

Stand-on Floor Scale

USER MANUAL **MS6110**



Please keep the instruction manual at hand and follow instruction for use.

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I. Explanation of Graphic Symbols on Label/Packaging

Text/Symbol	Meaning
\triangle	Caution, consult accompanying documents before use
Ā	Separate collection for waste of electrical and electronic equipment, in accordance with Directive 2002/96/EC. Do not dispose of device with everyday waste
•••	Name and address of device manufacturer, and year/country of manufacture
	Carefully read user manual before installation and usage, and follow instructions for use.
<u> </u>	Medical electrical device, Type B applied part
<u> </u>	Medical electrical device, Type BF applied part
REF	Device catalogue number / model number
EC REP	Name and address of authorized representative in the European Union
MD	Device is a medical device. Text indicates device category type
LOT	Manufacturer's batch or lot number for device
SN	Device's serial number
UDI	Device's Unique Device Identifier
е	Verification Scale Interval. Value expressed in units of mass. Used to classification and verification of an instrument.
€ 2460	Device conforms to (EU) 2017/745 Regulation on Medical Devices. Fourdigit number is identifier for medical device Notified Body

	Device complies with EC directives (verified models only)
C€ M200122	 M: Conformity label in compliance with Directive 2014/31/EU for non-automatic weighing instruments 20: Year in which conformity verification was performed and the CE label was applied. (ex: 16=2016) 0122: Identifier for metrology Notified Body
	Device is a Class III scale in compliance with Directive 2014/31/EU (verified models only)
	Name and address of entity importing device (if applicable)
À →文	Name and address of entity responsible for translating Information For Use (if applicable)
CON.	Event counter confirming how many times device has been calibrated (if applicable)
	Device conforms to Taiwan National Communications Commission(NCC) approval
Æ	Device conforms to U.S. Federal Communications Commission regulations
변 <u>M 20</u> 8506	Device complies with UK non-automatic weighing instruments regulations 2016 (verified models only) M: Conformity label in compliance with Non-automatic Weighing instruments Regulations 2016 20: Year in which conformity verification was performed and the UKCA label was applied. (ex: 20=2020) 8506:Identifier for metrology approved body
UK	Device complies with all UK applicable product legislation
$\bigcirc - \bigcirc - \bigcirc +$	Device's polarity of power.

[&]quot;In case of differences, icon on device itself takes precedence"

II. Copyright Notice

Copyright Notice Charder Electronic Co., Ltd.

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Charder Electronic Co., Ltd. No. 103, Guozhong Rd., Dali Dist., Taichung City, 41262 Taiwan

III. Safety Notes

A. General Information

Thank you for choosing this Charder Medical device. It is designed to be easy and straightforward to operate, but if you encounter any problems not addressed in this manual, please contact your local Charder service partner.

Before beginning operation of the device, please read this user manual carefully, and keep it in a safe place for reference. It contains important instructions regarding installation, proper usage, and maintenance.

Intended Purpose

This medical device is designed to be used in accordance with national regulations, to measure weight within specifications, for weight-related usage by professionals.

Clinical Benefit

Measurement results can be used by professionals to diagnose (and monitor) weight-related issues.

Intended medical indications/contraindications

Measurement: patient's body weight. No known contraindications to measurement of body weight.

Intended patient profile

- (a) Age: no restrictions
- (b) Weight: no restrictions within device weight capacity
- (c) Patient Conditions: require measurement of body weight. Able to stand independently without support.

Intended user profile

- (a) At least 20 years old
- (b) Minimum knowledge:
 - To be able to read at a high-school level and understand Arabic numerals (e.g. 1, 2, 3, 4...)
 - Basic hygiene knowledge
 - Trained in device's operation
 - Read the instruction manual

- (c) Language
 - Able to read the language of instruction manual and on-screen instructions
- (d) Qualifications
 - No special certifications or qualifications required

Residual Risk Evaluation

- (a) All foreseeable risks have been evaluated and considered acceptable. Generally speaking, the most likely risk caused by incorrect usage of the device is less accurate measurement (or inability to use device to acquire measurement), which does not pose imminent physical risk to patient or user.
- (b) Benefit-risk ratio is considered acceptable. Stand-on floor scales are an important option for measuring patients. Usage of device is unlikely to result in harm to user or patient.

General Handling

- Device should be placed on stable, flat, solid, non-slippery surface.
- Usage on soft surfaces (ex: carpet) may result in inaccurate results.
- Ensure all parts are properly locked and tightened before operating the device.
- Device is intended to measure one subject at a time.

Safety Instructions

Before putting device into use, please read this user manual carefully. It contains important instructions for installation, usage, and maintenance of device.

The manufacturer shall not be liable for damages caused by failure to heed the following instructions:

- Batteries should be kept away from children. If swallowed, promptly seek medical assistance.
- Expected service life: 5 years.
- Always comply with appropriate regulations when using electrical components under increased safety requirements.
- Improper installation will render the warranty null and void.
- Ensure voltage marked on power supply matches mains power supply.
- The device is intended for indoor use only.
- Observe permissible ambient temperatures for use
- Device meets requirements for electromagnetic compatibility. Do not exceed the maximum values specified in the applicable

standards.

Environmental

 All batteries contain toxic compounds; batteries should be disposed of via designated competent organizations. Batteries should not be incinerated.

Cleaning

- Device surface should be cleaned using alcohol-based wipes.
 Corrosive cleansing liquids should not be used. Pressure-washers should not be used.
- Do not use large amounts of water when cleaning the device, as it may cause damage to the internal electronics.
- Always disconnect device from mains power before cleaning.

Maintenance

Please contact your local Charder distributor for regular maintenance and calibration, regular checking of accuracy is recommended; frequency to be determined by level of use and state of device.

Warranty/Liability

- If Charder is responsible for a fault or defect present upon receipt of the unit, Charder shall either repair the fault, or supply a replacement unit. Should the repairs or replacement delivery fail, statutory provisions shall be valid. The period of warranty shall be two years, beginning on the date of purchase. Please retain your receipt as proof of purchase.
- No responsibility shall be accepted for damage caused through any of the following reasons: unsuitable or improper storage or use, incorrect installation or commissioning by the owner or third parties, natural wear and tear, changes or modifications, incorrect or negligent handling, chemical, electrochemical, or electrical interference, unless damage is attributable to negligence on the part of Charder.
- This device does not contain any user-maintained parts. All maintenance, technicalinspections, and repairs should be conducted by an authorized Charder service partner, using original Charder accessories and spare parts. Charder is not liable for any damages arising from improper maintenance or usage. Dismantlement of the device will void the warranty.

Disposal

This product is not to be treated as regular household waste, but should be taken to a designated collection points for electronics. Further information should be provided by local waste disposal authorities.



- Only the original adapter should be used with the device. Using an adapter other than the one provided by Charder may cause malfunction.
- Do not touch the power supply with wet hands.
- Do not crimp the power cable, and avoid sharp edges.
- Do not overload extension cables connected to the device.
- Route cables carefully, to avoid tripping.
- Keep device away from liquids.
- Do not remove the plug by yanking on the cable.
- Use only a correctly wired (100-240VAC) outlet, and do not use a multiple outlet extension cable.
- Do not under any circumstances dismantle or alter the device, as this could result in electric shock or injury as well as adversely affect the precision of measurements.
- Do not place the device in direct sunlight, or in close proximity to an intense heat source. Excessively high temperatures may damage the internal electronics.

Incident Reporting

Any serious incident that has occurred in relation to the device should be reported to the manufacturer, EU representative (if device is used in EU member state), and competent authority of user/subject's member state.

B. EMC GuidanceandManufacturer's Declaration

Guidance and manufacturer's declaration-electromagnetic emissions

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

Emission test	Compliance	Electromagnetic environment-guid ance
RF emissions CISPR 11	Group 1	The product uses RFenergy only for its internal function. Therefore, its RF emissions are very low and are not likely to cause any interference in nearby electronicequipment.
RF emissions CISPR 11	Class A	The product is suitable for use in all establishments other than domesticand those directly connected to a low voltage
Harmonic emissions IEC 61000-3-2	Class A	power supply network which supplies buildings used for domestic purposes.
Voltage fluctuations /flicker emissions IEC 61000-3-3	Compliance	

Guidance and manufacturer's declaration-electromagnetic immunity

The product is intended for use in the electromagnetic environment specified below. The customer or

the user of the product should assure that it is used in such an environment.

	IEC 60601		Electromagnetic	
Immunity test	test level	Compliance level	environment-quida	
test	ICSI ICVCI		nce	
Electrostati	±8 kV contact	±8 kV contact	Floors should be wood,	
C	±2 kV, ±4 kV, ±8	±2 kV, ±4 kV, ±8	concrete or ceramic tile. If	
discharge(E			floors are covered with	
SD) IEC	kV, ±15 kV air	<u>kV, ±15 kV air</u>	synthetic material, the relative	
61000-4-2			humidity should be	
			at least 30%	
Electrical	+ 2kV for	+ 2kV for	Mains power quality should	
fast	power supply	power supply	be that of a typical	
transient/	lines	lines	commercial or hospital	
burst IEC			environment.	
61000-4-4				
Surge IEC	+ 1kV line(s) to line(s)	_	Mains power quality should	
61000-4-5	+ 2kV line(s) to earth	+ 2kV line(s) to earth	be that of a typical	
			commercial or	
			hospital environment.	
Voltage Dips,	<u>0% UT for 0,5</u>	0% UT for 0,5 cycle	Mains power quality should	
short interruptions	cycle	0% UT for 1 cycle	be that of a typical	
and voltage	0% UT for 1 cycle	700/ LIT/000/ Jim in	commercial or hospital	
variations on	700/ UT/200/ dia	70% UT(30% dip in	environment. If the user of	
power supply input lines IEC	70% UT(30% dip	UT) for 25cycles	the product requires	
61000-4-11	in UT) for 25cycles	00/ LIT for F o	continued operation during	
61000-4-11	0% UT for 5 s	<u>0% UT for 5 s</u>	power mains interruptions, it is recommended that the	
	0% 01 101 5 8		product be powered from	
			anuninterruptible power	
			supply or a battery.	
Power	30 A/m	30 A/m	The product power	
frequency(50,	<u>55. VIII</u>	33.4111	frequency magnetic	
60 Hz)			fields should be at	
magnetic field			levels characteristic of	
IEC 61000-4-8			a typical location in a	
			typical commercial or	
			hospitalenvironment.	
NOTE UT is the a.c. mains voltage prior to application of the testlevel.				

Guidance and manufacturer's declaration-electromagnetic immunity

The product is intended for use in the electromagnetic environment specified below.

The customer or the user of the product should assure that is used in such and environment.

	·		
Immunity test	IEC 60601 test level	Compliance level	Electromagnetic environment-guidance

Conducte d RF IEC 61000-4- 6	3 Vrms 150 KHz to 80 MHz 6 V in ISM bands between 0,15 MHz and 80 MHz 80 % AM at 1 kHz	80 % AM at 1	Portable and mobile RF communications equipment should be used no closer to any part of the product including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter.
Radiated RF IEC 61000-4- 3	3 V/m 80MHz to 2,7 GHz	kHz 3 V/m 80MHz to 2,7 GHz	Recommended separation distance: $d = 1, 2 \sqrt{p}$ $d = 1, 2 \sqrt{p}$ 80MHz to 800 MHz $d = 2, 3 \sqrt{p}$ 800MHz to 2,7GHz Where P is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer and d is the recommended separation distance in metres (m). Field strengths from fixed RF transmitters, as determined by an electromagnetic site survey, ^a should be less than the compliance level in each frequency range. Interference may occur in the vicinity of equipment marked with the following symbol:

NOTE1 At 80 MHz and 800 MHz, the higher frequency rangeapplies.

NOTE2

Theseguidelinesmaynotapplyinallsituations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.

- a Field strengths from fixed transmitters, such as base stations for radio (cellular/cordless) telephones and land mobile radios, amateur radio, AMandFM radiobroadcast and TV broadcast cannot be predicted theoretically with accuracy. To assess the electromagnetic environment due to fixed RF transmitters, an electromagnetic site survey should be considered. If the measured field strength in the location in which the product is used exceeds the applicable RF compliance level above, the product should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as re-orienting or relocating the product.
- b Over the frequency range 150 kHz to 80 MHz, field strengths should be less than 3V/m.

Recommended separation distance between portable and mobile RF communications equipment and the product

The product 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 preventelectromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and the product as recommended below, according to the maximumoutput power of the communications equipment.

Rated maximum	Separation distance according to frequency of transmitter m			
output power of transmitte r W	150 kHz to 80 MHz d =1,2√P	80 MHz to 800 MHz d =1,2 \sqrt{P}	800 MHz to 2,7 GHz d =2,3 \sqrt{P}	
0,01	0,12	0,12	0,23	
0,1	0,38	0,38	0,73	
1	1,2	1,2	2,3	
10	3,8	3,8	7,3	
100	12	12	23	

For transmitters rated at a maximum output power not listed above, the recommended separation distance d in metres (m) can be estimated using the equation applicable to the frequency of the transmitter, where p is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer.

NOTE1 At 80 MHz and 800 MHz, the separation distance for the higher frequency range applies.

NOTE2 These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.

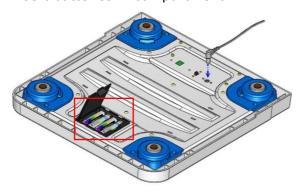
IV. Installation

A. Assembly

Device does not require assembly, and can be used once power is supplied.

B. Inserting Batteries

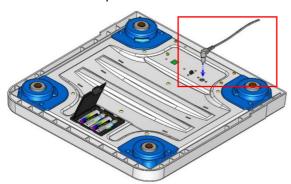
Insert batteries in compartment



2. Turn on power to confirm that battery is correctly installed.

C. Using Adapter

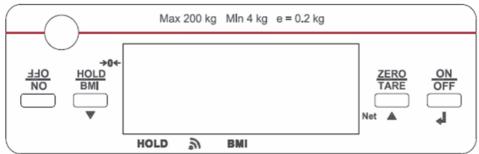
1. Connect adapter to device before connecting to mains power supply



2. Disconnect adapter from mains power supply before unplugging adapter pin from device.

V. Indicator

A. Indicator and Key Functions



Key Function

- 1. ON/OFF: Power button. Weighing results will face patient when device is initially powered on.
- 2. HOLD/BMI: Press once to Hold (determine stable weighing value used when weight is unstable). Press and hold for 3 seconds to enter Body Mass Index (BMI) calculation mode.
- 3. ZERO/TARE: Press to Zero or Tare weight.

B. Display layout

1. **O**: Weight is stable

2.

: Weight is negative

3. +O+: Device is at zero

5. HOLD: Hold function is active

6. BMI: BMI function is active

7. Net: Current result is net weight

VI. Using Device

A. Basic Operation

Switch on the device using **[ON/OFF]** key. (To turn off device, press and hold **[ON/OFF]** key for 3 seconds) The device will automatically perform self-calibration, displaying software version.

Once "0.00 kg" appears on indicator, device is ready for measurement.

Note: If "0.00 kg" does not display on indicator, press **[ZERO/TARE]**key to zero the device. This function can be used for weight within $\pm 2\%$ of full capacity.

Guide subject to stand on device. After the weight has stabilized, the "stable" symbol will appear on indicator.

Note: If subject's weight exceeds scale capacity (including tare), indicator will display "Err" prompt due to overload.

B. Hold

The hold function determines average weight, designed to be used if subject's weight will not stabilize (ex: an active child).

Note: if fluctuation is too severe, average weight determination will be difficult and hold may not function correctly

- 1. Switch on the device normally.
- 2. Press the **[HOLD/BMI]** key. "HOLD" will be displayed on the indicator.
- 3. Guide subject to stand on device.
- 4. After a few seconds, the average weight will be displayed on the indicator. This weight will be locked at this point, subject can leave measurement platform.
- 5. To release the locked weight, press the **[HOLD/BMI]** key again to return to the device to normal mode. (arrow next to "HOLD" on indicator will disappear)

Note: Hold function can be activated before or after subject stands on device. However, if subject finds it difficult to stand still, we recommend activating Hold after subject stands on device. Hold function will not function under 2 kg.

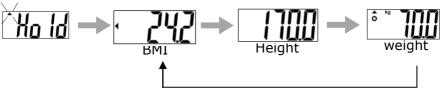
C. Tare

The tare function allows the user to deduct the weight of objects from the device's measurement result.

- 1. Place object that needs to be tared onto measurement platform.
- 2. Press **[ZERO/TARE]**key after stable symbol appears on indicator. Display will indicate "0.00 kg".
- 3. Guide subject (plus tared object) to stand on device. Conduct measurement.
- 4. To clear tare value, remove all objects from measurement platform, and press [**ZERO/TARE**]key.

D. Body Mass Index (BMI)

- 1. In normal mode, press and hold the **[HOLD/BMI]** key to enter BMI mode.
- 2. Display will show last input height. Left-most digit will flash.
- 3. Adjust height value using **[TARE]**(increase ↑) and **[HOLD/BMI]** (decrease ↓) keys. (press and hold to speed up)
- 4. Press [ON/OFF] key to confirm height.
- 5. Proceed to weigh subject as usual. Indicator will display weight and BMI after measurement.
- 6. Press **[HOLD/BMI]** to return to normal mode.



Category	BMI (kg/m²)	Risk of obesity-related disease
Under	< 18.5	Low
Normal	18.5-24.9	Average
Over	24.9-29.9	Slightly Increased
Obese I	30.0-34.9	Increased
Obese II	35.0-39.9	High
Obese III	> 40	Very High

(World Health Organization adult BMI standards)

VII. Device Setup

When the device is switched on, press and hold the **[ZERO/TARE]** key for 6 seconds, until the display shows the "SETUP", followed by "A_OFF" (first option in setting menu).

In device setup:

[HOLD/BMI] select menu option [ZERO/TARE] confirm selection



Auto-off: Instruct device to shut off automatically after a certain period of time.

Auto off options: 60 sec / 120 sec / 180 sec / 240 sec / 300 sec / off

Press [HOLD/BMI] to toggle between time options, and [ZERO/TARE] to confirm selection.



Adapter Auto-Off: Select whether auto-off will be activated when device is plugged into adapter.

Press [HOLD/BMI]to toggle between ON/OFF, and [ZERO/TARE] to confirm selection.



Hold Stop: When Hold Stop is "on", Hold will deactivate after subject leaves measurement platform.

Press [HOLD/BMI] to toggle between time options, and [ZERO/TARE] to confirm selection.



Date Setting: press [ZERO/TARE] to increase. Press [HOLD/BMI] to move to next digit. Press [ON/OFF] to confirm setting and move to next input. (e.g. after inputting Year, press [ON/OFF] to input month).

Order is YYYY.MM.DD.HH.DD (Year, Month, Day, Hour, Minute) (24 hour format).



Bluetooth (optional): If device has Bluetooth module installed, Bluetooth function can be turned on or off.

Press [HOLD/BMI] to toggle between ON/OFF, and [ZERO/TARE] to confirm selection.



Wi-Fi (optional): If device has Wi-Fi module installed, Wi-Fi function can be turned on or off.

Press [HOLD/BMI]to toggle between ON/OFF, and [ZERO/TARE] to confirm selection.



Wi-Fi Setting (optional): If device has Wi-Fi module installed, this option will appear. If "Auto" is selected, weight measurement will be automatically sent to connected printer or device. If "PKEY" is selected, transfer will occur manually only after **[PRINT]** key is pressed.

VIII. Wireless Connection

If the device has the wireless or bluetooth module installed, the indicator can transmit measurement results wirelessly. Please see Charder wireless or bluetooth software instructions for details.

IX. Troubleshooting

Product Defects

Charder's warranty is effective for the original purchaser of this device, subject to the terms and conditions listed in the Warranty Program & Return Policy.

- 1. If Charder is responsible for a fault or defect present upon receipt of the unit, Charder shall either repair the fault, or supply a replacement unit. Should the repairs or replacement delivery fail, statutory provisions shall be valid. The period of warranty shall be two years, beginning on the date of purchase. Please retain your receipt as proof of purchase.
- 2. No responsibility shall be accepted for damage caused through any of the following reasons: unsuitable or improper storage or use, incorrect installation or commissioning by the owner or third parties, natural wear and tear, changes or modifications, incorrect or negligent handling, chemical, electrochemical, or electrical interference, unless damage is attributable to negligence on the part of Charder.

If device is not covered under warranty, a service maintenance charge will apply, plus cost of replacement parts.

Before contacting your local Charder distributor for repair service, we recommend considering the following troubleshooting procedures:

Self-inspection

1. Device will not power on

- If battery power is depleted, replace with new batteries
- If batteries are not used, check if the AC power adapter is plugged into the device properly. Check if power adapter is plugged into mains properly

2. Indicator showing "0000" ZERO SPAN out of range

- Interference due to factors such as RF disturbance or ground vibration. Relocate device to location without interference and try again
- Unstable platform feet adjust platform feet according to bubble level indication (clockwise to retract, counter-clockwise to extend) and try again
- External objects interfering with measurement platform. Clear platform of objects and try again

- Device may not function properly on soft surfaces such as carpets or lawns. Relocate device to location with solid, stable floor
- If the steps above cannot resolve the problem, re-calibration may be required to correct weighing accuracy

3. Connection failure for data transmission to PC or printer

- Ensure wires are connected correctly between indicator and PC or printer
- Ensure printer is supplied with power. Ensure PC software is set up properly as indicated in this manual

Distributor support required

If the following errors occur, we recommend contacting your local Charder distributor for repair or replacement services:

1. Device will not power on

- Faulty on/off key
- Broken or damaged wires causing short circuit or faulty connection
- Safety fuse burnout
- Faulty AC Adapter

2. Indicator damage

- Possible hardware defects include: uneven brightness in LCD screen, blurred text, smeared rainbow screen, incorrect decimal display
- Unable to save or read data
- Indicator shows "ERRL" after device is switched on
- Keys not responding
- Buzzer malfunction

Error Messages

Error Messages	В	A 1:
Error Message	Reason	Action
LobAt	Low battery warning Voltage of battery is too low to operate device	Replace batteries, or plug in AC adapter
Err	Overload Total load exceeds device's maximum capacity	Reduce weight on measurement platform and try again
Err5	Counting Error Signal from loadcells too high or low	Error normally caused by faulty loadcell or wiring. Please contact distributor
00000	Zero count over calibration zero range +10% while power on	Remove weight from device and try again. If error persists, please contact distributor
00000	Zero count under calibration zero range -10% while power on	Remove weight from device and try again. If error persists, please contact distributor
ErrRd	Program Error Fault with device software	Please contact distributor

X. Product Specifications

A. Device Information

5.4		1400	440	
Model		MS6110		
Weight			200 kg x 0.2 kg	
Measurement	Accuracy	±1	.5e	
	OIML	Clas	s III	
	LCD Screen	1.0-inch LCD so	creen (5 digits)	
Dimensions (Standard)	Overall	345(W) x 350(D) x 65(H) mm	
(Device Weight	2.7 kg		
Key Functions		On/Off, Zero/Tare, Hold/BMI		
Data Trar	smission	Wireless module (optional)		
		NOTE : Device should be connected to network by qualified distributors only.		
Power	Supply	4 AA batteries / adapter		
Operation E	nvironment	+5°C~+35°C		
		15% / 85% RH 700 hPa ~1060 hPa		
Optional A	ccessories	Thermal Printer		
Standard A	ccessories	User manual x1, Power Adapter x1, USB cable x1		

B. Power Adapter Standards



The device is only compatible with the power adapters specified in the dashed block below.

AMP VOLTAGE	DRAWING NO.	CE APPROVED TYPE NO. / MODEL NO.	TYPE	Adapter plug
12V 2A	CD-AD-00041	UES24LCP-120200SPA	US	
	CD-AD-00041	UES24LCP-120200SPA	EU	
	CD-AD-00041	UES24LCP-120200SPA	UK	90 - degree
	CD-AD-00041	UES24LCP-120200SPA	AU	

Notes		

Notes		

XI. Declaration of Conformity

This product has been manufactured in accordance with the harmonized European standards, following the provisions of the below stated directives:

C € 2460	(EU) 2017/745 Regulation on Medical Devices	
CE M year	2014/31/EU Non-automatic Weighing Instruments Directive (OIML models only)	

RoHS Directive 2011/65/EU and Delegated Directive (EU) 2015/863

Radio Equipment Directive 2014/53/EU

(applicable if wireless module is used)

Part 15 of the Federal Communications Statement Rules

This device may not cause harmful interference.

This device must accept any interference received, including interference that may cause undesired operation.

Please see separate document showing on sticker of device for above markings.

Authorized EU Representative:



Manufactured by:



Charder Electronic Co., Ltd. No.103, Guozhong Rd., Dali Dist., Taichung City 41262, Taiwan

CD-IN-1386 [152490] 08/2024