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:· HEINE Direct Ophthalmoscopes



CE

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MED 112681 **2020-06-02**

V-200.00.152

ENGLISH

HEINE Direct Ophthalmoscopes

These instructions apply to the following products of the HEINE Direct Ophthalmoscope series: BETA200 LED, BETA200, BETA200 M2, BETA200 S LED, BETA200 S, K180, mini3000 LED, mini3000.

Please read and follow these instructions for use and keep them for future reference.

Intended Use

HEINE Direct ophthalmoscopes are intended for transient examination of the media (cornea, aqueous humour, lens, vitreous humour) and retina of the eye. The instruments feature an optical examination system and an illumination unit powered by a battery or rechargeable battery. The device may only be used by qualified medical personnel and in professional healthcare facilities.

For U.S. only:
Federal law restricts this device to sale by or on the order of a Physician or Practitioner.

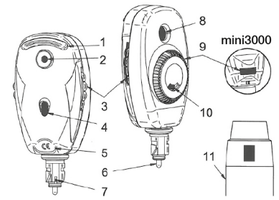
Warnings and Safety Information

CAUTION! This symbol indicates potential hazardous situations. Ignoring the corresponding instructions may lead to dangerous situations of mild to moderate extent.
(Background color yellow; foreground color black).

NOTE! This symbol indicates valuable advice in terms of set up, operation or maintenance, as applicable. Notes are important, but not related to hazardous situations.

Product overview

1 Spectacle rest
2 Viewing aperture
3 Lens wheel
4 Diopter readout (plus = green or black, minus = red)
5 Thumbrest
6 HEINE XHL / LED bulb
7 Connector
8 Viewing window
9 Aperture wheel
10 Filter selector (interference red-free filter)
11 HEINE Handle



Lens wheel (3) correction values

BETA200 LED, BETA200, BETA200 M2, K180
+ in 1D steps: 1-10 | 15 | 20 | 40 | D
- in 1D steps: 1-10 | 15 | 20 | 25 | 35 D

BETA200 S LED, BETA200 S
+ in 1D steps: +1 through +38 D
- in 1D steps: -1 through -36 D

mini3000 LED, mini3000
+ | 1 | 2 | 3 | 4 | 6 | 8 | 10 | 15 | 20 D
- | 1 | 2 | 3 | 4 | 6 | 8 | 10 | 15 | 20 D

Setting up

To set up the HEINE devices, screw the instrument head into the HEINE battery handle or plug it on the HEINE rechargeable handle.

Verify that the lamp voltage complies with the supply voltage of the handle. The coloured marking on the bottom of the lamp shows you the lamp voltage:

White ring = HEINE XHL® 2.5 V bulb
only for use with the HEINE BETA® Battery handle

Red ring = HEINE XHL® 3.5 V bulb
Black ring = HEINE LED illumination
only for use with the HEINE BETA® rechargeable handle or HEINE® wall unit

This does not apply to the HEINE ophthalmoscopes of the mini3000 series. Their lamp voltage is always 2.5 V independently of the handle.

Operation

During the examination, place your index finger on the lens wheel (3) and start the examination using the smallest aperture setting. Place your thumb in the thumbrest (5) or on the attachment clip (mini3000 instruments). In the diopter readout (4), you can read the set diopter value of the lens (negative values are shown in red). You can also operate the aperture wheel (9) with your index finger. With the filter selector (10) you can switch on the red-free filter.

Depending on the instrument, you can choose between different apertures:



From left to right: MicroSpot, medium aperture, large aperture, blue filter, fixation star with polar coordinates, fixation star, slit aperture, hemispot, red-free filter.

HEINE Ophthalmoscopes are intended for a transient examination of less than < 2 minutes with a 15 minutes break until the next application.

Please hold the device as close to the eye as possible.

The setup and operation of the HEINE handles are described in a separate instruction document.

If you realize a significant drop of brightness or a flashing of the lightning immediately after switching on the device, please change the batteries.

Hygienic reprocessing

The instruction is available:
- at www.heine.com
- in a paper version which you can request from the address listed

Maintenance

Changing the light source

Verify that the lamp voltage complies with the supply voltage of the handle.
Allow the device to cool down before changing the bulb.

BETA200 LED / BETA200 S LED / mini3000 LED
With these ophthalmoscopes, the LED cannot be changed.

BETA200 / BETA200 S / BETA200 M2 / K180 / mini3000
Detach the ophthalmoscope from the handle and remove the bulb (6). Insert the new bulb until it locks into place. The lug must fit inside the groove of the guide tube.

Service

The device has no components serviceable by the end-user.

General Notes

The warranty for the entire product is invalidated if non-genuine HEINE products or non-original parts are used and if repairs or modifications are made to the device by persons not authorized by HEINE. For more information, please visit www.heine.com.

The expected life cycle, when the device is normal used and the warning and safety information as well as the maintenance instructions are observed, is up to 7 years. Beyond this period, the product may continue to be used if it is in a safe and good condition.

For the examination please use as little light as possible. To do so, you can use the fixation star as it is combined with a neutral grey filter which reduces the brightness to 30 %. Please ensure to switch off the light after each examination.

General Warnings

Check the correct operation of the device before use! Do not use the device if there are visible signs of damage.

Do not use the device in fire- or explosive risk area (e.g. oxygen saturated or anaesthetic environments).

This product is not allowed to enter or be used in areas with strong magnetic fields e.g. MRI scanners.

Do not modify the device.

Use only original HEINE parts, spare parts, accessories and power sources.

Repairs shall only be carried out by qualified persons.

Light exposure hazard

Because prolonged intense light exposure can damage the retina, the use of the device for ocular examination should not be unnecessarily prolonged, and the brightness setting should not exceed what is needed to provide clear visualization of the target structures.

The retinal exposure dose for a photochemical hazard is a product of the radiance and the exposure time. If the value of radiance were reduced in half, twice the time would be needed to reach the maximum exposure limit.

While no acute optical radiation hazards have been identified for direct or indirect ophthalmoscopes, it is recommended that the intensity of light directed into the patient's eye be limited to the minimum level which is necessary for diagnosis. Infants, aphakes and persons with diseased eyes will be at greater risk. The risk may also be increased if the person being examined has had any exposure with the same instrument or any other ophthalmic instrument using a visible light source during the previous 24 hours. This will apply particularly if the eye has been exposed to retinal photography.

CAUTION – The light emitted from this instrument is potentially hazardous. The longer the duration of exposure, the greater is the risk of ocular damage. Exposure to light from this instrument when operated at maximum intensity will exceed the recommended maximum exposure (RME) of 2.2 J/cm², unless additional action is taken by the user to minimize exposure, after ___* (see table below, column "Recommended maximum").
The risk of retinal injury at an exposure of 2.2 J/cm² is not high, but because some patients may be more susceptible than others, caution is advised if this radiant exposure value is exceeded. However, because of a significant risk of injury at exposures exceeding 10 J/cm², the user should avoid exposures longer than ___* (see table exposure guidelines, column "Maximum").

Disposal

The product must be recycled as separated electrical and electronic devices. Please observe the relevant state-specific disposal regulations.

The appendix contains following tables
- Electromagnetic disturbances – Requirements and tests
- Technical specification
- Exposure guidelines
- Explanation of the used symbols

Electromagnetic disturbances – Requirements and tests	
The device is intended for use in the electromagnetic environment specified below. The customer or the user of the device should assure that it is used in such environments.	
Statement for the operational environments	Inside hospitals except for: near active HF surgical equipment and the RF shielded room of an ME system for magnetic resonance imaging, where the intensity of EM disturbances are high.
Performance features of the ME system that have been determined to be essential to the performance	None
Warning	Use of this equipment adjacent to or stacked with other equipment should be avoided because it could result in improper operation. If such use is necessary, this equipment and the other equipment should be observed to verify that they are operating normally.
	Use of accessories, transducers and cables other than those specified or provided by the manufacturer of this equipment could result in increased electromagnetic emissions or decreased electromagnetic immunity of this equipment and result in improper operation.
	Portable RF communications equipment (including peripherals such as antenna cables and external antennas) should be used no closer than 30 cm (12 inches) to any part of the direct ophthalmoscope, including cables specified by the manufacturer. Otherwise, degradation of the performance of this equipment could result.
A list of all cables, transducers and other accessories that are relevant for the EMC compliance	As applicable, see attachment
Test	**Compliance**
RF emissions CISPR11	Group 1 Class B
Harmonic Emissions	Passed*
Voltage Fluctuations/Flicker	Passed*

*n/a: „Not applicable“ in the internally powered mode

Technical specification

Environmental conditions for operation	+10 °C to +35 °C 30 % to 75 % rel. humidity 700 hPa to 1060 hPa
Environmental conditions for storage	+5 °C to +45 °C 45 % to 80 % rel. humidity 500 hPa to 1060 hPa
Environmental conditions for transport	-20 °C to +50 °C 45 % to 80 % rel. humidity 500 hPa to 1060 hPa
Nominal voltage	3.0 V – 3.7 V
Nominal current	XHL: max. 760 mA LED: max. 350 mA
Protection class	Internally powered
Device classification according to ISO 10942	Group: B
Device classification according to ISO 15004-2	Group: 2

Exposure guidelines*

Instrument type	EN ISO 15004-2:2007	ANSI Z80.36-2016	
		Recommended maximum	Maximum (high risk)
BETA200 LED BETA200 S LED	8 m 00 s	2 m 25 s	11 m 00 s
BETA200 BETA200 M2 BETA200 S	10 m 00 s	2 m 25 s	11 m 00 s
K180	2 m 00 s	4 m 44 s	21 m 29 s
mini3000 LED	47 m 11 s	7 m 36 s	34 m 33 s
mini3000	5 m 00 s	7 m 08 s	33 m 09 s

*) distance 10 mm from instrument to patient

XHL Xenon Halogen Bulbs

Type	2.5 V	3.5 V
BETA200 S/200/200 M2	X-001.88.069	AV: X-002.88.070 TL: X-002.88.072
K180	X-001.88.084	AV: X-002.88.086 TL: X-002.88.085
mini3000	X-001.88.106	

Erläuterung der verwendeten Symbole
Explanation of utilized symbols
Explication des symboles utilisés
Explicación de los símbolos utilizados
Spiegazione dei simboli utilizzati

CE	CE-Kennzeichnung kennzeichnet die Übereinstimmung mit der Europäischen Medizinprodukterichtlinie 93/42 EWG The CE mark indicates that the product complies with the European medical device directive 93/42/EEC. Le marquage CE indique la conformité à la directive européenne 93/42/CEE relative aux dispositifs médicaux. El marcado CE indique la conformidad con la directiva europea 93/42 /CEE relativa a los productos sanitarios. Il marchio CE indica la conformità con la direttiva europea sui dispositivi medici 93/42 CEE
REF	Katalog- oder Bestellnummer Catalogue- or order number Numéro de catalogue ou de commande Número de catálogo o de pedido Codice catalogo e di dell'ordine numero
	Hersteller Manufacturer Fabricant Fabricante Produttore
	Herstelldatum Date of manufacture Date de fabrication Fecha de fabricación Data di produzione
	Getrennte Sammlung von Elektro- und Elektronikgeräten (Europäische WEEE Richtlinie) Product bearing this symbol may not be disposed of together with general household waste, but instead requires separate disposal according to local provisions. (European Waste Electrical and Electronic Equipment Directive, WEEE) Tri sélectif des appareils électriques et électroniques (Directive européenne DEEE) Desechado separado de aparatos eléctricos y electrónicos (Directiva Europea RAEE) Raccolta differenziata di apparecchi elettrici ed elettronici (direttiva europea RAEE)
°C	Zulässiger Temperaturbereich in °C für Lagerung und Transport Temperature limits in °C for storage and transport Plage de température admise en °C pour le stockage et le transport Rango de temperatura permitida en °C para almacenar y transportar el producto Temperatura ammessa in °C per conservazione e trasporto
°F	Zulässiger Temperaturbereich in °F für Lagerung und Transport Temperature limits in °F for storage and transport Plage de température admise en °F pour le stockage et le transport Rango de temperatura permitida en °F para almacenar y transportar el producto Temperatura ammessa in °F per conservazione e trasporto

	Zulässige Luftfeuchtigkeit für Lagerung und Transport Humidity limitation for storage and transport Humidité admise pour le stockage et le transport Humedad del aire permitida para almacenar y transportar el producto Umidità atmosferica ammessa durante il trasporto e la conservazione
	Zulässiger Luftdruck für Lagerung und Transport Pressure limitation for storage and transport Pression atmosphérique admise pendant le transport et le stockage Presión de aire permitida para almacenar y transportar el producto Pressione atmosferica ammessa durante il trasporto e la conservazione
	Vorsicht Bruchgefahr! Fragile, handle with care! Fragile ! Manipuler avec soin Atención. Frágil. Attenzione: pericolo di rottura!
	Trocken lagern! Keep dry! Conserver au sec ! Conservar en un lugar seco! Evitare ambienti umidi!
	Gebrauchsanweisung verbindlich befolgen. (Hintergrundfarbe: blau, Vordergrundfarbe: weiß) Follow instructions for use! (Background color: blue, foreground color: white.) Suivre le mode d'emploi. (Couleur de fond : bleu ; couleur du premier plan : blanc) Seguir obligatoriamente las instrucciones de uso. (Color de fondo: azul, color de primer plano: blanco) Attenersi obbligatoriamente alle istruzioni per l'uso. (Colore dello sfondo: blu, colore in primo piano: bianco)
UDI	Unique Device Identification