

USER MANUAL **HM200P**

Portable Height Stadiometer

Explanation of Text/Symbols on Device Label/Packaging

Text/Symbol	Meaning
\triangle	Caution, consult accompanying documents before use
	Separate collection for waste of electrical and electronic equipment, in accordance with Directive 2002/96/EC. Do not dispose of device with everyday waste
444	Name and address of device manufacturer, and year/country of manufacture
	Carefully read user manual before installation and usage, and follow instructions for use.
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
€ 2460	Device conforms to 93/42/EEC as amended by 2007/47/EC Medical Device Directive. Four digit number is identifier for medical device Notified Body
	Name and address of entity importing device (if applicable)
A →文	Name and address of entity responsible for translating Information For Use (if applicable)

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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 height within specifications, for height-related usage by professionals.

Clinical Benefit

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

General Handling

- Device should be placed on stable, flat, solid, non-slippery surface.
- Ensure all parts are properly locked and tightened before operating the device.

Safety Instructions

■ The device has an expected service life of 5 years when correctly handled, serviced, and periodically inspected in accordance with manufacturer's instructions.

Cleaning

■ Device surface should be cleaned using alcohol-based wipes. Corrosive cleansing liquids should not be used.

Warranty/Liability

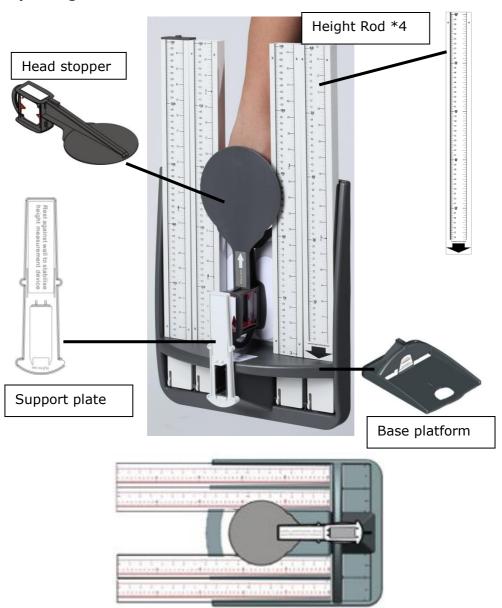
- The period of warranty shall be eighteen (18) months, beginning on the date of purchase. Please retain your receipt as proof of purchase.
- No responsibility shall be accepted for damage caused through any of the following reasons: unsuitable or improper storage or use, incorrect installation or commissioning by the owner or third parties, natural wear and tear, changes or modifications, incorrect or negligent handling.

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.

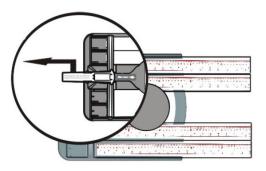
II. Installation

Unpacking the HM200P

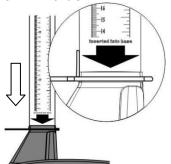


Assembly Instructions

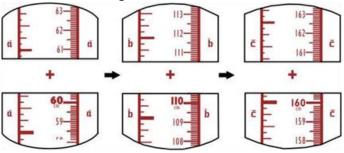
1. Remove support plate



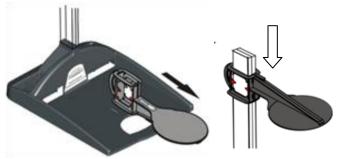
2. Insert Height Rods into base platform in order



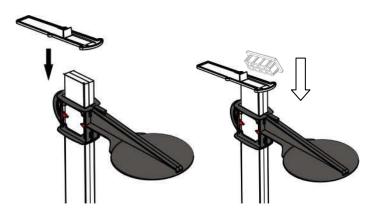
3. Ensure that height rods are inserted in the correct order



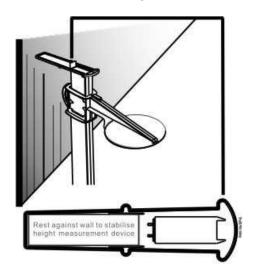
4. Mount head stopper onto height rod



5. Place support plate on top of height rod

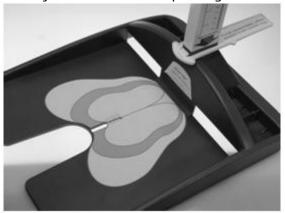


6. Place HM200P against the wall for stability.

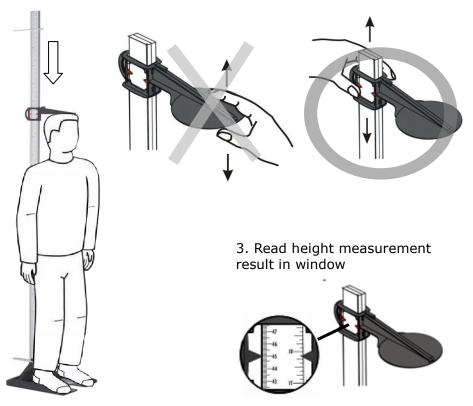


III. Using Device

1. Subject should stand up straight with feet within the foot markings



2. Subject's back should tough the height measurement rod slightly. Lower stopper until it touches top of subject's head



IV. Product Specifications

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Model		НМ200Р			
Height Measurement	Range	14-205 cm 5.5-81 in			
	Graduation	1 mm 1/16 in			
	Accuracy	±10 mm			
Dimensions	Overall	350(W) x 400(D) x 2430(H) mm			
Device Weight (approximate)		2.0 kg / 4.4 lb (carry case 0.6 kg / 1.3 lb)			
Operation Temperature & Humidity		5°C~35°C			
Optional Accessories		AR-2481, AR-00001 carrying bag			
Standard Accessories		User manual x1			

Notes			
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V. Declaration of Conformity

Manufacturer hereby declares that this product is in conformity with the regulations and standards outlined in the following directives:

93/42/EEC as amended by 2007/47/EC
Medical Device Directive
Classification: Class I with measuring function

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

Authorized EU Representative:



Obelis s.a.

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Manufactured by:

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