cuffs, Disinfectable one piece cuff have a tufted strip on one side and hooks on the other. This ensures that the cuffs can be easily opened and closed quickly and repeatedly. On all the models (with the exception of sanaphor®), the cuffs are calibrated, i.e. provided with measurement lines. To make sure that you have chosen the right cuff size, check to see whether the white index line is in the range between the arrows after the cuff has been put on. If the index line fails to reach this range, the cuff is too small. If it is beyond the range, the cuff is too large. Precise blood pressure readings can be obtained only if the correct cuff size is used.

Nylon-Velcro cuffs (Two Piece Reusable Velcro Cuff), The following cuff sizes are available for all models (except sphygmotensiophone, sanaphon® and exacta®), see cuff table:
Disinfectable one piece cuff (One Piece Reusable Velcro Cuff One Tube/Two Tube), The following cuff sizes are available for all models (except sphygmotensiophone,sanaphon, ri-san self-measurement and exacta), see cuff table:

sanaphon® and ri-san® (Two Piece Reusable D-Ring Cuff Self-Mesurament): Measure your arm circumference to make sure that it lies within the range indicated on the sizes are available: child, adult, large adult and thigh. These sizes correspond to the circum the above table.

B. Cotton-velcro cuffs self-measurement [Two Piece Reusable Cuff sphygmotensiophone]:
Our cotton-velcro cuffs have a tufted strip on one side and hooks on the other. This ensures that the cuff can be easily opened and closed quickly and repeatedly. Measure your arm circumference to make sure that it lies within the range indicated on the cuff.

The following cuff sizes are available for the models listed below, see cuff table:

C. Cotton hook cuffs (Two Piece Reusable Hook Cuff):

circumference of your arm to make sure that it lies within the range indicated on the cuff fabric. Measure the Circumference of your arm to make sure that it lies within the range indicated on the cuff.

The following cuff sizes are available for the models listed below: R1 shock-proof®, minimus® III, minimus® III, big ben Round / Square (all versions) and ri-san , siehe cuff table:

D. Cotton bandage cuffs (Two Piece Reusable Bandage Cuff):
There is a bandage strip and a hook on one side of the bandage cuff. To attach the cuff, simply insert the hook into the bandage strip. Measure the circumference of your arm to make sure that it lies within the range indicated on the cuff.

The following cuff sizes are available for the models listed below: R1 shock-proof*, minimus* II, minimus* III, bigben Round / Square [all versions] and ri-san*, see cuff table:

Patient position for intended use: sit comfortably, legs not crossed, back and arm supported, center of cuff on the upper arm at the level of the right cardiac atrium; the patient should be as relaxed as possible and should refrain from talking during the measuring process; approx. 5 minutes should elapse before the first measurement is taken.

Close the valve by turning the deflation screw clockwise (except in the case of ri-san*).

Putting on the cuff: Put the cuff on in such a way that the lower edge of the cuff is approx. 2 to 3 cm above the crook of the arm [level of the right cardiac atrium) or approx. 5 cm above the knee joint. Make sure that the marking sign is placed above the artery. The white index strip should be within the marked index range.

- index range.
 After putting on the cuff, inflate the cuff to approx. 20 mmHg above the expected systolic blood pressure value [- the upper value] with the aid of the ball.
 When measuring blood pressure, the position of the operator is normally in front of the patient or at the patient's side.
- e. ended to use phase V of the Korotkoff sounds (K5) for auscultatory measurement in adults, to
- use phase V of the Korotkoff sounds (KA) for auscultatory measurement in children aged 3 to 12 years, and to use phase V of the Korotkoff sounds (KA) for auscultatory measurement in children aged 3 to 12 years, and to use phase V of the Korotkoff sounds (K5) for auscultatory measurement in pregnant female patients, except if the Korotkoff sounds can be heard during deflation of the cuff if this is the case, K4 is
- to be used.

 Place the chestpiece of the stethoscope, preferably our model anestophon, catalog no. 4177-01 4177-05, above the artery underneath the cuff.

 For the units intended for home use, no separate stethoscope is needed, since the chestpiece is integrated in the cuff. If a unit intended for home use is used, the membrane of the chestpiece installed in the cuff must be placed above the artery. The cuff is put on by pulling the free end of the cuff through the metal retainer and closin the cuff with the aid of the hook-and-loop fastener.

- retainer and closing the cuff with the aid of the hook-and-loop fastener.

 A Nylon-velorc cuffs: Close the cuff using the Velcro hook-and-loop fastener.

 B. Bandage cuffs: Fasten the bandage cuff by hooking the hook onto the bandage strap.

 C. Hook cuffs: In the case of a hook cuff, the metal hook is hooked onto the small metal rods of the cuff cover.

 To enable blood pressure measurement, open the deflation screw by turning it counterclockwise. The de
 flation rate should ideally range between 2 and 3 mmHg/s and can be set by delicate adjustment of the
 screw. Visual check of the deflation rate: The needle must move through 1 to 1.5 scale marks per second on
 the scale. After completion of the measurement, open the valve completely for rapid deflation of the cuff.
 The rif-sam? model is equipped with a push-button valve. Activate this valve in such a way that the ideal
 deflation rate between 2 and 3 mmHg/s is achieved. To completely deflate the cuff, press the button all the
 way to the stop position.

 When the upper blood pressure value (systole) has been reached, rhythmic beats can be heard.
 Systole = The upper blood pressure value is the value that results when the heart contracts and the blood is
- - ssels. essure value (diastole) has been reached, the beats fall silent.

again filling up with blood.

The blood pressure measurement has been completed.

We would like to point out to you that a unit intended for home use does not replace regular visits to the doctor and that only the doctor can accurately analyze your measured values.

General information

General information
The goal of cleaning and disinfection of medical products is the protection of patients, users and third persons and conserving the value of the medical products. On account of the product design and the used material, no defined limit of maximum processing cycles can be fixed. The lifetime of the medical products depends on their function and on a appropriate treatment of the devices. Before returning faulty products for repair they must have gone through the described reprocessing process.

tb can be cleaned outside with a humid cloth until optical cleanness is given.

ATTENTION!

Never place the manometer in liquid! This item is not approved for automated reprocessing and sterilization.

2. Cuffs
Cotton and Nylon velcro cuff (latex and latex free)
Cleaning:

Cleaning: After removing the bladder, wipe the nylon-velcro covers with a damp cloth. Alternatively, these can be washed with soap and cold water like all the other cuffs. If you decide on the latter course, rinse the cuffs with clear water afterwards and let them air dry. Wipe the bladder and tubes with a damp cloth.

Disinfection:After removing the bladder, wash the cuff covers in cold water to which disinfectant has been added. After-wards, let them air dry. Only disinfectants with approved efficiency and in accordance with the national sta dards can be used. The bladder and tubes can be be wiped with a cotton cloth moistened with ethanol.

Disinfectable one piece cuff
Cleaning:
The cuff can be wiped with a damp cloth. Alternatively, it can be washed with soap and cold water like all the other cuffs. Please rinse the cuff with clear water afterwards. In addition this cuff can be washed at up to 60° C in the washing machine. Before next use, please ensure that no liquid is remaining in the cuff. This can affect the measurement results negatively and it can damage the manometer technology.

Disinfection:The cuff can be completely inserted into liquid disinfectant. Only disinfectants with approved efficiency and in accordance with the national standards can be used. Before next use, please ensure that no liquid is remaining in the cuff. This can affect the measurement results negatively and it can damage the manometer technology.

IMPORTANT! IDP VALUAN : Do not iron nylon-velcro cuffs, Disinfectable one piece cuff. Never expose the cuffs to intensive solar radiation! Never touch the cuff covers or bladders with a sharp instrument, since this could cause damage!

Maintenance The product does not require any maintenance.

om the manometer and hold the manometer in a vertical position. When the still at 0 on the scale, the instrument has been adjusted properly. If the pointer is below or above 0, the instru-ment must be recalibrated. Either take it to an authorised **Riester** dealer or send it to us.

Monitoring of instruments
All countries except for Germany:
The respective legal provisions apply for all countries, except for Germany. The reference manometer, which is used for calibration, must be traceable to national and international measurement standards.

Technical Data

mal environmental conditions under which the error containing in a relative air assurement conditions:

10°C (50°F) to 40°C (104°F) at a relative air humidity of 85 % (non-condensing) rage conditions:

-20°C (-4°F) to 70°C (158°F) at a relative air humidity of 85 % (non-condensing) risions:

Aneroid, desk, wall, stand and anaesthetic model Round scale

Round scale Increments of 2 mm Hg 0 to 300 mm Hg 0 to 300 mm Hg

No stop pin I or 2, depending on the particular model

Air-release valve that can be regulated

nation on disposal from your appropriate local facility or from your local env-

Auf dem Gerät bzw. Auf der Verpackung finden sich folgende Symbole. The following symbols are depicted on the packaging. Les symboles suivants figurent sur l'appareil ou sur son emballage. En et dispositivo o en su embalaje encontrará los siguientes símbolos.

	LATEX	XX XX
Two Piece Reusable Wefcro Cuff LOT Inside the Cover	₩	₩
Newborn	5-7,5cm	5-8cm
Infant	7,5 - 13 cm	8-13cm
Child	13 - 20 cm	13-17cm
Small Adult	17 - 26 cm	17-24cm
Adult	24 - 32 cm	24-34cm
	32 - 48 cm	34-44cm
Thigh	42 - 50 cm	42-50cm
Thigh XL	50 - 70 cm	50-70cm
One Piece Reusetible/ClatoGistTQbe/Tube Tibe Tube LOT On the Cover		
Newborn		5 - 8 cm
Infant		8 - 13 cm
Child		13 - 17 cm
Small Adult		17 - 24 cm
Adult		24 - 32 cm
		32 - 41 cm
Two Piece Reusable Cuff Sphygmotensiophone One Inside the Cover		
Infant	7,5 -13 cm	
Child	13 - 20 cm	13 - 20 cm
Adult	24 - 32 cm	24 - 32 cm
Two Piece Reusable Hook Cuff LOT Inside the Cover		
Adult	24 - 32 cm	24 - 32 cm
Two Piece Reusable Bandage Cuff Lot Inside the Cover		
Adult	24 - 32 cm	24 - 32 cm

WARRANTY

This product has been manufactured under the strictest quality standards and has undergone a thorough final quality check before leaving our factory. We are therefore pleased to be able to provide a warranty of 2 years from date of purchase on all defects, which can verifiably be shown to be due to material or manufacturing faults. A warranty claim does not apply in the case of improper handling. All defective parts of the product will be replaced repaired free of charge within the warranty period. This does not apply to wearing parts. For R1 shock-proof, we grant 5 years, for precisa N shock-proof, 2 years for the calibration, which is required by CE-certification. A warra claim can only be granted if this Warranty Card has been completed and stamped by the dealer and is enclosed whe product. Please remember that all warranty claims have to be made during the warranty period. We will, of course, be pleased to carry out checks or repairs after expiry of the warranty period at a charge. You are also welcome to request a provisional cost estimate fromus free of charge. In case of a warranty claim or repair, pleas return the Riester product with the completed Warranty Card to the following address:

Seriennummer bzw. Chargennummer, Serial number or batch number Numméro de série/de lot, Número de serie o de lote Серийний номер или номер партии, Numero di Serie risp. numero di carica

Datum, Date, Date, Fecha, дата, Data,

Stemnel und Unterschrift des Eachhändlers

rsteller / Manufactured by / Fabricant / Fabricante / фабрика́нт / Zulässiger Temperaturbereich in °C für Lagerung und Transport Temperature limits in °C for storage and transport Limites de temperature, en°C, lors du stockage et du transport ímites de temperatura en °C para almacenamiento y trans опустимый диапазон температур в °C при хранении и нспортировке rvallo di temperatura ammesso in °C per la conservazione e il tra: Zulässiger Temperaturbereich in °F für Lagerung und Transport Temperature limits in °F for storage and transport Limites de temperature, en °F, Iors du stockage et du transport Limites de temperature an °F para almacenamiento y transporte Допустимый диапазон температур в °F при хранении и транспортировке Intervallo di temperatura ammesso in °F per la conservazione e il traspor Zulässige Luftfeuchtigkeit für Lagerung und Transport Humidifty limitation for storage and transport Limites d'humidife pendant le stockage et le transport Limite de humedad para almacenamiento y transporte Допустимая впажность воздуха при хранении и транспортировке Umidità dell'aria ammessa per la conservazione e il trasporto **(%)** Markierungszeichen auf der Manschette für die Lage der Markierungszeichen auf der Manschette für die Lage der Manschette auf der Arterie.
Marking sign on the cuff to indicate the position of the cuff above the artery. Marquage sur le brassard permettant le positionnement de ce dernier sur l'artère Marca de referencia en el brazalete para ubicación del brazalete sobre la arteria.
Merки на манжете обозначают требуемое положение манжеты на артерии.
Simbolo гірогатов su bracciale che indica la posizione del bracciale stesso sull'arteria. "Ф" Gebrauchsanweisung befolgen Follow the operating instructions Subrre les instructions Seguir las instrucciones del manual Соблюдать руководство по применению Rispettare le istruzioni per l'uso

CE-Kennzeichnung kennzeichnet die Übereinstimmung mit der Europäischen Medizinprodukterichtlinien 93/42 EWG DE marking indicates compliance with European Medical Devices Directiv

(3)42 EEU
Le marquage CE indique que le produit est conforme à la directive eupoéenne relative aux dispositifs médicaux 93/42/CFF

ropéenne rélative aux dispositifs médicaux 93/42/CEE
Marca CE que indica la conformidad con la directiva sobre productos sanita
rios europeos 93/42 СЕЕ
Мариморика ОЕ подтверждает соответствие европейской Директиве п
продуктам медицинского назначения 93/42 ESC
Il contrassegno CE definisce la conformità con la Direttiva europea sui
Dispositivi Medici 93/42 CEE

Seriennummer / Serial number / Numéro de série / número de serie / юмер се́рии / Numero di serie

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rocken Lagern eep dry Jaintenir au sec Хранить в сухом месте Conservare in luogo asciutto Grüner Punkt (länderspezifisch) Grüner Punkt (country-specific) Point Vert (spécifique au pays) Punto verde (especifico del país) Маржировка «Зепеная точка» (действует в конкретной стране) Punto Verde (specifico del pase) LATEX Latexfree Sans latex Senza lattice **без латекса** Sin látex Achtung! Gebrauchsanweisung befolgen. Caution! Follow the operating instructions. Attention! Suivez les instructions d'utilisation (3) Attenzione! Seguire le istruzioni operative! Внимание! Следуйте инструкциям по эксплуатации. Atención! Siga las instrucciones del manual de uso! _OT-Nummer / YY = steht für den jeweiligen Monat / XXXX = steht für das ' / YY = stands for the respective month / XXXX = stands for the LOT YY/XXXX rispettivo anno LOT-номер / YY = обозначает месяц производства / XXXX = обозначает год производства Notareno de lote / YY = representa el mes respectivo / XXXX = representa el año respectivo €0\$ erstellungsdatum ate of Manufacture ate de fabrication ata di produzione Дата производства Fecha de manufactur Ŵ