

HEINE BETA® 200 / BETA® 200 S LED Ophthalmoscopes

BETA 200: -35 / +40
27 diopter steps
BETA 200 S: -36 / +38
74 single diopter steps

Patented, unique, stepless dimming from 3 % to 100 % with practical one-finger operation.



Reduction of reflections thanks to the “Aspherical Optical System” (AOS) *exclusively from HEINE.

Optical components are mounted on a cast aluminum frame.

LED HQ
LED NOW IN HEINE QUALITY.

DATA	
Description	HEINE BETA 200 / 200S LED Ophthalmoscope
Catalogue Number	C-008.30.100, C-008.30.120
Version / Date	V03 / 16.10.2018

GENERAL	
Product variants	BETA 200 LED Ophthalmoscope BETA 200S LED Ophthalmoscope
Material	synthetics, metal, glas
REACH/RoHS	compliant
Phthalate	product is phthalate free
Latex	product is latex free
Biocompatibility	compliant
Surface	synthetics, metal, glas
Environmental conditions operation	temperature: +10 °C to +35 °C, relative humidity: 30 % to 75 %, air pressure: 700 hPa to 1060 hPa
Environmental conditions storage	temperature: +5 °C to +45 °C, relative humidity: 45 % to 80 %, air pressure: 500 hPa to 1060 hPa
Environmental conditions transport	temperature: -20 °C to +50 °C, relative humidity: 45 % to 80 %, air pressure: 500 hPa to 1060 hPa
Guarantee	5 years
Instructions for use	Deutsch, English, Français, Español, Italiano, Svenska, Nederlands, Português, Dansk, Suomi *
Operating elements	lens wheel, aperture wheel, filter switch
Display	indirect illuminated index of refraction
Power Supply	HEINE rechargeable handles (3.5 V), HEINE EN200 wall transformer, HEINE EN200-1 wall transformer
Accessories	n/a
Patents	n/a

MECHANICAL	
Weight	73 g / 114 g (incl. packaging) 75 g / 116 g (incl. packaging)
Dimensions product	98 x 48 x 28 mm (height x width x depth)
Dimensions packaging	108 x 42 x 68 mm (length x height x depth)
Connections	AV for rechargeable handle
Aufdrucke	examiner-sided: BETA 200 LED, HEINE Logo, CE examiner-sided: BETA 200S LED, HEINE Logo, CE patient-sided: Symbole, HEINE made in Germany AV connector: data matrix code, SN, www.heine.com

ELECTRICAL	
Input voltage	3.0-3.7 V DC
Current consumption	max. 350 mA
Operation time	ca. 7 h using fully loaded Li-ion L rechargeable battery (X-007.99.383)
Safety class	internal power supply
Fuse	n/a

OPTICAL	
Type	LED illumination (HQ) 3.5V fixed in the instrument
Luminous flux**	typ. 0.4 lm
Illuminance** (in 200 mm distance)	typ. 600 lx +/- 150 lx
Color temperature	3500 K +/- 500 K
Color rendering index	typ. CRI ≥ 90, high R9
Lifetime	typ. 42,000 h
Classification according to ISO 10942	group B
Classification according to ISO 15004-2	group 2
Apertures	6 luminous field apertures with separate red-free filter: Slit, fixation star with polar coordinates, cobalt blue filter, large spot, small spot, hemispot 7 luminous field apertures with separate red-free filter: Cobalt blue filter, fixation star with polar coordinates, large spot, small spot, pinhole, slit, hemispot
Lens / Diopter	27 lens / 27 diopter steps (-35 D to +40 D) 28 lens / 74 single-diopter steps (-36 D to +38 D)

* further languages on request
** at 3.7 V supply voltage



CLEANING, DISINFECTION

Procedure	Wipe cleaning and wipe disinfection with agents recommended in the instructions for use. Please consider the detailed information in the accompanying documents!
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CODES

Customs Code	90185090	90185090
EAN/GTIN	4053755182534	4053755182541
HIBC	+E229C008301000017	+E229C0083012000019

Regulatory

Product classification (EU)	Class I
Product classification (USA)	Class 2
Product classification (Canada)	Class I
UMDNS code	12-817
GMDNS code	12817
Regulation number (FDA)	886.1570
Product code (FDA)	HLJ

Fulfills the requirements of Directives & Standards

ISO 13485	Medical devices - Quality management systems - Requirements for regulatory purposes
Directive 93/42/EEC	Concerning medical devices
IEC 60601-1	Medical electrical equipment: General requirements for basic safety and essential performance
IEC 60601-1-2	Medical electrical equipment - Part 1-2: General requirements for basic safety and essential performance -Collateral standard: Electromagnetic disturbances - Requirements and tests
ISO 14971	Medical devices - Application of risk management to medical devices
IEC 60601-1-6	Medical electrical equipment - Part 1-6: General requirements for basic safety and essential performance - Collateral standard: Usability
IEC 62366-1	Medical devices - Part 1: Application of usability engineering to medical devices
DIN EN 15004-1	Ophthalmic instruments – Fundamental requirements and test methods – Part 1: General requirements applicable to all ophthalmic instruments
DIN EN 15004-2	Ophthalmic instruments – Fundamental requirements and test methods – Part 2: Light hazard protection
IEC 60601-1-9	Medical electrical equipment - Part 1-9: General requirements for basic safety and essential performance - Collateral Standard: Requirements for environmentally conscious design
ISO 17664	Processing of health care products - Information to be provided by the medical device manufacturer for the processing of medical devices
ISO 2248	Packaging; complete, filled transport packages; vertical impact test by dropping
Directive (2011/65/EU) ROHS	On the restriction of the use of certain hazardous substances in electrical and electronic equipment
Directive (2012/19/EU) WEEE	On Waste Electrical and Electronic Equipment
Regulation (1907/2006) REACH	Registration, Evaluation, Authorization and Restriction of Chemicals

