

## SPECIFICATIILE TEHNICE (F4.1)

Numărul licitației: ocds-b3wdp1-MD-1551098298360;

**ID: 21005662**

Denumirea licitației: *Achiziționarea consumabilelor oftalmologice conform necesităților IMSP pentru anul 2019.*

Nr. Lot	Cod CPV	Descrierea bunurilor	Modelul articolului	Tara de origine	Producător	Specificatia tehnica deplina solicitata de catre autoritatea contractanta	Specificatia tehnica deplina propusa de catre ofertant	Standa rde de referin ta
22.	33100000-1	Casete pentru faecoemulsifi care, compatibile cu aparatul Geuder	G-60700	Germania	<b>Geuder AG</b>	<p>Casete pentru faecoemulsificare compatibile la aparatul Geuder Geuder (Seria S4-1320483, tip Megatron S4), steril/CertIFICATE de calitate:</p> <p>CertIFICATE de calitate:</p> <ul style="list-style-type: none"> <li>- Certificat CE, valabil sau declaratiie de conformitate, valabilă, în funcție de evaluarea conformității cu anexele corespunzătoare, confirmat prin semnătura Participantului.</li> <li>- Certificat ISO 13485 valabil pentru produsele oferite, confirmat prin semnătura Participantului</li> <li>- Termenul de garanție/valabilitate a bunurilor la momentul livrării va fi nu mai mic de 12 luni din termenul total de valabilitate a acestora.</li> </ul> <p>Documente confirmative:</p> <ul style="list-style-type: none"> <li>- Catalogul producătorului/prospecte/documente tehnice de confirmare a specificațiilor tehnice pentru produsul oferit, confirmat prin semnătura Participantului.</li> <li>- Instrucțiunea de utilizare a produsului, inclusiv copie sau original în limba engleză sau rusă și traducerea în limba de stat pentru produsul oferit la livrare, confirmată prin semnătura Participantului.</li> <li>- În ofertă se va indica codul produsului oferit pentru a putea fi identificat conform catalogului prezentat.</li> </ul>	<p>Casete pentru faecoemulsificare compatibile la aparatul Geuder Geuder (Seria S4-1320483, tip Megatron S4), steril/CertIFICATE de calitate:</p> <p>CertIFICATE de calitate:</p> <ul style="list-style-type: none"> <li>- Certificat CE, valabil sau declaratiie de conformitate, valabilă, în funcție de evaluarea conformității cu anexele corespunzătoare, confirmat prin semnătura Participantului. - anexat</li> <li>- Certificat ISO 13485 valabil pentru produsele oferite, confirmat prin semnătura Participantului - anexat</li> <li>- Termenul de garanție/valabilitate a bunurilor la momentul livrării va fi nu mai mic de 12 luni din termenul total de valabilitate a acestora.</li> </ul> <p>Documente confirmative:</p> <ul style="list-style-type: none"> <li>- Catalogul producătorului/prospecte/documente tehnice de confirmare a specificațiilor tehnice pentru produsul oferit, confirmat prin semnătura Participantului. - anexat</li> <li>- Instrucțiunea de utilizare a produsului, inclusiv copie sau original în limba engleză sau rusă și traducerea în limba de stat pentru produsul oferit la livrare, confirmată prin semnătura Participantului.</li> <li>- În ofertă se va indica codul produsului oferit pentru a putea fi identificat conform catalogului prezentat. - indicat</li> </ul>	CE, ISO



59.	33100000-1	Set de vitrectomie anterioară, compatibile cu aparatul Gender (Seria S4-1320483, tip Megatron S4)	G-46001 G-46009	Germania  Gender AG	<p>Set de vitrectomie anterioară pentru aparatul Gender (Seria S4-1320483, tip Megatron S4), steril</p> <p>Certificate de calitate:</p> <ul style="list-style-type: none"> <li>- Certificat CE, valabil sau declarație de conformitate, valabilă, în funcție de evaluarea conformității cu anexele corespunzătoare, confirmat prin semnătura Participantului.</li> <li>- Certificat ISO 13485 valabil pentru produsele oferite, confirmat prin semnătura Participantului</li> <li>- Termenul de garanție/valabilitate a bunurilor la momentul livrării va fi nu mai mic de 12 luni din termenul total de valabilitate a acestora.</li> </ul> <p>Documente confirmative:</p> <ul style="list-style-type: none"> <li>- Catalogul producătorului/prospecte/documente tehnice de confirmare a specificațiilor tehnice pentru produsul oferit, confirmat prin semnătura Participantului.</li> <li>- Instrucțiunea de utilizare a produsului, inclusiv copie sau original în limba engleză sau rusă și traducerea în limba de stat pentru produsul oferit la livrare, confirmată prin semnătura Participantului.</li> <li>- În ofertă se va indica codul produsului oferit pentru a putea fi identificat conform catalogului prezentat.</li> </ul>	<p>Set de vitrectomie anterioară pentru aparatul Gender (Seria S4-1320483, tip Megatron S4), steril, <b>comps din:</b></p> <ul style="list-style-type: none"> <li>- <b>Cutter unica folosinta G-46001</b></li> <li>- <b>Tub de infuzie unica folosinta pentru cutter. G-46009.</b></li> </ul> <p>Certificate de calitate:</p> <ul style="list-style-type: none"> <li>- Certificat CE, valabil sau declarație de conformitate, valabilă, în funcție de evaluarea conformității cu anexele corespunzătoare, confirmat prin semnătura Participantului. - anexat</li> <li>- Certificat ISO 13485 valabil pentru produsele oferite, confirmat prin semnătura Participantului - anexat</li> <li>- Termenul de garanție/valabilitate a bunurilor la momentul livrării va fi nu mai mic de 12 luni din termenul total de valabilitate a acestora.</li> </ul> <p>Documente confirmative:</p> <ul style="list-style-type: none"> <li>- Catalogul producătorului/prospecte/documente tehnice de confirmare a specificațiilor tehnice pentru produsul oferit, confirmat prin semnătura Participantului. - anexat</li> <li>- Instrucțiunea de utilizare a produsului, inclusiv copie sau original în limba engleză sau rusă și traducerea în limba de stat pentru produsul oferit la livrare, confirmată prin semnătura Participantului.</li> <li>- În ofertă se va indica codul produsului oferit pentru a putea fi identificat conform catalogului prezentat.</li> </ul>	CE, ISO
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85.	33100000-1	Vrf pentru faoemulsifi care, compatibil cu aparatul Gender	G-24005	Germania	<b>Gender AG</b>	<p>Vrf steril pentru faoemulsificare compatibile cu aparatul Gender (Seria S4-1320483, tip Megatron S4),</p> <p>Certificare de calitate:</p> <ul style="list-style-type: none"> <li>- Certificat CE, valabil sau declaratiie de conformitate, valabilă, în funcție de evaluarea conformității cu anexele corespunzătoare, confirmat prin semnătura Participantului.</li> <li>- Certificat ISO 13485 valabil pentru produsele oferite, confirmat prin semnătura Participantului</li> <li>- Termenul de garanție/valabilitate a bunurilor la momentul livrării va fi nu mai mic de 12 luni din termenul total de valabilitate a acestora.</li> </ul> <p>Documente confirmative:</p> <ul style="list-style-type: none"> <li>- Catalogul producătorului/prospecte/documente tehnice de confirmare a specificațiilor tehnice pentru produsul oferit, confirmat prin semnătura Participantului.</li> <li>- Instrucțiunea de utilizare a produsului, inclusiv copie sau original în limba engleză sau rusă și traducerea în limba de stat pentru produsul oferit la livrare, confirmată prin semnătura Participantului.</li> <li>- În ofertă se va indica codul produsului oferit pentru a putea fi identificat conform catalogului prezentat.</li> </ul>	<p>Vrf steril pentru faoemulsificare compatibile cu aparatul Gender (Seria S4-1320483, tip Megatron S4), <b>reutilizabil, din titan.</b></p> <p>Certificare de calitate:</p> <ul style="list-style-type: none"> <li>- Certificat CE, valabil sau declaratiie de conformitate, valabilă, în funcție de evaluarea conformității cu anexele corespunzătoare, confirmat prin semnătura Participantului. - anexat</li> <li>- Certificat ISO 13485 valabil pentru produsele oferite, confirmat prin semnătura Participantului - anexat</li> <li>- Termenul de garanție/valabilitate a bunurilor la momentul livrării va fi nu mai mic de 12 luni din termenul total de valabilitate a acestora.</li> </ul> <p>Documente confirmative:</p> <ul style="list-style-type: none"> <li>- Catalogul producătorului/prospecte/documente tehnice de confirmare a specificațiilor tehnice pentru produsul oferit, confirmat prin semnătura Participantului. - anexat</li> <li>- Instrucțiunea de utilizare a produsului, inclusiv copie sau original în limba engleză sau rusă și traducerea în limba de stat pentru produsul oferit la livrare, confirmată prin semnătura Participantului.</li> <li>- În ofertă se va indica codul produsului oferit pentru a putea fi identificat conform catalogului prezentat. — indicat</li> </ul>	CE, ISO
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Director Tehnomedica SRL,

Tatiana Roibu



# TEHNOMEDICA

Ciuflea 38/1 2001 Cisinou Moldova tel./fax 601 102, 601 087  
e-mail <tehnomedica\_md@yahoo.com>

**Către Centrul pentru Achiziții Publice Centralizate în Sănătate**

**În atenția Grupului de Lucru al Licitației nr. ocds-b3wdp1-MD-1551098298360;  
ID: 21005662 din 13.04.2019**

## Declaratie

### termen de valabilitate

Prin prezenta, declarăm, că pentru produsele oferite in LP **21005662** confirmăm, termenul de valabilitate a produselor la livrare va fi nu mai mic de 80% din termenul de valabilitate al acestuia garantat de producător si nu mai mic de 12 luni.

Cu respect,  
Director  
13.04.19



Tatiana Roibu

# bsi.



By Royal Charter

## EC Certificate - Production Quality Assurance

Directive 93/42/EEC on Medical Devices, Annex V

**No.**

**CE 575413**

Issued To:

**Geuder AG  
Hertzstraße 4  
69126 Heidelberg  
Germany**

In respect of:

**See certificate scope page.**

on the basis of our examination of the quality assurance system under the requirements of Council Directive 93/42/EEC, Annex V. The quality assurance system meets the requirements of the directive. For the placing on the market of class IIb and class III products an Annex III certificate is required.

For and on behalf of BSI, a Notified Body for the above Directive (Notified Body Number 0086):

Frank Lee, EMEA Compliance & Risk Director

First Issued: **29 June 2016**

Date: **29 June 2016**

Expiry Date: **25 May 2019**

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Page 1 of 2

Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive as demonstrated through the regular surveillance activities of the Notified Body. This approval excludes all products designed and/or manufactured by a third party on behalf of the company named on this certificate, unless specifically agreed with BSI.  
This certificate was issued electronically and is bound by the conditions of the contract.

Information and Contact: BSI, Kitemark Court, Davy Avenue, Knowlhill, Milton Keynes MK5 8PP. Tel: + 44 345 080 9000  
BSI Assurance UK Limited, registered in England under number 7805321 at 389 Chiswick High Road, London W4 4AL, UK.  
A member of BSI Group of Companies.



Certificate No: CE 575413

## Certificate Scope:

**Manufacture of non active, sterile and non sterile ophthalmic surgical products and accessories.**

**Produktion von nicht aktiven, sterilen und nicht sterilen ophthalmo-chirurgischen Produkten und Zubehör.**

**Those aspects of Annex V related to metrology in the manufacture of ophthalmic surgical measuring devices.**

**Die Aspekte des Anhangs V im Zusammenhang mit der Messfunktion bei der Herstellung der ophthalmo-chirurgischen Messinstrumente.**

**Those aspects of Annex V related to sterility in the manufacture of ophthalmic surgical irrigation/aspiration (I/A) instruments, single-use irrigation cannulas, single-use endoprobes, single-use fiber optics, single-use adapters for glass syringes and Bonn Injection Sets.**

**Die Aspekte des Anhangs V im Zusammenhang mit der Sterilität bei der Herstellung der ophthalmo-chirurgischen Spül- und Sauginstrumente, Einmal-Spülkanülen, Einmal-Endosonden, Einmal-Lichtleiter, Einmal-Adapter für Glasspritzen und Bonner Injektionssets.**

First Issued: **29 June 2016**

Date: **29 June 2016**

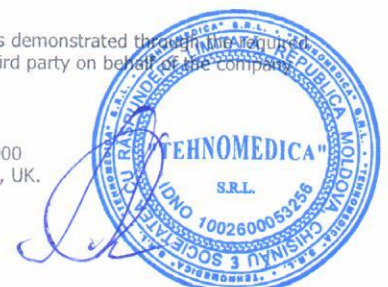
Expiry Date: **25 May 2019**

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Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive as demonstrated through the surveillance activities of the Notified Body. This approval excludes all products designed and/or manufactured by a third party on behalf of the manufacturer named on this certificate, unless specifically agreed with BSI.  
This certificate was issued electronically and is bound by the conditions of the contract.

Information and Contact: BSI, Kitemark Court, Davy Avenue, Knowlhill, Milton Keynes MK5 8PP. Tel: + 44 345 080 9000  
BSI Assurance UK Limited, registered in England under number 7805321 at 389 Chiswick High Road, London W4 4AL, UK.  
A member of BSI Group of Companies.



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# Certificate of Registration

## QUALITY MANAGEMENT SYSTEM - ISO 9001:2015

This is to certify that:

Geuder AG  
Hertzstraße 4  
69126 Heidelberg  
Germany

Holds Certificate No:

**FM 575411**

and operates a Quality Management System which complies with the requirements of ISO 9001:2015 for the following scope:

Design, development, manufacture, warehousing, distribution, installation and maintenance of ophthalmo-surgical instruments, active device systems and associated accessories, implants, sterilisation trays and containers.  
Warehousing and distribution of gases, silicone oils, dyes, perfluorocarbons and semifluorinated alkanes for ophthalmology.

For and on behalf of BSI:

Andrew Launn, EMEA Systems Certification Director

Original Registration Date: 2001-06-01

Latest Revision Date: 2018-09-14

Effective Date: 2017-05-26

Expiry Date: 2020-05-25



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Information and Contact: BSI, Kitemark Court, Davy Avenue, Knowlhill, Milton Keynes MK5 8PP. Tel: + 44 345 080 9000

BSI Assurance UK Limited, registered in England under number 7805321 at 389 Chiswick High Road, London W4 4AL, UK.



# bsi.



By Royal Charter

## Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 13485:2016 & EN ISO 13485:2016

This is to certify that:

Geuder AG  
Hertzstraße 4  
69126 Heidelberg  
Germany

Holds Certificate Number:

**MD 575412**

and operates a Quality Management System which complies with the requirements of ISO 13485:2016 & EN ISO 13485:2016 for the following scope:

Please see scope page.

For and on behalf of BSI:

Stewart Brain, Head of Compliance & Risk - Medical Devices

Original Registration Date: 2016-06-27

Latest Revision Date: 2018-12-13

Effective Date: 2017-05-26

Expiry Date: 2020-05-25

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Information and Contact: BSI, Kitemark Court, Davy Avenue, Knowlhill, Milton Keynes MK5 8PP. Tel: + 44 345 080 9000  
BSI Assurance UK Limited, registered in England under number 7805321 at 389 Chiswick High Road, London W4 4AL, UK.  
A Member of the BSI Group of Companies.





Certificate No: **MD 575412**

## Registered Scope:

Design, development, manufacture and control of manufacture, warehousing, distribution, installation and maintenance of active and non active, sterile and non sterile ophthalmic surgical devices/systems, instruments and accessories, ophthalmic implants, sterilization trays and containers.

Warehousing and distribution of silicone oils, dyes, perfluorocarbons and semifluorinated alkanes for use as liquid intraocular endotamponades and gas-based intraocular tamponades and vitreous substitutes for the area of ophthalmology.

Auslegung, Entwicklung, Produktion und Lenkung der Produktion, Lagerhaltung, Vertrieb, Installation und Instandhaltung von aktiven und nicht aktiven, sterilen und nicht sterilen ophthalmo-chirurgischen Geräten/Systemen, Instrumenten und Zubehör, ophthalmologischen Implantaten, Sterilisationsbehältern und Sterilisationscontainern.

Lagerhaltung und Vertrieb von Silikonölen, Färbemitteln, Perfluorcarbonverbindungen und semifluorierten Alkanen für die Verwendung als flüssige intraokulare Endotamponaden, gasförmige intraokulare Tamponaden und Glaskörperersatzstoffe im Einsatzbereich der Ophthalmologie.

Original Registration Date: 2016-06-27

Latest Revision Date: 2018-12-13

Effective Date: 2017-05-26

Expiry Date: 2020-05-25

This certificate was issued electronically and remains the property of BSI and is bound by the conditions of contract.

An electronic certificate can be authenticated [online](#).

Printed copies can be validated at [www.bsigroup.com/ClientDirectory](http://www.bsigroup.com/ClientDirectory)

Information and Contact: BSI, Kitemark Court, Davy Avenue, Knowlhill, Milton Keynes MK5 8PP. Tel: + 44 345 080 9000  
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# bsi.



By Royal Charter

## EC Certificate - Full Quality Assurance System

Directive 93/42/EEC on Medical Devices, Annex II excluding Section 4

**No.** CE 575415  
**Issued To:** Geuder AG  
Hertzstraße 4  
69126 Heidelberg  
Germany

In respect of:

**Design, development and manufacture of active and non active, sterile and non sterile ophthalmic surgical devices/systems, instruments, accessories and ophthalmic implants.**

**Auslegung, Entwicklung und Produktion von aktiven und nicht aktiven, sterilen und nicht sterilen ophthalmo-chirurgischen Geräten/Systemen, Instrumenten, Zubehör und ophthalmologischen Implantaten.**

on the basis of our examination of the quality assurance system under the requirements of Council Directive 93/42/EEC, Annex II excluding section 4. The quality assurance system meets the requirements of the directive. For the placing on the market of class III products an Annex II section 4 certificate is required.

For and on behalf of BSI, a Notified Body for the above Directive (Notified Body Number 0086):

Frank Lee, EMEA Compliance & Risk Director

First Issued: **29 June 2016**

Date: **29 June 2016**

Expiry Date: **25 May 2019**

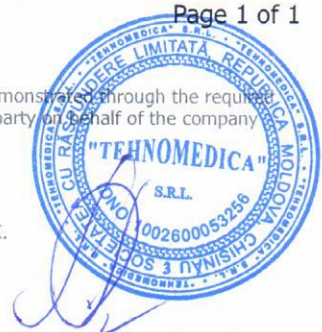
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Page 1 of 1

Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive as demonstrated through the required surveillance activities of the Notified Body. This approval excludes all products designed and/or manufactured by a third party on behalf of the company named on this certificate, unless specifically agreed with BSI.

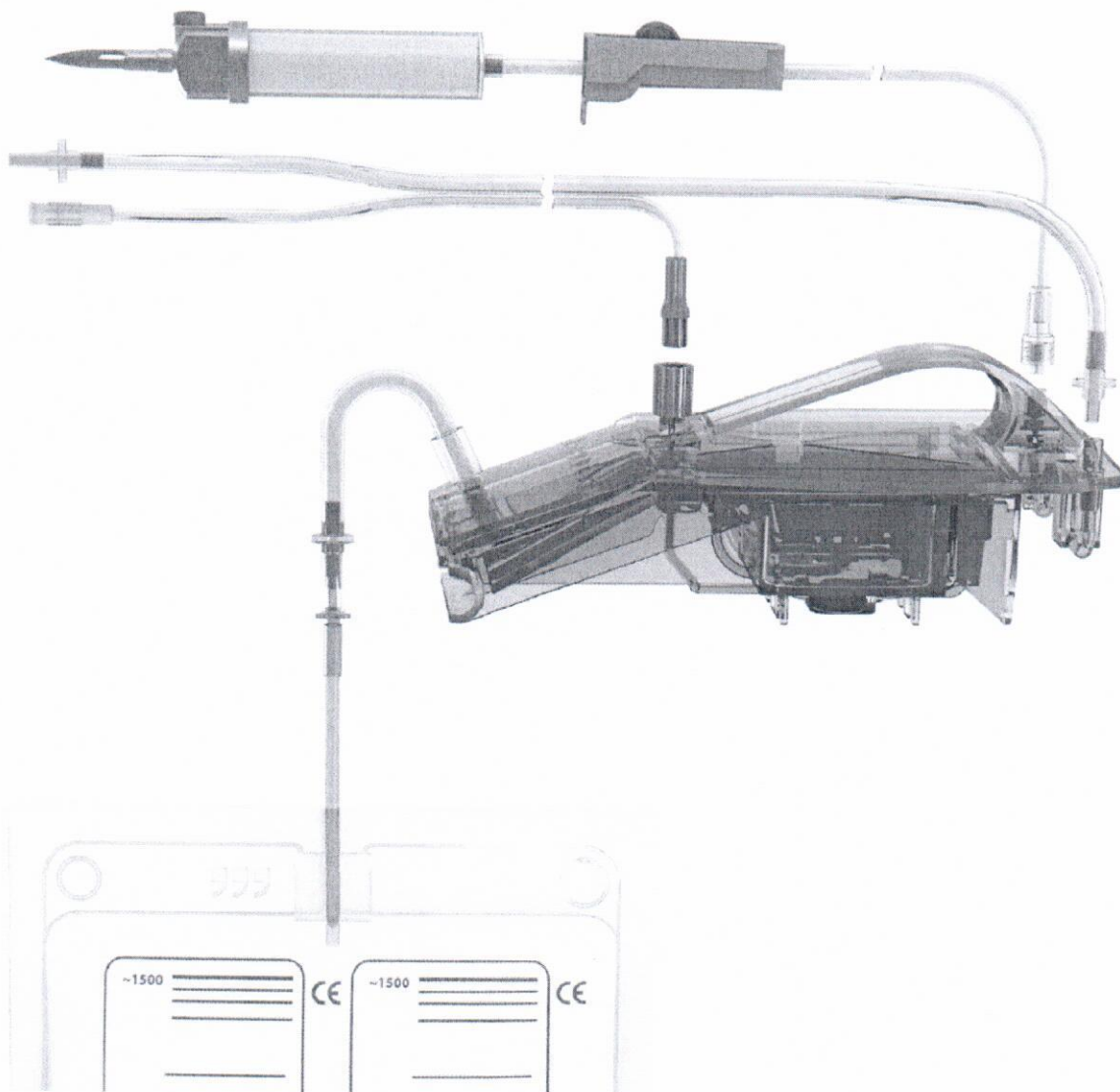
This certificate was issued electronically and is bound by the conditions of the contract.

Information and Contact: BSI, Kitemark Court, Davy Avenue, Knowlhill, Milton Keynes MK5 8PP. Tel: + 44 345 080 9000  
BSI Assurance UK Limited, registered in England under number 7805321 at 389 Chiswick High Road, London W4 4AL, UK.  
A member of BSI Group of Companies.



KASSETTEN - steril  
 CASSETTES - sterile

**Geuder**<sup>®</sup>  
 Precision made in Germany



**KASSETTE**  
 für megaTRON S3 V / VIP / S4<sup>HPS</sup>,  
 inkl. Irrigationsbesteck,  
 GEUDER Sicherheitskonnektor,  
 Auffangbeutel  
 VPE 20 Stück, **steril**

**CASSETTE**  
 for megaTRON S3 V / VIP / S4<sup>HPS</sup>,  
 with irrigation set,  
 GEUDER safety-connector,  
 drain bag  
 20 pcs. per box, **sterile**

**G-30161** **KASSETTE**  
 inkl. VA Schläuche

**CASSETTE**  
 incl. VA tubing

**G-60700** **HPS KASSETTE**  
 inkl. HPS VA Schlauch

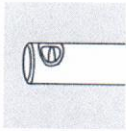
**HPS CASSETTE**  
 incl. HPS VA tubing

*Lot dd.*



UNO COLORLINE  
HIGH-SPEED VITREKTOREN - steril  
UNO COLORLINE  
HIGH SPEED VITREOUS CUTTER - sterile

**Geuder**<sup>®</sup>  
Precision made in Germany



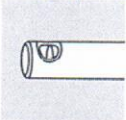
**EINMAL-  
HIGH-SPEED-VITREKTOR 20G**  
Uno Colorline ROT:  
20 Gauge / 0.9 mm  
pneumatisch,  
mit GEUDER Sicherheitskonnektor  
**steril**

**SINGLE-USE HIGH SPEED  
VITREOUS CUTTER 20G**  
Uno Colorline RED:  
20 gauge / 0.9 mm  
pneumatic,  
with GEUDER safety-connector  
**sterile**

**G-42001** VPE 1 Stück  
**G-46001** VPE 6 Stück

1 pc. per box  
6 pcs. per box

*Ref 59.*

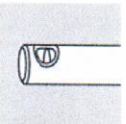


**EINMAL-  
HIGH-SPEED-VITREKTOR 23G**  
Uno Colorline GRÜN:  
23 Gauge / 0.64 mm  
pneumatisch,  
mit GEUDER Sicherheitskonnektor  
**steril**

**SINGLE-USE HIGH SPEED  
VITREOUS CUTTER 23G**  
Uno Colorline GREEN:  
23 gauge / 0.64 mm  
pneumatic,  
with GEUDER safety-connector  
**sterile**

**G-42301** VPE 1 Stück  
**G-46301** VPE 6 Stück

1 pc. per box  
6 pcs. per box



**EINMAL-  
HIGH-SPEED-VITREKTOR 25G**  
Uno Colorline BLAU:  
25 Gauge / 0.5 mm  
pneumatisch,  
mit GEUDER Sicherheitskonnektor  
**steril**

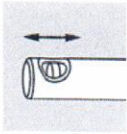
**SINGLE-USE HIGH SPEED  
VITREOUS CUTTER 25G**  
Uno Colorline BLUE:  
25 gauge / 0.5 mm  
pneumatic,  
with GEUDER safety-connector  
**sterile**

**G-42501** VPE 1 Stück  
**G-46501** VPE 6 Stück

1 pc. per box  
6 pcs. per box



**UNO COLORLINE HIGH-SPEED VITREKTOREN - steril**  
**UNO COLORLINE HIGH SPEED VITREOUS CUTTER - sterile**

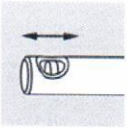


**EINMAL-HIGH-SPEED-VITREKTOR 20G MACH 2**  
**Uno Colorline ROT:**  
 20 Gauge / 0.9 mm  
 Doppelschneide, pneumatisch, mit GEUDER Sicherheitskonnektor steril

**SINGLE-USE HIGH SPEED VITREOUS CUTTER 20G MACH 2**  
**Uno Colorline RED:**  
 20 gauge / 0.9 mm  
 Doubleblade, pneumatic, with GEUDER safety-connector sterile

G-42002 VPE 1 Stück  
 G-46002 VPE 6 Stück

1 pc. per box  
 6 pcs. per box

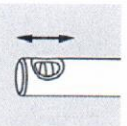


**EINMAL-HIGH-SPEED-VITREKTOR 23G MACH 2**  
**Uno Colorline GRÜN:**  
 23 Gauge / 0.64 mm  
 Doppelschneide, pneumatisch, mit GEUDER Sicherheitskonnektor steril

**SINGLE-USE HIGH SPEED VITREOUS CUTTER 23G MACH 2**  
**Uno Colorline GREEN:**  
 23 gauge / 0.64 mm  
 Doubleblade, pneumatic, with GEUDER safety-connector sterile

G-42302 VPE 1 Stück  
 G-46302 VPE 6 Stück

1 pc. per box  
 6 pcs. per box



**EINMAL-HIGH-SPEED-VITREKTOR 25G MACH 2**  
**Uno Colorline BLAU:**  
 25 Gauge / 0.5 mm  
 Doppelschneide, pneumatisch, mit GEUDER Sicherheitskonnektor steril

**SINGLE-USE HIGH SPEED VITREOUS CUTTER 25G MACH 2**  
**Uno Colorline BLUE:**  
 25 gauge / 0.5 mm  
 Doubleblade, pneumatic, with GEUDER safety-connector sterile

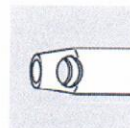
G-42502 VPE 1 Stück  
 G-46502 VPE 6 Stück

1 pc. per box  
 6 pcs. per box

**G-46009**

**EINMAL-INFUSIONSROHR 20G**  
**Uno Colorline ROT:**  
 20 Gauge / 0.9 mm  
 für 20 Gauge Vitrektoren  
 G-42001 und G-46001  
 VPE 6 Stück, steril

**SINGLE-USE INFUSION TUBE 20G**  
**Uno Colorline RED:**  
 20 Gauge / 0.9 mm  
 für 20 gauge vitreous cutters  
 G-42001 and G-46001  
 6 pcs. per box, sterile

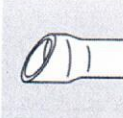



*Lot 59*

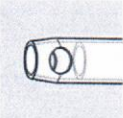

**ULTRASCHALLHANDGRIFFE UND ZUBEHÖR**  
 - wiederverwendbar  
**ULTRASONIC HANDPIECES AND ACCESSORIES**  
 - reusable

	<b>ULTRASCHALLSPITZE</b> Titan, Durchmesser 1.17 mm	<b>ULTRASONIC TIP</b> titanium, diameter 1.17 mm		
<b>G-24040</b>	15°	15°		
<b>G-24050</b>	30°	30°		
<b>G-24060</b>	45°	45°		

*Lot 85*

<b>G-24055</b>	<b>MEGA-ULTRASCHALLSPITZE</b> Titan, 30°, (DE-Patent 43 13 245), Durchmesser an der Spitze 1.26 mm	<b>MEGA ULTRASONIC TIP</b> titanium, 30°, (US-Patent 5,451,229), diameter at the tip 1.26 mm		
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<b>G-24101</b>	<b>INFUSIONSAUFSATZ</b> Silikon, gelb, für Ultraschall-Handgriff, (DE-Patent 40 08 594), Infusionsrohr Ø 1.8 mm	<b>INFUSION SLEEVE</b> silicone, yellow, for ultrasonic handpiece, (US-Patent 40 08 594), infusion tube Ø 1.8 mm		
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<b>G-24104</b>	<b>INFUSIONSAUFSATZ</b> Silikon, gelb, flexibles Infusionsrohr mit Silikonüberzug, (DE-Patent 197 00 809), Infusionsrohr Ø 1.8 mm	<b>INFUSION SLEEVE</b> silicone, yellow, with flexible, silicone coated infusion tube, (US-Patent 5,879,356), infusion tube Ø 1.8 mm		
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<b>G-24111</b>	<b>INFUSIONSAUFSATZ</b> gehärtetes Silikon, gelb für Ultraschall-Handgriff (DE-Patent 40 08 594) Infusionsrohr 1.8 mm	<b>INFUSION SLEEVE</b> hardened silicone, yellow-colored for ultrasonic handpiece (DE-Patent 40 08 594) infusion tube 1.8 mm		
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<b>G-24313</b>	<b>PEP 2.4 MM ULTRASCHALLSPITZE</b> Pure Efficiency Phaco für 2.4 mm Inzisionen Titan, 40°, 3 Stufen Design, grün	<b>PEP 2.4 MM ULTRASONIC TIP</b> Pure Efficiency Phaco for 2.4 mm incisions titanium, 40°, 3 step design, green		
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<b>G-24314</b>	<b>PEP 2.4 MM INFUSIONSAUFSATZ</b> Pure Efficiency Phaco für 2.4 mm Inzisionen, grün transparent, Silikon	<b>PEP 2.4 MM INFUSION SLEEVE</b> Pure Efficiency Phaco for 2.4 mm incisions, green translucent, silicone		
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<b>G-24310</b>	<b>PEP 2.2 MM ULTRASCHALLSPITZEN</b> Pure Efficiency Phaco für 2.2 mm Inzision Titan, 40°, 3 Stufen-Design, violett	<b>PEP 2.2 MM ULTRASONIC TIP</b> Pure Efficiency Phaco for 2.2 mm incision titanium, 40°, 3 step design, purple		
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