

### **EC** Certificate

Directive 98/79/EC Annex IV, excluding Sections 4 and 6
Full Quality Assurance System
In Vitro Diagnostic Medical Devices

Registration No.: HL 60128323 0001

Report No.: 21195407 011

Manufacturer: DRG Instruments GmbH

Frauenbergstr. 18 35039 Marburg Deutschland

**Products:** In vitro diagnostic reagents

(see attachment for products included)

Replaces EC Certificate, Registration No.: HL 60084590 0001

**Expiry Date:** 2023-04-02

The Notified Body hereby declares that the requirements of Annex IV, excluding section 4 and 6 of the directive 98/79/EC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex IV, section 5 of the aforementioned directive. For placing on the market of List A devices covered by this certificate an EC design-examination certificate according to Annex IV, section 4 and and LG4 Annex IV and LG4 Annex I

verification of manufactured products according to section 6 is required.

**Effective Date:** 2018-04-03

Date: 2018-04-03

Dipl./Ing//Sven Hoffmann

Notified Bod

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg

TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 98/79/EC concerning in vitro diagnostic medical devices with the identification number 0197.



Doc. 1/2, Rev. 0

# TÜV Rheinland LGA Products GmbH Tillystraße 2, 90431 Nürnberg

Attachment to Certificate

Registration No.: HL 60128323 0001 Report No.: 21195407 011

Manufacturer: DRG Instruments GmbH

Frauenbergstr. 18 35039 Marburg Deutschland

#### Products included:

Reagents and reagent products for evaluating the risk of trisomy 21:

- Pregnancy associated Plasma Protein-A ELISA, EIA-2397
- Free ß-HCG ELISA, EIA-4718

Reagents and reagent products for determining the tumoral marker PSA:

- Free PSA ELISA, EIA-4189
- Total PSA ELISA, EIA-3719
- Total PSA ELISA (HYBRID-XL), HYE-5370

Reagents and reagent products for diagnosing phenylketonuria:

- Phenylalanine PKU, ENZ-4502 and ENZ-4743

**Notified Body** 

Dipl.-Ing. Sen Hoffmann

Date: 2018-04-03



Doc. 2/2, Rev. 0

## TÜV Rheinland LGA Products GmbH Tillystraße 2, 90431 Nürnberg

Attachment to Certificate

Registration No.:

HL 60128323 0001

Report No.:

21195407 011

Manufacturer:

**DRG Instruments GmbH** 

Frauenbergstr. 18 35039 Marburg Deutschland

### Products included:

Reagents and reagent products for determining infectious diseases:

- Chlamydia trachomatis IgA, EIA-3461
- Chlamydia trachomatis IgG, EIA-3462
- Chlamydia trachomatis IgM, EIA-3463
- Toxoplasma gondii IgA, EIA-3683
- Toxoplasma gondii IgG, EIA-3519
- Toxoplasma gondii IgM, EIA-3520
- Rubella IgG, EIA-3510
- Rubella IgM, EIA-3511
- CMV IgG, EIA-3468
- CMV IgG, EIA-5778
- CMV IgM, EIA-3469

Date: 2018-04-03

10/020 h 04.08 @ TÜY, TUEV and TUV are registered trademarks. Utilisation and application requires prior approval

Notified Body

Dipl.-Ing. Sven Hoffmanr