



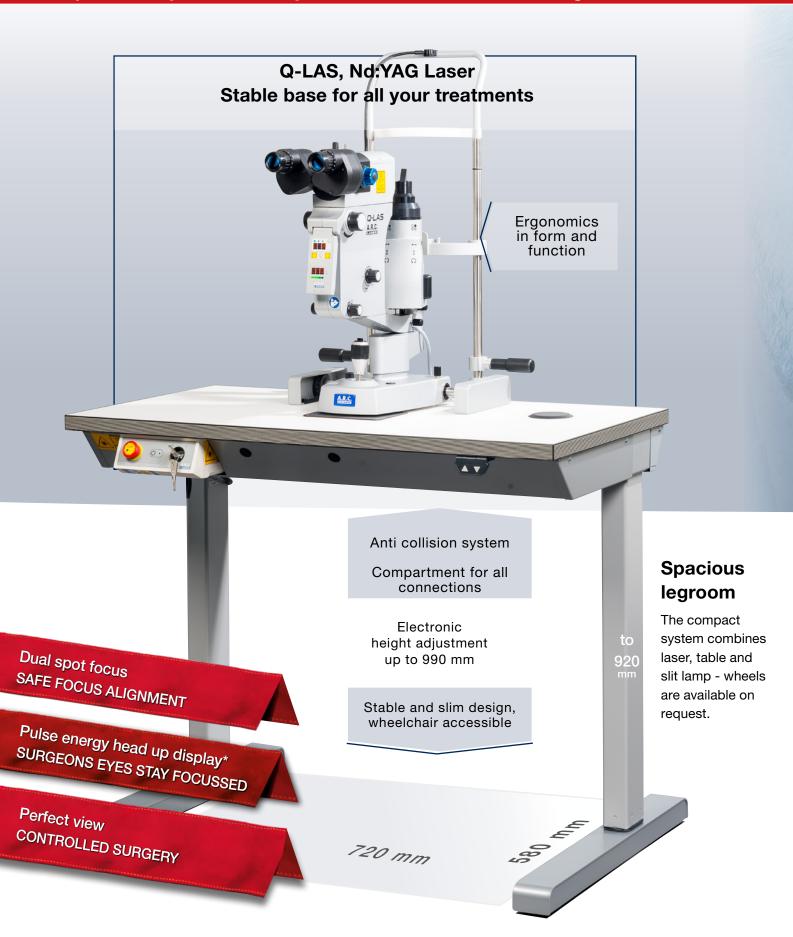
Q-LAS

THE STANDARD LASER
FOR IRIDOTOMY AND
CAPSULOTOMY
IN YOUR DAILY
ROUTINE



www.arclaser.de info@arclaser.de

Capsulotomy - Iridotomy Sleek columns - more legroom



^{*} This accessory is optional, and available on request only.





Premium eye safety

As a standard, our Q-LAS is equipped with a sharp selective safety filter for:

- true color
- detailed resolution

Your choice:

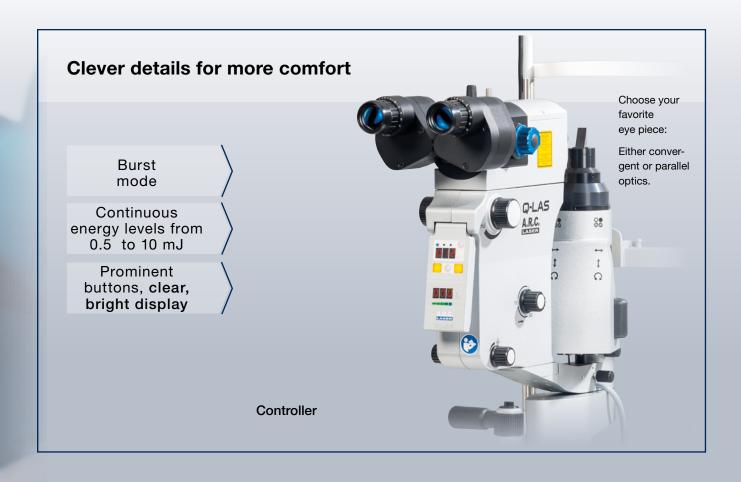
- brand contact glasses
- the optics: parallel-/convergent

Joystick

Height adjustment, slit lamp mobility and laser trigger in one

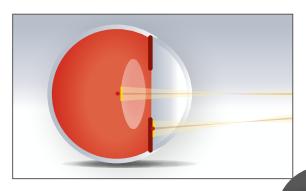
Q-LAS ADVANTAGES

quick & easy to focus



Slit lamp PCL5 Z

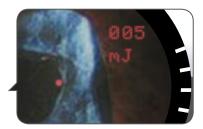
Specially coated optics with parallel or convergent tube provide a detailed view into the anterior segment



The **N**eutral **C**olor **D**esign protection filter is permanently installed.

Head-up Display (optional)

Check all the necessary values at a glance - without taking the eyes from the eyepiece.





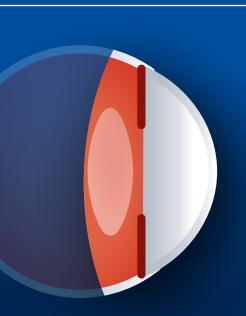
Q-LAS + PCL5

The perfect optic for capsulotomy and iridotomy



PCL5 Z

Designed for the anterior segment





Q-LASThe Nd:YAG laser for you.





Are you looking for the special and reliable solution for secondary cataract surgery? The Nd:YAG laser Q-LAS with PCL5 slit lamp leaves little room for alternatives.



DualSpot-Focus for perfect focussing



Ergonomic handling with prominent buttons



Vertical offset of the slit illumination for a larger visual field

Q-LAS combines economics, ergonomics and longevity in an innovative DESIGN.

MORE OPTIONS: VARIO.

2 high-class lasers on one table.

Nd:YAG + **KTP** laser SLT + KTP laser





SPECIFICATIONS

ND:YAG LASER Q-LAS

ND. IAG LASEN G-LAS	
Laser Wavelength	Q-switched, Nd:YAG, 1064 nm,
Output Energy (Laser)	0.5 mJ to 10 mJ - Single Pulse
Therapy beam pulse settings	0.1 mJ steps from 0.5 mJ (<4 ns) Burst mode 1, 2 or 3 Pulses Cone angle 16°, Spot size <10 µm Defocussing 150/300 µm, posterior
Beam Delivery	Coupling in slit lamp
Display / Control	LED Interface
Cooling	Internal, air
Aiming Beam	635 nm red < 1mW, adjustable
Power Requirement	100-240 V AC, 47/63 Hz, 90 VA
Weight / Dimensions with table and slit lamp	50 kg HWD <99 cm / 100 cm / 58 cm
Laser classification EN 60825-1	Therapy beam: 3B Aiming beam: 2

Alterations of the described features or pictured features are possible. Please keep updated on the current status before ordering.

A.R.C. Laser GmbH

Bessemerstr. 14 90411 Nuremberg

Germany

Subject to change without notice. © A.R.C. Laser 2020.

Risks and warnings: Q-LAS is intended solely for use by trained physicians. YAG laser: contraindicated for eyes with corneal pathologies and chronically elevated IOP. Risks include IOP rise, macular edema and retinal detachment. Refer to the operator manual for a complete list of intended use, contraindications and risks.



VISIBLE AND INVISIBLE LASER RADIATION

Avoid direct irradiation of eye or skin or scattered radiation. laser class: see technical specifications

Your local distributor:



+49 911 217 79 99

4 +49 911 217 79-0

info@arclaser.com www.arclaser.com







EC Certificate

Full Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4) (Devices in Class IIa, IIb or III)

No. G1 18 02 02848 003

Manufacturer:

A.R.C. Laser GmbH

Bessemerstr. 14 90411 Nürnberg **GERMANY**



Facility(ies):

A.R.C. Laser GmbH

Bessemerstr. 14, 90411 Nürnberg, GERMANY

Product Category(ies): Therapeutic and surgical lasers and associated sterile laser contact probes and sterile bare fibers

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.:

713127554

Valid from:

2018-07-01

Valid until:

2023-06-30

2018-04-24 Date,

Stefan Preiß

1. Punil



TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

Page 1 of 1







Certificate

No. Q5 002848 0004 Rev. 01

Holder of Certificate: A.R.C. Laser GmbH

> Bessemerstr. 14 90411 Nürnberg **GERMANY**

A.R.C. Laser GmbH Facility(ies):

Bessemerstr. 14, 90411 Nürnberg, GERMANY

Design and development, manufacturing, installation and service of therapeutic and surgical lasers and associated laser contact probes and bare fibers as well as slit lamps

Certification Mark:



Scope of Certificate: Design and development, manufacturing,

> installation and service of therapeutic and surgical lasers and associated laser contact probes and bare fibers as well as slit lamps

EN ISO 13485:2016 **Applied Standard(s):**

Medical devices - Quality management systems -

Requirements for regulatory purposes

(ISO 13485:2016) DIN EN ISO 13485:2016

The Certification Body of TÜV SÜD Product Service GmbH certifies that the company mentioned above has established and is maintaining a quality management system, which meets the requirements of the listed standard(s). All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: www.tuvsud.com/ps-cert?q=cert:Q5 002848 0004 Rev. 01

Report No.: 713202771

Valid from: 2021-07-16 Valid until: 2024-06-30

Christoph Dicks Date. 2021-07-16

Head of Certification/Notified Body

KONFORMITÄTSERKLÄRUNG / DECLARATION OF CONFORMITY



Name und Adresse der Firma Name and address of the firm

ARC Laser GmbH Bessemerstr. 14 90411 Nürnberg Germany

Wir erklären in alleiniger Verantwortung, dass / We declare under our sole responsibility that

das Medizinprodukt

Q-Las

the medical device

Q-switched Nd:YAG laser 1064 nm

Produktkombination / in combination with

Spaltlampe PCL5, Hersteller: A.R.C. Laser GmbH; Klassifizierung I

Slit lamp PCL5, Producer: A.R.C. Laser GmbH; of Class I

der Klasse / of class

II b, Regel 9 / II b, rule 9

Nach Anhang IX der Richtlinie 93/43/EWG / according to annex IX of direct. 93/42/EEC

Laserklasse / laser classification

3B

UMDNS

16-947

allen Anforderungen der Medizinprodukte-Richtlinie 93/42/EWG entspricht, die anwendbar sind / meets all the provisions of the directive 93/42/EEC which apply to it.

Angewandte harmonisierte Normen, nationale Normen oder andere normative Dokumente

Dem Gerät wurden die betreffenden harmonisierte Normen gemäß der Richtlinie 93/42/EWG Anhang I

zu Grunde gelegt.

Applied harmonised standards, national standards or other normative documents The device is based on the relevant harmonized standards of Directive 93/42/EEC Annex I

Konformitätsbewertungsverfahren Conformity assessment procedure MDD 93/42/EEC, Anhang II ohne (4) MDD 93/42/EEC, Annex II excluding (4)

Konformitätsbewertungsstelle (falls beigezogen) Notified Body (if consulted)

TÜV SÜD Product Service GmbH (CE 0123)

Ridlerstraße 65 / D-80339 München / Germany

Zertifikats Nr. / Registration No.

G1 18 02 02848 003

gültig bis / Valid through

30.06.2023

Nurnberg, 19,07,2018

M. Barten, QMB

90411 Nürnberg

A.R.C. Laser GmbH Bessemerstraße 14

Name und Funktion / Name and function