

EC Certificate of Conformity

The Notified Body

**MEDCERT Zertifizierungs- und Prüfungsgesellschaft für die Medizin GmbH
Pilatuspool 2 – 20355 Hamburg – Germany**

herewith certifies that the company:

**megro GmbH & Co. KG
Am Schornacker 30
46485 Wesel
Germany**

has introduced and maintains a quality assurance system for the products / product categories listed in the appendix.

The compliance of this quality assurance system with the below mentioned requirements of the **Council Directive 93/42/EEC** was verified by an audit:

Annex II without section 4

This certification is subject to surveillance by MEDCERT.

This certificate is valid until 01 March 2024

Report No.: 2239FS19F
Process No.: QS – 2239
Certificate No.: 2239GB410190924

Hamburg, 24 September 2019



MEDCERT Certification Body
(Lorenz Runge)

The certificate is only valid when provided entirely with all of its pages.
To verify the validity of this certificate, contact info@medcert.de.

MEDCERT Identification Number: 0482

Form F10010005e EN / Rev. 10 / 2019.05.22



Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
www.zlg.de
ZLG-BS-237.10.15

Appendix of EC Certificate of Conformity

Process No.: QS – 2239

Certificate No.: 2239GB410190924

List of products / product categories included in the scope of certificate**Disinfectants**

– End of list –

This appendix is integral part of the above-referenced certificate.
The certificate is only valid when provided entirely with all of its pages.
To verify the validity of this certificate, contact info@medcert.de.

MEDCERT Identification Number: 0482



EC Certificate of Conformity

The Notified Body

**MEDCERT Zertifizierungs- und Prüfungsgesellschaft für die Medizin GmbH
Pilatuspool 2 – 20355 Hamburg – Germany**

herewith certifies that the company:

**megro GmbH & Co. KG
Am Schornacker 30
46485 Wesel
Germany**

has introduced and maintains a quality assurance system
**concerning the conformity of the medical devices with the metrological
requirements**

for the products / product categories listed in the appendix.

The compliance of this quality assurance system with the below mentioned requirements of the
Council Directive 93/42/EEC was verified by an audit:

Annex V

This certification is subject to surveillance by MEDCERT.

This certificate is valid until 01 March 2024

Report No.: 2239FS19F
Process No.: QS – 2239
Certificate No.: 2239GB416190924

Hamburg, 24 September 2019


MEDCERT Certification Body
(Lorenz Runge)

The certificate is only valid when provided entirely with all of its pages.
To verify the validity of this certificate, contact info@medcert.de.

MEDCERT Identification Number: 0482

Form F10010014e EN / Rev. 8 / 2019.05.22



Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
www.zlg.de
ZLG-BS-237.10.15

Appendix of EC Certificate of Conformity

Process No.: QS – 2239

Certificate No.: 2239GB416190924

List of products / product categories included in the scope of certificate

- **Blood pressure monitors (aneroid)**
- **Bladder syringes**

– End of list –

This appendix is integral part of the above-referenced certificate.
The certificate is only valid when provided entirely with all of its pages.
To verify the validity of this certificate, contact info@medcert.de.

MEDCERT Identification Number: 0482

Form F10010014e EN / Rev. 8 / 2019.05.22



Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
www.zlg.de
ZLG-BS-237.10.15

EC Certificate of Conformity

The Notified Body

**MEDCERT Zertifizierungs- und Prüfungsgesellschaft für die Medizin GmbH
Pilatuspool 2 – 20355 Hamburg – Germany**

herewith certifies that the company:

**megro GmbH & Co. KG
Am Schornacker 30
46485 Wesel
Germany**

has introduced and maintains a quality assurance system
for the aspects concerned with securing and maintaining sterile conditions

for the products / product categories listed in the appendix.

The compliance of this quality assurance system with the below mentioned requirements of the
Council Directive 93/42/EEC was verified by an audit:

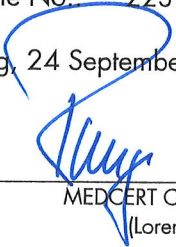
Annex V

This certification is subject to surveillance by MEDCERT.

This certificate is valid until 01 March 2024

Report No.: 2239FS19F
Process No.: QS – 2239
Certificate No.: 2239GB415190924

Hamburg, 24 September 2019



MEDCERT Certification Body
(Lorenz Runge)

The certificate is only valid when provided entirely with all of its pages.
To verify the validity of this certificate, contact info@medcert.de.

MEDCERT Identification Number: 0482

Form F10010014e EN / Rev. 8 / 2019.05.22



Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
www.zlg.de
ZLG-BS-237.10.15

Appendix of EC Certificate of Conformity

Process No.: QS – 2239

Certificate No.: 2239GB415190924

List of products / product categories included in the scope of certificate

- **Urine bags**
- **Skin markers**
- **Rectal tubes**
- **Urological catheters**
- **Sterile medical drapes**
- **Tweezers for single use**
- **Bandaging materials**
- **Bladder syringes**
- **Copolymer examination gloves**

– End of list –

This appendix is integral part of the above-referenced certificate.
The certificate is only valid when provided entirely with all of its pages.
To verify the validity of this certificate, contact info@medcert.de.

MEDCERT Identification Number: 0482

Form F10010014e EN / Rev. 8 / 2019.05.22



Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
www.zlg.de
ZLG-BS-237.10.15

EC Certificate of Conformity

The Notified Body

**MEDCERT Zertifizierungs- und Prüfungsgesellschaft für die Medizin GmbH
Pilatuspool 2 – 20355 Hamburg – Germany**

herewith certifies that the company:

**megro GmbH & Co. KG
Am Schornacker 30
46485 Wesel
Germany**

has introduced and maintains a quality assurance system for the products / product categories listed in the appendix.

The compliance of this quality assurance system with the below mentioned requirements of the **Council Directive 93/42/EEC** was verified by an audit:

Annex V

This certification is subject to surveillance by MEDCERT.

This certificate is valid until 01 March 2024

Report No.: 2239FS19F
Process No.: QS – 2239
Certificate No.: 2239GB414190924

Hamburg, 24 September 2019


MEDCERT Certification Body
(Lorenz Runge)

The certificate is only valid when provided entirely with all of its pages.
To verify the validity of this certificate, contact info@medcert.de.

MEDCERT Identification Number: 0482

Form F10010005e EN / Rev. 10 / 2019.05.22



Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten www.zlg.de
ZLG-BS-237.10.15

Appendix of EC Certificate of Conformity

Process No.: QS – 2239

Certificate No.: 2239GB414190924

List of products / product categories included in the scope of certificate

- **Blood lancets**
- **Medical electric thermometers**
- **Scalpels and scalpel blades**
- **Vacuum bottles**
- **Nasal oxygen cannulas**
- **Stitch cutters**
- **Stitch cutter sets**
- **Acupuncture needles**
- **Oxygen catheters**
- **Suction catheters**
- **Disinfectants**

– End of list –

This appendix is integral part of the above-referenced certificate.
The certificate is only valid when provided entirely with all of its pages.
To verify the validity of this certificate, contact info@medcert.de.

MEDCERT Identification Number: 0482



Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
www.zlg.de
ZLG-BS-237.10.15