

Chair Scale

USER MANUAL **MS5480**



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 |
|-------------------|---|
| \wedge | Caution, consult accompanying documents before use |
| X | 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 |
| E | Carefully read user manual before installation and usage, and follow instructions for use. |
| Ŕ | Medical electrical device, Type B applied part |
| × | 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. |
| CE 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)

CEM200122
 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) |
| A ⇒₹ | 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) |
| NI | Device conforms to Taiwan National Communications Commission(NCC) approval |
| FC | Device conforms to U.S. Federal Communications Commission regulations |
| ĽK <u>M</u> 208506 | 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 CA | Device complies with all UK applicable product legislation |
| \ominus \oplus \oplus | Device's polarity of power. |

"In case of differences, icon on device itself takes precedence"

Copyright Notice Charder Electronic Co., Ltd.

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Charder reserves the right to correct misprints in the manual without prior notice, and modify the exterior of the device for quality purposes without customer consent.



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

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. Unable 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. Chair scales are an important option for measuring patients. Usage of device is unlikely to result in harm to user or patient.

General Handling

- Ensure all parts are properly locked and tightened before operating the device.
- Measurement accuracy requires the subject's feet, back, and head to be straightly aligned. Please note that height can vary throughout the day
- **CAUTION**: Do not use next to equipment that may cause electromagnetic or other types of interference.

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:

- The device has an expected service life of 5 years when correctly handled, serviced, and periodically inspected in accordance with manufacturer's instructions.
- Improper installation will render the warranty null and void.
- Observe permissible ambient temperatures for use

Cleaning

Device surface should be cleaned using alcohol-based wipes.

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

- The period of warranty shall be eighteen(18) months, 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.

Measurements for physically disabled persons.

- Physically disabled persons should not attempt to take measurements alone, but instead should have their caretakers assist them in using the device.
- Footrest can only be used when subject is sitting in chair. To avoid injury, subject should refrain from standing on footrest, as device may tip over if used incorrectly.





The collapsible frame should be handled with caution. Keep fingers, hands, or other parts of the body clear when folding or unfolding frame, to avoid injury.



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 Guidance and Manufacturer'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 o | f the product should | assure that it is used in such an environment. |
| Emission test Compliance | | Electromagnetic |
| Emission test | Compliance | environment-guidance |
| RF emissions CISPR 11 | Group 1 | The product uses RF energy only for its internal function. Therefore, its RF emissions are very low and are not likely to cause any interference in nearby electronic equipment. |
| RF emissions CISPR 11 | Class A | The product is suitable for use in all establishments other than domestic and those |
| Harmonic emissions IEC 61000-3-2 | Class A | directly connected to a low voltage 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.

| The customer or the user of the product should assure that it is used in such an environment. | | | |
|---|--|--|--|
| Immunity test | IEC 60601 test level | Compliance level | Electromagnetic environment-guidance |
| Electrostatic discharge(ESD) IEC 61000-4-2 Electrical fast | $\frac{\pm 8 \text{ kV contact}}{\pm 2 \text{ kV}, \pm 4 \text{ kV}, \pm 8}$ kV, ±15 kV air ± 2kV for | $ \pm 8 \text{ kV contact} \\ \pm 2 \text{ kV, } \pm 4 \text{ kV, } \pm 8 \\ \text{kV, } \pm 15 \text{ kV air} \\ \pm 2 \text{ kV for} $ | Floors should be wood, concrete or ceramic tile. If floors are covered with synthetic material, the relative humidity should be at least 30% Mains power quality should |
| transient/burst IEC 61000-4-4 | power supply lines | power supply lines | be that of a typical commercial or hospital environment. |
| Surge IEC 61000-4-5 | <u>+</u> 1kV line(s) to line(s) <u>+</u> 2kV line(s) to earth | \pm 2kV line(s) to earth | be that of a typical commercial or hospital environment. |
| Voltage Dips, short interruptions and voltage variations on power supply input lines IEC 61000-4-11 | <u>0% UT for 0,5</u> <u>cycle</u> <u>0% UT for 1 cycle</u> <u>70% UT(30% dip</u> in UT) for 25cycles <u>0% UT for 5 s</u> | <u>0% UT for 0,5 cycle</u> <u>0% UT for 1 cycle</u> <u>70% UT(30% dip in</u> <u>UT) for 25cycles</u> <u>0% UT for 5 s</u> | Mains power quality should be that of a typical commercial or hospital environment. If the user of the product requires continued operation during power mains interruptions, it is recommended that the product be powered from an uninterruptible power supply or a battery. |
| Power frequency (50, 60 Hz) magnetic field IEC 61000-4-8 | <u>30 A/m</u> | 30 A/m | The product power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercial or hospital environment. |
| NOTE UT is the a.c. mains voltage prior to application of the test level. | | | |

| 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 |
| Conducted RF IEC 61000-4-6 | 3 Vrms 150 KHz to 80 MHz | 3 Vrms 150 KHz to 80 MHz | Portable and mobile RF communications equipment should be used no closer to any part of the product including cables, |
| | 6 V in ISM bands between 0,15 MHz and 80 MHz 80 % AM at 1 kHz | 6 V in ISM bands between 0,15 MHz and 80 MHz 80 % AM at 1 kHz | 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 <u>80MHz to 2,7 GHz</u> | 3 V/m <u>80MHz to 2,7 GHz</u> | 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 <i>P</i> is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer and <i>d</i> 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 ^b |
| | | | Interference may occur in the vicinity of equipment marked with the following symbol: |

NOTE1 At 80 MHz and 800 MHz, 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.

- a Field strengths from fixed transmitters, such as base stations for radio (cellular/cordless) telephones and land mobile radios, amateur radio, AM and FM radio broadcast and TV broadcast cannot be predicted theoretically with accuracy. To assess the electromagnetic environment due to fixed 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 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.

| Rated maximum | Separation distance according to frequency of transmitter | | | |
|--------------------------------|---|---------------------------------------|---|--|
| output power of transmitter | | m | | |
| W | 150 kHz to 80 MHz d =1,2√ <i>P</i> | 80 MHz to 800 MHz d =1,2√ <i>P</i> | 800 MHz to 2,7 GHz d =2,3√ <i>P</i> | |
| 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.

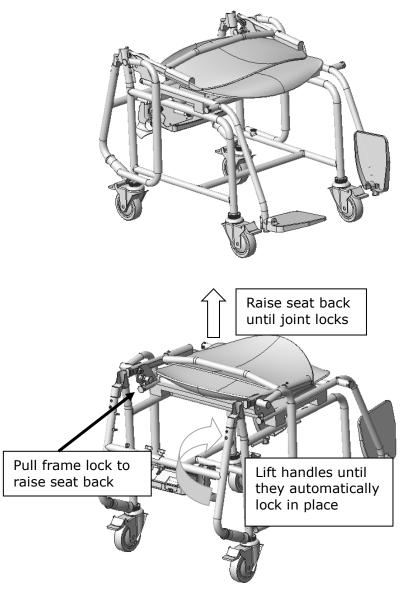
NOTE1 At 80 MHz and 800 MHz, the separation distance for the higher frequency range applies.

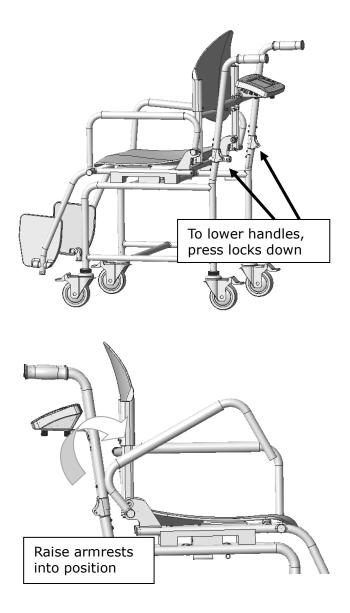
NOTE2 These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.

IV. Installation A. Unfolding device

Device should arrive in folded form.

Rotate footrests into position.







Engaging wheel brakes

Wheel brakes must be engaged prior to measurement. Press down on brakes to engage.



B. Using adapter and charging battery

The rechargeable battery should be recharged at least once every 3 months, regardless of if the device has been used. Battery can be charged by plugging device's exclusive adapter into AC Connector Port.

After a long period in storage (e.g. >3 months), the battery should run a full cycle (charge/discharge) to allow it to restore full capacity.

Ensure rechargeable battery housing is installed and inserted properly into the compartment.



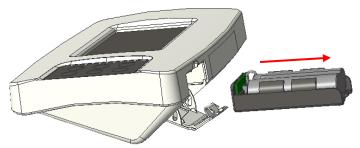
If prompt displays on the LCD, please charge battery promptly to avoid battery damage.

C. Replacing Rechargeable Battery Pack

1. Open battery housing cover



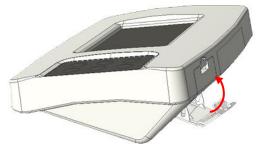
2. Accessing batteries



3. Place new battery pack into housing, and insert into indicator



4. Close battery housing compartment cover. Turn on power to confirm that battery is correctly installed.



D. Optional Accessories

This device is compatible with a variety of accessories. Please consult their respective user manuals for assembly instructions:

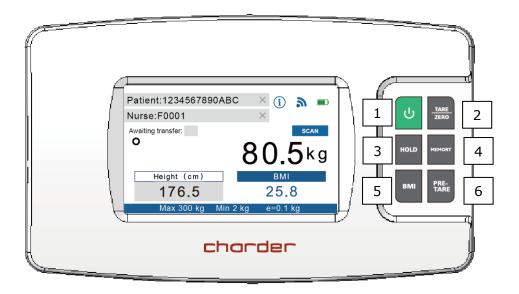
a. Barcode Scanner

■ AS-9300 Infrared Barcode Scanner

V. Indicator

A. Indicator and Key Functions

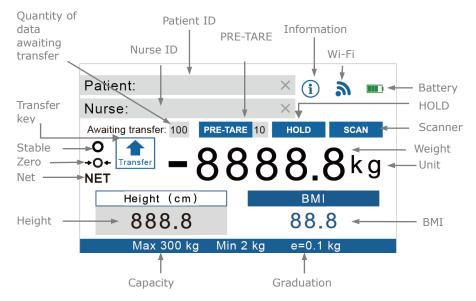
I



Key Function

| С | POWER | Power on or power off. |
|---------|-------|---|
| TARE/ZE | RO | Reset display to 0.0 kg display. Press and hold for 6 seconds to enter device settings. |
| HOLD | | Determine stable weighing value - used when weight is unstable. |
| MEMOR | Y | Save pre-tare values (up to 10 sets can be stored in device memory). |
| BMI | | Calculation of Body Mass Index. |
| PRE-TAR | RE | Pre-tare the known weight of an object (ex: chair) before beginning measurement. |

B. Display layout



Definitions

Quantity of data awaiting transfer: If device is not connected wirelessly, measurement results will be temporarily stored in device. Once device is connected, operator can press **Transfer** to send results wirelessly. After transfer is complete, number will revert to "0"

PRE-TARE: If Pre-Tare function is active, this indicates which pre-tare value is being used.

HOLD: Will appear if Hold is active. (Hold needs to be activated in order to save and transfer results)

SCAN: Will appear if compatible barcode scanner is plugged into device

Transfer: After measurement is completed, height/weight result can be transferred wirelessly (if Hold is active). Press **[Transfer]** to send results.

Wi-Fi: Indicator will reflect current Wi-Fi connectivity status.

disconnected



VI. Basic Operation

1. Switch on the device using ${{f U}}$ key. The device will automatically perform self-calibration

2. Once "0.0" appears on indicator, device is ready for use

NOTE: If "0.0" does not display on indicator, press **[TARE/ZERO]** key to zero the device.

A. Basic Operation

Direct Wi-Fi data transfer

NOTE: If results do not need to be transferred after measurement, this step can be skipped.

1. The device acts as an Access Point

that can be connected to via Wi-Fi. To

ensure that the phone/tablet/PC

Nurse:F0001 Awaiting transfer: Band Solution Height (cm) 176.5 Max 300 kg Min 2 kg e=0.1 kg

Patient:1234567890ABC

G

connects to the correct device, first identify the device's MAC Address by clicking

| Back Information | | |
|------------------|-------------------|--|
| MODEL N | MS4980 | |
| SERIAL N | T20000123 | |
| Time | 2021/03/04 09:40 | |
| UP Code | UP-00126 | |
| FW Ver. | V1.03 | |
| Wifi MAC | 70:1d:08:06:2c:36 | |
| Capacity | 300x0.1kg | |
| BMP_Ver. | 201120 | |
| G in Time | 1 | |

2. The "name" of the device's Wi-Fi Access Point will be "DP4800_(MAC Address)" The default password to connect to the device is "00000000"

NOTE: The Model No. displayed in Information will vary depending on the

device model.

3. After the phone/tablet/PC is connected to the device, the wireless

symbol will change from



4. Device is now ready to send results wirelessly to phone/tablet/PC.5. Before or after measurement, press the **[HOLD]** key. "HOLD" will be displayed on the indicator. HOLD must be active, to transfer results.

NOTE: by default, patient ID, weight, and height must be filled in to transfer results. Otherwise, **[Transfer]** button will not appear. To allow transfer of "incomplete" results, please change settings (press and hold **[TARE/ZERO]** key for 6 seconds to enter settings).

B. Weight measurement

- 1. Engage wheel brakes, raise footrests, and raise armrests.
- 2. Turn on device

3. Guide subject to sit on chair.Place subject's feet upon footrests. After a few seconds, the average weight will be displayed on the indicator. This weight will be locked - at this point, subject can stand up from chair. 4. If BMI calculation is unnecessary, press **[Transfer]** button to send results wirelessly. If device is not currently connected, results will temporarily be stored in device memory (number of records saved indicated by 'Awaiting transfer'). After transfer is complete, number will revert to "0"

C. BMI calculation



CLEAR

1.Press the **[BMI]** key to enter BMI mode.

cm

3

6

9

Ente

5

8

0

4

7

~

2. Enter height using numeral keys (ex: to input 170 cm, press 1-7-0-0). Press **[CLEAR]** key to re-input.

3. After inputting height, press **[Enter]** to confirm.

4. Proceed to weigh subject as usual. Indicator will display weight, height, and BMI.

5. To transfer results, ensure that HOLD is active, and press **[Transfer]** button to send results wirelessly. If device is not currently connected, results will temporarily be stored in device memory (number of records saved indicated by 'Awaiting transfer'). After transfer is complete, number will revert to "0"

| 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)

D. 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 chair.

2. Press **[TARE/ZERO]** key after stable symbol appears on indicator. Display will indicate "0.0".

3. Guide subject (plus tared object) to be weighed upon chair. Conduct measurement.

4. To clear tare value, remove all objects from chair, and press **[TARE/ZERO]** key.

E. Pre-Tare

The Pre-Tare function is used to subtract the known weight of a substance prior to weighing. The device can store 10 sets of pre-tare values in memory. Once pre-tare weights have been stored, they can be recalled by pressing the **[MEMORY]** key.

Input Pre-Tare Value

| DESCRIPTION | EXAMPLE |
|--|--|
| Press [PRE-TARE] key. Input pre-tare weight value, starting from the left | Back Pre-tare 0.0, kg 1 2 3 CLEAR 4 5 6 7 8 9 ← 0 Enter |
| Enter pre-tare weight using 0~9 keys. Ex: to pre-tare 5.0 kg of weight, press 0-0-5-0. Ex: to pre-tare 13.5 kg of weight, press 0-1-3-5. Press [Enter] key to confirm the pre-tare weight. | Back Pre-tare 0 1 2 3 CLEAR 4 5 6 7 8 9 ← 0 Enter Back Pre-tare Back Pre-tare 1 2 3 CLEAR 5 0 [6] 7 8 9 CLEAR 7 8 9 € 0 Enter |
| Device will return to measurement mode. Indicator will display minus sign to the left of pre-tare weight value. | Patient: Image: Constraint of the second |

Save a Pre-Tare value

| DESCRIPTION | EXAMPLE |
|---|--|
| Press and hold [MEMORY] key for 3 seconds. Input the number for this pre-tare setting (between 01-10). Ex: To save memory set 3, press 0-3. | Back MEMORY 0.0 kg 1 2 3 GLEAR 4 5 6 MEMORY 3 |
| Press the weight value box on the screen (marked in the red box to the right) Enter pre-tare weight using 0~9 keys. Ex: to pre-tare 13.5 kg of weight, press 0-1-3-5. | Back MEMORY 0 1 2 3 CLEAR 4 5 6 MEMORY 3 7 8 9 ← 0 Enter |
| Press [Enter] key to store pre-tare weight (saved to memory set 3, in this example) | Back MEMORY 123 123 456 789 ← 0 |
| PRE-TARE ID will show Pre-Tare memory set 3 as active, and deduct the pre-tare value (13.5 kg, in this example) from the displayed weight result. | Patient: Nurse: Awaiting transfer: PRE-TARE 3 PRE-TARE 3 - 1 3.5 k g Height (cm) Max 300 kg Min 2 kg e=0.1 kg |

Recall Pre-Tare Value

| DESCRIPTION | EXAMPLE |
|--|---|
| Press [MEMORY] key to toggle between Pre-Tare settings saved in device (between 01-10). | Patient: × (i) 為 ■ Nurse: × |
| NOTE : to add new Pre-Tare value, please press and hold [MEMORY] key for 3 seconds to enter settings (see "Save a Pre-Tare value" above) | Awaiting transfer: PRE-TARE 3 - 13.5 kg Height (cm) BMI Max 300 kg Min 2 kg e=0.1 kg |

Press and hold **[TARE/ZERO]** key for 6 seconds to enter General Setting mode.

| EXIT General | | | |
|---------------|------|------------------|------|
| Auto Off Time | 180s | G-Compensation | |
| Backlight | High | H.M. Calibration | |
| Buzzer | On | Height Capacity | High |
| Data Transfer | On | URL Host | |
| Date/Time | | Auto Hold | On |
| Wifi Setting | | Auto Transfer | On |

Press menu options on the touchscreen to adjust settings.

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

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

Backlight: adjust backlight brightness.

Options: Low / Mid / High

Buzzer: When function is turned on, beeping noise will be made when: indicator is turned on, keys are pressed, and weight is stable.

Options: Enable / Disable

Data Transfer: If enabled, all data fields (patient ID, weight, height) need to be completed to transfer data. If fields are incomplete (ex: only weight, no height or ID), data will not be transferred.

Options: Enable / Disable

Date/Time: Set device time. (Format: YYYY/MM/DD HH:M)

WiFi Settings: Send results via direct transfer or via network (set Access Point if selected)

G-Compensation: Authorized distributor can adjust gravity compensation value (password required)

H.M. Calibration: Calibrate ultrasonic height stadiometer

Height Capacity: Default setting is "High". This setting should be adjusted by distributor only - changing it may affect measurement accuracy.

URL Host: Set IP address (ex: 192.168.0.1). Please note that if server is restarted, another IP may be automatically assigned. If IP change occurs, please re-input correct IP once more to complete settings.

Auto Hold: Determine if hold function will be automatically activated on start-up

Auto Transfer: Determine if results will be automatically transferred after measurement completion

VIII. Connecting scale to receiving device

The scale is designed to transfer results wireless to receiving device. Please consult instruction manual for receiving device.

Connection directly to Electronic Medical System should be conducted by qualified distributors/administrators only.

IX. Troubleshooting

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 power adapter is plugged into the device properly. Check if power adapter is plugged into mains properly

2. Indicator showing "00000" 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

Error Messages

| Error Messages | |
|--|--|
| Error Message | Action |
| Low battery Please replace new batteries or plug the AC adaptor for operation. | Please charge battery using adapter, or replace battery |
| Overload Please reduce the loading and try again. | Maximum weight exceeded. Reduce weight on platform before attempting measurement |
| Loadcell error Please contact your nearest Authorized Dealer for further technician service & repair. | If problem persists, please contact distributor |
| Zero count over calibration zero range Plese re-calibrate this instrument. | Re-calibration may be required. If problem persists, please contact distributor |
| Zero count under calibration zero range Plese re-calibrate this instrument. | Re-calibration may be required. If problem persists, please contact distributor |
| ADC error Please contact your nearest Authorized Dealer for further technician service & repair. | If problem persists, please contact distributor |

| Model | | MS5480 | |
|--|--------------------------------|---|--|
| Display | | DP4800 | |
| | Capacity | 250kgx 0.1kg 550 lb x 0.2 lb | |
| Weight | Accuracy | ±1.5e | |
| Measurement | Unit | kg / lb | |
| | LCD Screen | Color LCD touchscreen | |
| | Overall | 590(W) x 1020(D) x 960(H) mm | |
| Dimensions | Seat | Height: 470mm Width: 460 mm Back Height: 480 mm | |
| Armrest | | Height: 660 mm | |
| Device | Weight | 17.6 kg | |
| Key Fu | nctions | Power, Tare/Zero, Hold, Memory, BMI, Pre-Tare | |
| Data Transmission | | USB, Wireless NOTE : Device should be connected to | |
| | | network by qualified distributors only | |
| Power Supply Rechargeable battery pack / a | | Rechargeable battery pack / adapter | |
| Operation Environment | | +5℃~+35℃ 15%~85% RH 700 hPa ~1060 hPa | |
| Standard A | ccessories | User manual*1, Power Adapter*1 | |
| Optional A | al Accessories Barcode Scanner | | |

B. Power Adapter Standards



Only the original adapter should be used with the device. Using an adapter other than the one provided by Charder may cause malfunction.

Amp Voltage: 5V/2A Drawing No: CD-AD-00023

| AMP VOLTAGE | DRAWING NO.: | CE APPROVED TYPE NO. / MODEL NO.: | ТҮРЕ | Adapter plug |
|----------------|-----------------|--------------------------------------|------|-----------------|
| 5V 2A | AD-00023 | UES12LCP-050200SPC | US | |
| 5V 2A | AD-00023 | UES12LCP-050200SPC | EU | |
| 5V 2A | AD-00023 | UES12LCP-050200SPC | UK | |
| 5V 2A | AD-00023 | UES12LCP-050200SPC | AU | |

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XI. Declaration of Conformity

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

| CE 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:



Obelis s.a. Bd Général Wahis, 53 B-1030 Brussels Belgium

Manufactured by:



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

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