

ENGLISH

ELECTRONIC HANGING SICK LIFT SCALE

MOD. RS – lb version





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By choosing the WUNDER mod. **RS** professional electronic hanging scale, you have purchased a high precision instrument. Since over 40 years Wunder has placed its experience at the service of health. This instrument is compliant with national standards in hospitals and clinics with medical class Im with measurement function and is calibrated in conformity with accuracy class III.

<u>1. GENERAL RULES</u>

Carefully read this manual before using the instrument as it supplies important indications concerning OPERATING SAFETY AND MAINTENANCE.

WUNDER reserves the right to modify the images in the following manual, only if they are purely aesthetic modifications and do not affect the safety and performance of the instrument, without communicating the updates promptly. **Conventions:** The following symbols have been used in this manual:

CC 0476	MEDICAL DEVICE IN COMPLIANCE WITH COMMUNITY DIRECTIVE 93/42/EEC			
С Є М 0474 М	INSTRUMENT SUITABLE FOR LEGAL USE, IN COMPLIANCE WITH DIRECTIVE 2014/31/EU AND EUROPEAN STANDARD EN45501			
	INSTRUMENT IN COMPLIANCE WITH NAWI METRIC DIRECTIVE ACCURACY CLASS III 90/384 - 2014/31/UE AND THE EN45501 EU STANDARD			
	ATTENTION! PLACED BEFORE DETERMINING PROCEDURES. COMPLIANCE FAILURE CAN HARM THE OPERATOR OR PATIENT OR DAMAGE THE PRODUCT			
X	WASTE DISPOSAL IN COMPLIANCE WITH 2012/19/UE DIRECTIVE			
Ť	TYPE B PARTS SUPPLIED Image: Battery power			
-	INDICATION OF WEIGHT FUNCTIONALITY	→0←	INDICATION OF STABLE WEIGHT	
(())	POSSIBLE INTERFERENCES NEAR THE INSTRUMENT		DUAL INSULATION (CLASS II)	
	READ THIS MANUAL CAREFULLY BEFORE USING THE INSTRUMENT			
	MANUFACTURER: WUNDER SA.BI. SRL – VIA VECCHIA PER MONZA, 20 – TREZZO S/ADDA (MI), ITALY			



<u>2. SAFETY</u>

Operators must read this manual carefully, follow the instructions contained therein and become familiar with the correct use and maintenance procedures of the instrument.

This manual contains important information for the installation, use and maintenance of the dynamometer.

The manufacturer assumes no responsibility for direct or indirect damage, including loss of profits, or for any other commercial damage that may result from the use of the product that does not comply with the instructions in this manual. Keep this manual and the declaration of conformity for consultation and support of personnel training:

- Do not overload the instrument beyond the maximum flow value.
- Do not apply loads abruptly.
- Do not use sharp or pointed objects to press the keys.
- Do not attempt to open the instrument.
- Do not remove the seals on the instrument.
- Do not short-circuit the battery terminals.

• Use only the power supply provided by Wunder and before use check the compatibility between the local mains voltage and the adapter's rated voltage (if equipped).

- Periodically check the integrity of the instrument power cable and do not come into contact with hot appliances.
- Make sure that the power cord does not create a risk of getting in the way or getting caught.
- Before cleaning the instrument, unplug the power cord.
- Do not immerse the instrument in water or other liquids.
- Have the maintenance operations and the subsequent metric checks done regularly (see paragraph).
- If on wheels, make sure that the instrumentation can not move accidentally. Use the parking brake while the patient sits up and stands up and help people who do not stand up properly.

NOTE: The medical device requires special precautions regarding electromagnetic compatibility and must be installed and used according to the information supplied with the accompanying documents.



\triangle		
	ATTENTION!	IMPORTANT DIRECTIONS

	<u>A.</u>	TTENTION! IMPORTANT DIRECTIONS	
	The assembling of RS dyr	namometer must be done by skilled operators only and before the use be	
\wedge	sure that all hanging sick l	ift elements, dynamometer and sling are properly assembled, to weight the	
	patient on safety condition	ns. A faulty assembling of these elements could cause the risk of falling with	
	serious consequences for	the patient.	
	AFTER THE ASSEMBLY AN	ID BEFORE PILLING THE PATIENT, THE OPERATOR MUST CHECK IF:	
\wedge	※ THE PATIENT IS PROT	ECTED WITH SOFT PARTS TO AVOID IMPACTS OR SERIOUS DAMAGES.	
	IN PARTICULAR, PLA	CE THE PATIENT MATTRESSES, PILLOWS, BED OR ANY SOFT ELEMENT TO	
	GUARANTEE THE BES	T SAFETY OF THE PATIENT WHILE CHECKING THE WEIGHT INTO	
	SUSPENSION		
	※ The RS dynamometer a	and the harness are in VERTICAL POSITION	
	※ The operator must ma	ke sure that the weighing is carried out with raising unhealthy patients.	
	※ The operator who duri	ng the lifting and patient weighing phase MUST ENSURE that the patient	
	raises with the attached d	ynamometer is stationary on a level surface.	
	※ IT IS ABSOLUTELY FO	RBIDDEN to move before, during and after the weighing phase, the patient	
	raises with dynamome	eter mod. RS300 and the harnessed patient	
	※ The operator must ALV	NAYS accompany and hold the patient's harness during the patient lifting	
	phase to prevent torsions	and sudden movements of the dynamometer and harness that can cause	
	 breakage and irreparable damage, with the risk of serious consequences for the patient. MONOT MOVE, TRANSPORT or ROTATE the patient raises with a dynamometer with the patient in charge. When the weighing phase is completed, the patient must be removed from the harness and 		
	moved with other devices in safety		
360°		means of RS dynamometer, the operator must be sure that the patient	
	remains still, to avoid any	instruments torsion and the following uncorrect weighing.	
NO TORSION	After weighing phase the	operator must <u>ALWAYS</u> follow and hold the sling of the patient to avoid	
3607	torsion and sudden movement of both dynamometer and sling that could cause breakages and		
	irreparable damages, with the risk of serious consequences for the patient.		
NO TORSION			
1	2		
	CUSCINO	<u>ATTENTION</u>	
	DI SICUREZZA	To measure the patient ALWAYS IN SAFETY	
		The operator MUST place a cushion	
J.	7	Under the patient raised	
0		<u>stat die patent faisea</u>	





- When using electrical components under increased safety requirements, always comply with the appropriate regulations.

- Improper installation will render the warranty null and void.
- Ensure the voltage marked on the power supply unit matches your mains power supply.
- This device is designed for use indoors.
- Observe the permissible ambient temperatures for use
- The device meets the requirements for electromagnetic compatibility. Do not exceed the maximum values specified in the applicable standards.
- Ensure that the patient does not lean against the device risk of falling! If you have any problem, contact your local service partner.
- The maximum scale loading must not be exceded. The device beeps when loading exceeds its maximum capacity.

2.1 Intended use

This device is intended to be used for the suspension weighing of patients for general diagnostic purposes. **Environment of use**: in hospitals and specialized medical clinics. The installation room must be equipped with an electrical system that complies with the regulations in force. It is recommended to use the device in environments not exposed to magnetic interference.

Personnel destined to use the product: specialized operators and doctors who are aware of all the safety procedures for correct use.

Control and Responsibility: the medical device must be used under the supervision of a qualified doctor or qualified maintenance personnel and periodic checks that are aware of all safety procedures.

Limitations of use: this medical device can only be used as described in this manual

Useful life of the product: 7 years



2.2 Manufacturer's guide and declaration – Electromagnetic Immunity

The electronic scales **RS** model is scheduled for operation in the electromagnetic environment specified below.

The customer and the user should ensure that it is used in that environment.

Guide and Statement of manufacturer – Electromagnetic emissions			
Emission test	IEC 60601 Conformity	Electromagnetic environment guidance	
RF Emission CISPR11	Group 1	RS model uses RF energy only for its internal function. Therefore, its RF emissions are very low and are not likely to cause any interference in nearby electronic equipment.	
RF Emission CISPR11	Class B	The product is suitable for use in all	
Harmonic emission IEC 61000-3-2	Class A	establishments, including domestic establishments	
Voltage fluctuations / flicker emissions IEC 61000-3-3	Compliant	and those directly connected to the public low-voltage power supply network that supplies buildings used for domestic purposes.	

Guidance and manufacturer's declaration - Electromagnetic Immunity			
Immunity test	IEC 60601 Compliance level	Electromagnetic environment guidance	
Electrostatic discharges (ESD) IEC 61000-4-2	± 8 kV contact ±2, ±4, ±8, ±15 kV air	The floors should be made of wood, concrete or ceramic. If the floors are covered in synthetic material, the relative humidity should be at least 30%.	
Electrical fast transient / burst IEC 61000-4-4	± 2kV for power supply lines ± 1kV for input/output lines	The power supply should be of the type used typically in commercial or hospital environments.	
Surge IEC 61000-4-5	± 1kV line(s) to line(s) ± 2kV line(s) to earth	The power supply should be of the type used typically in commercial or hospital environments.	
Voltage dips, short interruptions and voltage variations on power supply input lines IEC 61000-4-11	0% UT for 0,5 cycle 0% UT for 1 cycle 70% UT (30% dip in UT) for 25s 0% UT for 5 s Note: U_T is the A.C. main voltage prior to application of the test level	The power supply should be of the type used typically in commercial or hospital environments. If the user requires continued operation, it is recommended that the product is powered from an uninterruptible power supply or a battery.	
Power frequency (50, 60 Hz) Magnetic field IEC 61000-4-8	30 A/m	The product power frequency magnetic fields should be at levels of a typical location in a typical commercial or hospital environment.	

ATTENTION!

The medical device requires particular electromagnetic compatibility precautions and must be installed and used according to the information provided in this manual.



Manufacturer's guide and declaration - Electromagnetic emissions			
Immunity test	IEC 60601 Compliance Level	Electromagnetic environment-guidance	
Conducted RF	3Vrms 150kHz to 80MHz (for	Portable and mobile RF communications equipment should	
IEC 61000-4-6	appliances that are not life	be used no closer to any part of the product including cables,	
	supporting)	than the recommended separation distance calculated from	
		the equation applicable to the frequency of the transmitter.	
		Recommended separation distance	
		d = 1.2 √ P	
		d = 1.2 √ P from 80 MHz to 800 MHz	
		d = 2.3 √ P from 800 MHz to 2.5 GHz	
	3 V/m	${\bf P}$ is the maximum output power rating of the transmitter in	
Radiated RF	80MHz to 2,7 GHz	watts (W) according to the transmitter manufacturer and ${f d}$ is	
IEC 61000-4-3	(for appliances that are not	the recommended separation distance in metres (m). Field	
	life equipment)	strengths from fixed RF transmitters, as determined by an	
		electromagnetic site survey ¹ , should be less than the	
		compliance level in each frequency range ² . Interference may	
		occur in the proximity of equipment marked with the	
		following symbol:	

¹ From 80 MHz to 800 MHz is applied the higher frequency range.

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

- a) The intensity of the field for fixed transmitters such as base stations for radio, mobile and cordless phones and land radio mobile, amateur radio, radio transmitters in the AM and FM and TV transmitters cannot be predicted theoretically with accuracy. To establish an electromagnetic environment due to fixed RF transmitters, it should consider the electromagnetic survey of the site. If the field strength measured at the place where you use the instrument exceeds the applicable level of compliance of the above, the device should be observed to verify normal operations. If you notice abnormal performance, it may take additional measures such as a different orientation of the device or re-locate it.
- b) The field strength over a frequency range of 150 kHz to 80 MHz should be less than 3 V/m.



Recommended separation distance between RS Scale and mobile RF communications equipment

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

Output power rating	Separation distance according to frequency of transmitter (m)		
of the 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	0,38	0,38	0,73
1	1,2	1,2	2,3
10	3,8	3,8	7,3
100	12	12	23

For transmitters with maximum rated power output not reported above, the recommended separation distance **d** in meters (m) can be calculated using the equation applicable to the frequency of the transmitter, where **P** is the maximum rated power output of the transmitter in Watt (W) according to the manufacturer of the transmitter.

Note: ¹ From 80 MHz to 800 MHz is applied the higher frequency range.

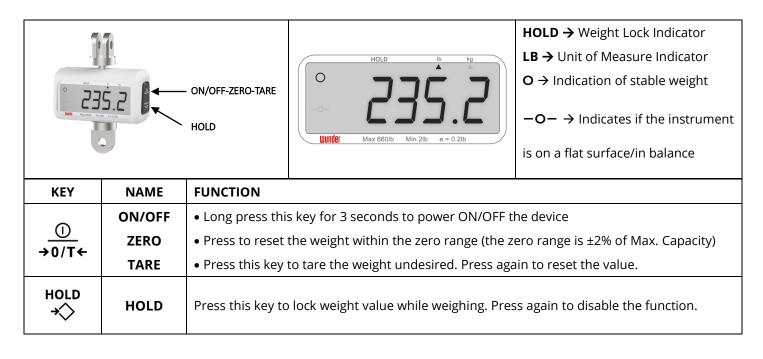
Note: ² These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.

Model	RS300
Factory	Wunder Sa. Bi. Srl - Trezzo sull'Adda (MI) Italy
Max Capacity : Division	Max 660 lb : e=0.2 lb (min = 2 lb)
OIML	Class III
Unit Weight	lb (OIML)
Display	LCD 1inches with 5 digits
Dimension	4.72 x 2.75 x 6.3 inches
Key & Functions	ON/ZERO-OFF-TARE, HOLD
Power	6 alkalin batteries mini-stilo code AAA
Operating Temperature	32 °F / 104 °F
RoHS2	Comply with Directive 2011/65/CE (included delegated directive EU 2015/863)

<u>3. TECHNICAL FEATURES</u>



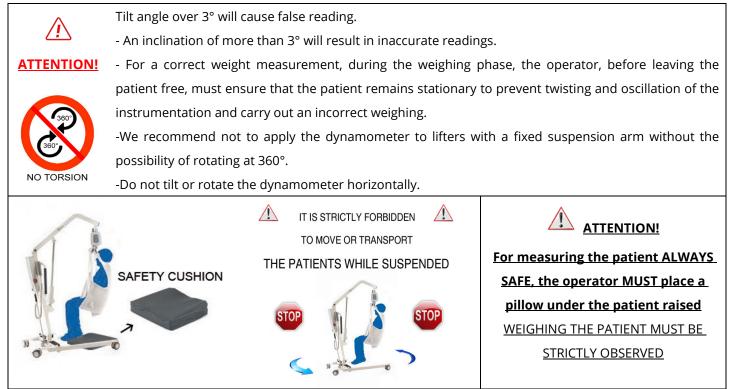
<u>4. KEY PANEL</u>



5. WEIGHING OPERATION

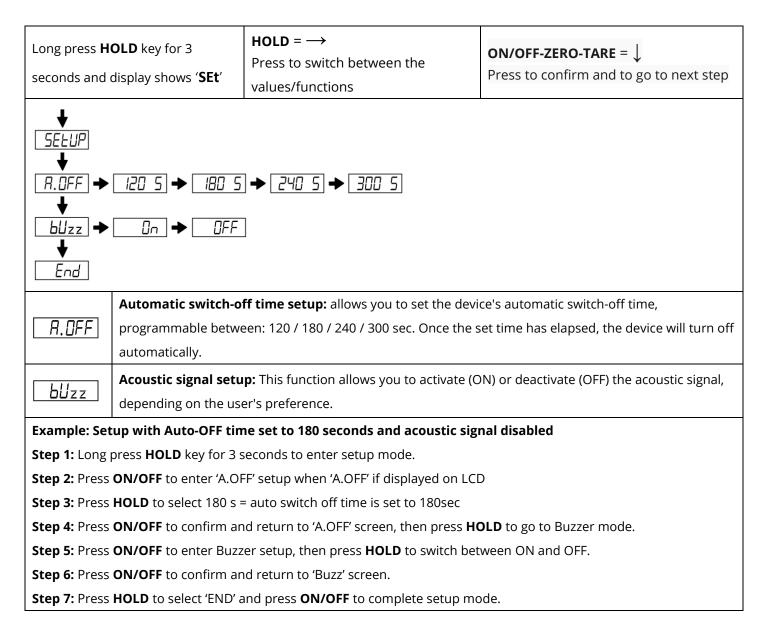
Before reading detailed instructions on how to use all the weighing functions, please read the following guidelines:

- Always be sure that the display shows `Zero` before use, if it doesn't then press the **ZERO** key.
- The Professional Medical Hoist scale is designed to detect when a stable weight is achieved, your reading should be taken at this point.





6. FUNCTION SETTINGS





7. INSTRUCTION FOR BATTERY INSTALLATION

RS uses six AAA size alkaline batteries. Please read the following instruction before using the scale.

1. Find the battery cover on the backside of the scale	2. Remove the battery cover.
3. Take out the battery case	4. Installing new AAA size batteries.
5. Reinstalling the battery case.	6. Installing the battery cover.

8. WARNING INFORMATION

1. Low battery indication This warning show that the voltage of battery is very low, please change a new one.	LobAt
2. High Zero The loading is over limit when power on, please reduce the loading.	00000
3. Low Zero The loading is under limit when power on, please increasing the loading.	00000
4. Overload or Counting error	
The loading is over limit when power on, please reduce the loading and try again.	Fre
If the trouble still exists, please call the service.	
5. EEPROM error	١٢
The programs of the scale is in error, please call the service.	נוונ

9. CLEANING & MAINTENANCE

ATTENTION!

IT IS RECOMMENDED TO KEEP THE INSTRUMENT SUPPLIED UNDER CONTROL WITH A CORRECT PERIODIC MAINTENANCE

We recommend having this check carried out by qualified personnel to carry out the operation. For further information please contact the WUNDER Customer Service Technical Service, which is at your disposal. For a better and longer life of the product it is good to periodically perform a thorough general cleaning.

Periodically perform (at least once a year) functional checks of the RS dynamometer, as follows:

- Control of mechanical parts, hooks, pins, screws, etc.
- Keyboard functional control,
- Control of the Abs enclosure
- Battery check
- Weight control (according to metrological standards DL N.517 and DM N.182).

The instrument must be cleaned with a soft cloth, dampened with water or neutral detergent, avoiding the use of solvents or abrasive substances. Do not use large amounts of water while cleaning the scales, as it may cause damage to the electrical components of the balance. **Always disconnect the scales from the power supply before cleaning.** In case of prolonged use of the instrument, remove the batteries from the terminal and cover the instrumentation to keep it intact. During transport, be careful not to subject the instrument to shocks or excessive mechanical stress. In case of repair or assistance, contact your dealer or an authorized service center by contacting <u>service2@wunder.it</u> or <u>sales@wunder.it</u>

The instrument is sold approved with a first metrological check (plate with M). A subsequent calibration is always necessary if one or more safety stamps are damaged or the display shows abnormal weights.

We recommend that qualified personnel perform maintenance. The WUNDER technical assistance service is at your disposal.

⚠ ATTENTION!

In some countries, calibration can only be performed by an authorized / qualified agent.

Contact your distributor for more information



10. SCRAPPING AND WASTE DISPOSAL

If set aside for a long period, protect those parts which could be damaged due to dust build-up.

Scrapping

When you decide to no longer use this item, we recommend making it unusable. We also recommend making those parts which could be sources of danger harmless

Waste disposal 2012/19/UE

This product complies with the **EU Directive 2012/19/UE**. The symbol of the crossed-out waste bin on the appliance indicates that the product, needing to be treated separately from household waste, at the end of its useful life must be completed in a separate collection facility for electric and electronic appliances or returned to the dealer upon purchase of a new equivalent appliance. The user is responsible for bringing the appliance to an appropriate collection structure at the end of its life. Appropriate separate collection and sending the appliance for recycling, treatment and environmentally compatible waste disposal contributes to avoid possible negative effects on the environment and health and favours the recycling of the materials the product is made of. For more detailed information regarding available collection systems, contact your local waste disposal service or the shop where the product was purchased. As consumers, you are obliged by law to return used or dead batteries. You may deposit old batteries at public collection spots in your town or else with any battery dealer who has placed specific collectors for this purpose. Even when scrapping electric and electronic appliances, they must be removed and deposited in specific collectors.

NOTE: The following symbols indicate the presence of harmful substances.

X

Pb Pb = batteries containing Lead Cd Cd = batteries containing Cadmium Hg Hg = Batteries containing Mercury



Do not throw electric parts and used batteries away with household waste. Dispose of the batteries by means of your closest collection centres.



11. WARRANTY-LIABILITY

WUNDER products are guaranteed under warranty. The warranty is valid for faults or malfunctions of the instrument due to manufacturing or material defects, and WUNDER reserves the right to repair or replace the article. In the event that the repairs of the fault or the delivery of the replacement unit do not have a positive outcome, legal measures are to be considered valid. The term of guarantee is two years, taking effect on the date of purchase. The manufacturer shall not be held liable for damage deriving from any of the following causes: improper or unsuitable storage or use; improper installation or placement in service by the owner or a third party; normal wear and tear from use; changes or modifications; unsuitable or negligent manipulation; excessive use; chemical, electrochemical or electrical interference or interference from humidity; unless such causes may be attributed to negligence on the part of Wunder.

12. IDENTIFICATION PLATES

The year of manufacture is indicated on the applied metrological plate. Ex: **M 19 = 2019, 20 = 2020** ... and so on.



Device class Im with measuring function according to the directive 93/42 CEE



Repetition of the metrological verification

The instrument is sold with a first metrological check (plate with 'M'). We recommend that qualified personnel perform maintenance. The WUNDER technical assistance service is at your disposal.



In some countries, periodic verification can only be performed by an authorized / qualified body. Contact your distributor for further information.