Product information

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Glutamate ELISA

RUO

REF **DEE2400R**

Σ **96**

Demeditec Diagnostics GmbH
Lise-Meitner-Strasse 2
24145 Kiel – Germany
www.demeditec.com

Table of contents

1. Introduction

1.1 Intended use and principle of the test

Enzyme Immunoassay for the quantitative determination of L-glutamate in urine and various biological samples.
After extraction and derivatisation Glutamate is quantitatively determined by ELISA. The subsequent competitive ELISA uses the microtiter plate format. The antigen is bound to the solid phase of the microtiter plate. The derivatized analyte concentrations in the standards, controls and samples compete with the solid phase bound analytes for a fixed number of antibody binding sites. After the system is in equilibrium, free antigen and free antigen-antibody complexes are removed by washing. The antibody bound to the solid phase is detected by an anti-rabbit IgG-peroxidase conjugate using TMB as a substrate resulting in a colour reaction. The reaction is monitored at a wavelength of 450 nm.
Quantification of unknown samples is achieved by comparing their absorbance with a reference curve prepared with known standard concentrations. Manual processing of the ELISA is recommended. The use of automatic laboratory equipment is the responsibility of the user.
This product is not intended to clinical diagnoses.

1.2 Background

Glutamate, also known as glutamic acid, is one of the most important excitatory neurotransmitter in the central nervous system (CNS). It is released presynaptically and it binds postsynaptically to specific receptors for glutamate. The enzyme glutamic acid decarboxylase is able to convert L-glutamate in the CNS by decarboxylation to γ-aminobutyric acid (GABA), which acts as an inhibitory neurotransmitter.

2. Procedural cautions, guidelines, warnings and limitations

2.1 Procedural cautions, guidelines and warnings

(1) This kit is intended for professional use only. Users should have a thorough understanding of this protocol for the successful use of this kit. Only the test instruction provided with the kit is valid and must be used to run the assay. Reliable performance will only be attained by strict and careful adherence to the instructions provided.
(2) The principles of Good Laboratory Practice (GLP) must be followed.
(3) In order to reduce exposure to potentially harmful substances, wear lab coats, disposable protective gloves and protective glasses where necessary.
(4) All kit reagents and specimens should be brought to room temperature and mixed gently but thoroughly before use. For dilution or reconstitution purposes, use deionized, distilled, or ultra-pure water. Avoid repeated freezing and thawing of reagents and specimens.
(5) The microplate contains snap-off strips. Unused wells must be stored at 2 – 8 °C in the sealed foil pouch with desiccant and used in the frame provided. Microtiter strips which are removed from the frame for usage should be marked accordingly to avoid any mix-up.
(6) Duplicate determination of sample is highly recommended.
(7) Once the test has been started, all steps should be completed without interruption. Make sure that the required reagents, materials, and devices are prepared for use at the appropriate time.
(8) Incubation times do influence the results. All wells should be handled in the same order and time intervals.
(9) To avoid cross-contamination of reagents, use new disposable pipette tips for dispensing each reagent, sample, standard and control.
(10) A standard curve must be established for each run.
(11) The controls should be included in each run and fall within established confidence limits. The confidence limits are listed in the QC-Report provided with the kit.
(12) Do not mix kit components with different lot numbers within a test and do not use reagents beyond expiry date as shown on the kit labels.
(13) Avoid contact with Stop Solution containing 0.25 M H_2SO_4. It may cause skin irritation and burns. In case of contact with eyes or skin, rinse off immediately with water.
(14) TMB substrate has an irritant effect on skin and mucosa. In case of possible contact, wash eyes with an abundant volume of water and skin with soap and abundant water. Rinse contaminated items before reuse.
(15) For information about hazardous substances included in the kit please refer to Safety Data Sheet (SDS). The Safety Data Sheet for this product is made available directly on the website of the manufacturer or upon request.
(16) Kit reagents must be regarded as hazardous waste and disposed of according to national regulations.
(17) In case of any severe damage to the test kit or components, the manufacturer has to be informed in writing, at the latest, one week after receiving the kit. Severely damaged single components must not be used for a test run. They must be stored properly until the manufacturer decides what to do with them. If it is decided that they are no longer suitable for measurements, they must be disposed of in accordance with national regulations.

2.2 Limitations

Any inappropriate handling of samples or modification of this test might influence the results.

2.2.1 Interfering substances and proper handling of specimens

Avoid excess of acid: excess of acid might exceed the buffer capacity of the dilution buffer.

A pH of 5.0 during the extraction is mandatory.

Urine

Please note the sample preparation stabilization of the urine sample! It cannot be excluded that high acid concentrations lead to incorrect results. Up to 20 µl 6 M HCl per 1 ml urine no influence on the results was observed.

2.2.2 Drug and food interferences

There are no known substances (drugs, food) which ingestion interferes with the measurement of glutamate level in the sample.

2.2.3 High-Dose-Hook effect

No hook effect was observed in this test.

3. Storage and stability

Store kit and reagents at 2 – 8 °C until expiration date. Do not use kit and components beyond the expiry date indicated on the kit labels. Once opened, the reagents are stable for 2 months when stored at 2 – 8 °C. Once the resealable pouch of the ELISA plate has been opened, care should be taken to close it tightly again including the desiccant.

4. Materials

4.1 Contents of the kit

BA D-0024	**REAC-PLATE**	**Reaction Plate –** ready to use
Content:	1 x 96 well plate, empty, in a resealable pouch	
BA D-0090	**FOILS**	**Adhesive Foil** – ready to use
Content:	Adhesive foils in a resealable pouch	
Number:	1 x 4 foils	
BA E-0030	**WASH-CONC** **50x**	**Wash Buffer Concentrate** – concentrated 50x
Content:	Buffer with a non-ionic detergent and physiological pH	
Volume:	1 x 20 ml/vial, purple cap	
BA E-0040	**CONJUGATE**	**Enzyme Conjugate** – ready to use
Content:	Goat anti-rabbit immunoglobulins conjugated with peroxidase	
Volume:	1 x 12 ml/vial, red cap	
Description:	Species is goat	
BA E-0055	**SUBSTRATE**	**Substrate** – ready to use
Content:	Chromogenic substrate containing 3,3',5,5'-tetramethylbenzidine, substrate buffer and hydrogen peroxide	
Volume:	1 x 12 ml/vial, black cap	
BA E-0080	**STOP-SOLN**	**Stop Solution** – ready to use
Content:	0.25 M sulfuric acid	
Volume:	1 x 12 ml/vial, grey cap	
BA E-2410	**AS** **GLUT**	**Glutamate Antiserum** – ready to use
Content:	Rabbit anti-glutamate antibody in buffer with proteins and non-mercury preservative, blue coloured	
Volume:	1 x 6 ml/vial, blue cap	

BA E-2413	ASSAY-BUFF **Assay Buffer** – ready to use
Content:	Buffer with alkaline pH
Volume:	1 x 20 ml/vial, yellow cap
Hazard pictograms:	 GHS08 GHS07
Signal word:	Danger
Hazardous ingredients:	Boric acid
Hazard statements:	H360FD May damage fertility. Suspected of damaging the unborn child.
Precautionary statements:	P201 Obtain special instructions before use. P280 Wear protective gloves, protective clothing, eye protection, face protection. P308+P313 IF exposed or concerned: Get medical advice/attention. P501 Dispose of contents/container to an authorised waste collection point.
Additional statements:	Restricted to professional users.
BA E-2428	EQUA-REAG **Equalizing Reagent** – lyophilized
Content:	Lyophilized protein
Volume:	1 vial, brown cap
Description:	Species is bovine
BA E-2431	GLUT **Glutamate Microtiter Strips** – ready to use
Content:	1 x 96 wells (12x8) antigen precoated microwell plate in a resealable pouch with desiccant
BA E-2442	EXTRACT-PLATE 48 **Extraction Plate** – ready to use
Content:	2 x 48 well plate, precoated with cation exchanger in a resealable pouch
BA E-2446	D-REAGENT **D-Reagent** – ready to use
Content:	Crosslinking agent in dimethylsulfoxide
Volume:	1 x 3 ml/vial, white cap
Hazard pictograms:	 GHS07
Signal word:	Warning
Hazardous ingredients:	Glutaraldehyde
Hazard statements:	H317 May cause an allergic skin reaction.
Precautionary statements:	P261 Avoid breathing mist/vapours/spray. P280 Wear protective gloves. P333+P313 If skin irritation or rash occurs: Get medical advice/attention. P501 Dispose of contents/container to an authorised waste collection point.
BA E-2458	Q-BUFFER **Q-Buffer** – ready to use
Content:	TRIS buffer
Volume:	1 x 20 ml/vial, white cap
BA E-2460	DILUENT **Diluent** – ready to use
Content:	Buffer with sodium acetate
Volume:	1 x 20 ml/vial, green cap

BA E-2787	NAOH	**NaOH** – ready to use
Content:	Sodium hydroxide solution	
Volume:	1 x 2 ml/vial, purple cap	
Hazard pictograms:	 GHS07	
Signal word:	Warning	

4.2 Calibration and Controls

Standards and Controls – ready to use

Cat. no.	Component	Colour/Cap	Concentration [µg/ml]	Concentration [µmol/l]	Volume/ Vial
BA E-2401	STANDARD A	white	0	0	4 ml
BA E-2402	STANDARD B	yellow	0.6	4.08	4 ml
BA E-2403	STANDARD C	orange	2	13.6	4 ml
BA E-2404	STANDARD D	blue	6	40.8	4 ml
BA E-2405	STANDARD E	grey	20	136	4 ml
BA E-2406	STANDARD F	black	60	408	4 ml
BA E-2451	CONTROL 1	green	Refer to QC-Report for expected value and acceptable range.		4 ml
BA E-2452	CONTROL 2	red			4 ml
Conversion:	glutamate [µg/ml] x 6.8 = glutamate [µmol/l]				
Content:	Acidic buffer with non-mercury preservatives, spiked with a defined quantity of glutamate.				

4.3 Additional materials required but not provided in the kit

- Water (deionized, distilled, or ultra-pure)
- Absorbent material (paper towel)

4.4 Additional equipment required but not provided in the kit

- Calibrated precision pipettes to dispense volumes between 10 – 100 µl; 12.5 ml
- Microtiter plate washing device (manual, semi-automated or automated)
- ELISA reader capable of reading absorbance at 450 nm and if possible 620 – 650 nm
- Microtiter plate shaker (shaking amplitude 3 mm; approx. 600 rpm)
- Vortex mixer

5. Sample collection, handling and storage

Various biological samples can be used for L-Glutamate determination. The assay was validated for human urine samples.

Urine

Spontaneous urine (second morning urine) stabilized with 10 µl 6 M HCl per 1 ml of urine sample should be used. The measurement results are related to the creatinine content of the sample.
Storage: up to 6 hours at 18 – 25 °C; up to 14 days at 2 – 8 °C; up to 6 months at < -15 °C. Repeated freezing and thawing should be avoided. Avoid exposure to direct sunlight.

6. Test procedure

Allow all reagents and samples to reach room temperature and mix thoroughly by gentle inversion before use. Number the Extraction Plate, Reaction Plate and microwell plates (Microtiter Strips which are removed from the frame for usage should be marked accordingly to avoid any mix-up). Duplicate determinations are recommended.
The binding of the antisera and of the enzyme conjugate and the activity of the enzyme are temperature dependent. The higher the temperature, the higher the absorption values will be. Varying incubation times will have similar influences on the absorbance. The optimal temperature during the enzyme immunoassay is between 20 – 25 °C.
If the product is prepared in parts, unused wells in Reaction and Extraction Plates should be covered to avoid contamination. After preparation, the used wells must be labelled to prevent double use.
During the overnight incubation at 2 – 8 °C with the antiserum, the temperature should be uniform all over the ELISA plate to avoid any drift and edge-effect.

⚠ *The use of a microtiter plate shaker with the following specifications is mandatory: shaking amplitude 3 mm; approx. 600 rpm. Shaking with differing settings might influence the results.*

6.1 Preparation of reagents and further notes

Wash Buffer
Dilute the 20 ml Wash Buffer Concentrate **WASH-CONC** **50X** with water to a final volume of 1000 ml.
Storage: 2 months at 2 – 8 °C

Equalizing Reagent
Reconstitute the **EQUA-REAG** with **12.5 ml** of **ASSAY-BUFF**.
Reconstituted Equalizing Reagent which is not used immediately has to be stored in aliquots for max. 2 months at < -15 °C and may be thawed only once.

D-Reagent
The **D-REAGENT** has a freezing point of 18.5 °C. Make sure that the **D-REAGENT** has reached room temperature and forms a homogeneous, crystal-free solution.

Glutamate Microtiter Strips
In rare cases residues of the blocking and stabilizing reagent can be seen in the wells as small, white dots or lines. These residues do not influence the quality of the product.

Extraction Plate
In rare cases residues of the cation exchanger can be seen in the wells as small, black dots or lines. These residues do not influence the quality of the product.

6.2 Preparation of samples

The Glutamate ELISA is a flexible test system for various biological sample types and volumes. It is not possible to give a general advice how to prepare the samples. However, the following basics should help the researcher to adapt the protocol to his specific needs:

- Avoid excess of acid: excess of acid might exceed the buffer capacity of the dilution buffer. A **pH of 5.0** during the extraction is mandatory.
- It is advisable to perform a **Proof of Principle** to determine the recovery of glutamate from the samples. Prepare a stock solution of glutamate. Add small amounts (to change the native sample matrix as less as possible) of the stock solutions to the sample matrix and check the recovery.
- The sample volume determines the sensitivity of this test. Determine the sample volume needed to determine glutamate in your sample by testing different amounts of sample volumes.
- If a sample volume **< 100 µl** is used, water (deionized, distilled, or ultra-pure) has to be added to a final **volume of 100 µl.**

If you need any support in establishing a protocol for your specific purposes, do not hesitate to contact the manufacturer directly!

6.3 Preparation of samples – Extraction

1.	Pipette **100 µl** of the **standards, controls** and **urine samples** into the appropriate wells of the **EXTRACT-PLATE** **48**.
2.	Add **100 µl** of the **DILUENT** to all wells. Cover plate with **FOILS** and **shake** for **10 min** at **RT** (20 – 25 °C) on a **shaker** (approx. 600 rpm).
3.	Use **25 µl** for the subsequent **derivatization**!

6.4 Derivatization

1.	Pipette **25 µl** of the **extracted standards, controls** and **urine samples** into the appropriate wells of the **REAC-PLATE**.
2.	Pipette **10 µl** of **NAOH** into all wells.
3.	Pipette **50 µl** of the **Equalizing Reagent** into all wells.
4.	Pipette **10 µl** of the **D-REAGENT** into all wells.
5.	Cover plate with **FOILS** and shake for **2 h** at **RT** (20 – 25 °C) on a **shaker** (approx. 600 rpm).
6.	Pipette **75 µl** of the **Q-BUFFER** into all wells.
7.	Shake for **10 min** at **RT** (20 – 25 °C) on a **shaker** (approx. 600 rpm).
8.	Use **25 µl** for the **ELISA!**

6.5 Glutamate ELISA

1.	Pipette **25 µl** of the **prepared standards, controls and urine samples** into the appropriate wells of the **Glutamate Microtiter Strips** Ш **GLUT.**
2.	Pipette **50 µl** of the **AS GLUT** into all wells and mix shortly.
3.	Cover plate with **FOILS** and incubate for **15 – 20 h** (overnight) at **2 – 8 °C**.
4.	Remove the foil. Discard or aspirate the content of the wells. Wash the plate **3 x** by adding **300 µl** of **Wash Buffer, discarding** the content and **blotting dry each time** by tapping the inverted plate on absorbent material.
5.	Pipette **100 µl** of the **CONJUGATE** into all wells.
6.	Incubate for **30 min** at **RT** (20 – 25 °C) on a **shaker** (approx. 600 rpm).
7.	Discard or aspirate the contents of the wells and wash the plate **3 x** by adding **300 µl** of **Wash Buffer, discarding** the content and **blotting dry each time** by tapping the inverted plate on absorbent material.
8. ⚠	Pipette **100 µl** of the **SUBSTRATE** into all wells and incubate for **20 – 30 min** at **RT** (20 – 25 °C) on a **shaker** (approx. 600 rpm). ***Avoid exposure to direct sunlight!***
9.	Add **100 µl** of the **STOP-SOLN** to each well and shake the microtiter plate to ensure a homogeneous distribution of the solution.
10.	**Read** the absorbance of the solution in the wells within 10 minutes, using a microplate reader set to **450 nm** (if available a reference wavelength between 620 nm and 650 nm is recommended).

7. Calculation of results

Measuring range	Glutamate	
	Urine	0.26 – 60 µg/ml

The standard curve, which can be used to determine the concentration of the unknown samples, is obtained by plotting the absorbance readings (calculate the mean absorbance) of the standards (linear, y-axis) against the corresponding standard concentrations (logarithmic, x-axis) using a concentration of 0.001 µg/ml for Standard A (this alignment is mandatory because of the logarithmic presentation of the data). Use non-linear regression for curve fitting (e.g. 4-parameter, marquardt).

⚠ *This assay is a competitive assay. This means: the OD-values are decreasing with increasing concentrations of the analyte. OD-values found below the standard curve correspond to high concentrations of the analyte in the sample and have to be reported as being positive.*

The concentrations of the samples (100 µl undiluted sample used) and controls can be read directly from the standard curve.

⚠ In case < 100 µl sample volume was used, concentrations of the samples taken from the standard curve have to be multiplied by a correction factor:

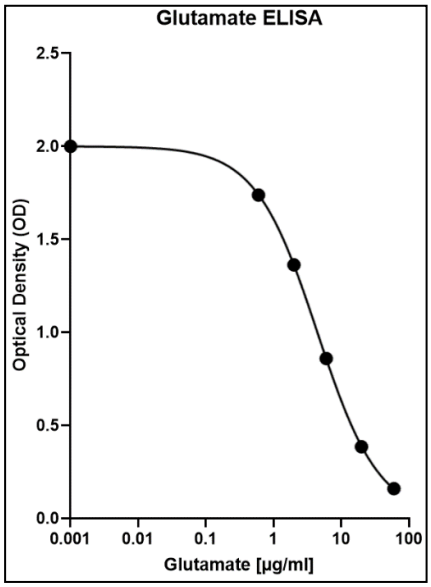
$$\text{Correction factor} = \frac{\text{100 µl (volume of standards)}}{\text{sample volume (µl)}}$$

Samples found with concentrations higher than the highest standard (Standard F) should be diluted accordingly with water (deionized, distilled, or ultra-pure) and must be re-assayed. For the calculation of the concentrations this dilution factor has to be taken into account.

Conversion:
Glutamate [µg/ml] x 6.8 = Glutamate [µmol/l]

7.1 Typical standard curve

⚠ *Example: Do not use for calculation!*



8. Control samples

The confidence limits of the kit controls are indicated on the QC-Report.

9. Assay characteristics

Various biological samples can be used for L-Glutamate determination. The assay was validated for human urine samples.

9.1 Performance data

Analytical Sensitivity	
	Glutamate
Limit of Blank (LOB)	0.11 µg/ml
Limit of Detection (LOD)	0.17 µg/ml
Limit of Quantification (LOQ)	0.26 µg/ml

Analytical Specificity (Cross Reactivity)	
Substance	**Cross Reactivity [%]**
	Glutamate
L-Glutamine	< 0.4
Glycine	< 0.4
β-Alanine	< 0.4
L-Alanine	< 0.4
L-Aspartic Acid	< 0.4
GABA	< 0.4
5-Amino-n-valeric Acid	< 0.4

Precision							
Intra-Assay				**Inter-Assay**			
Sample	n	Mean ± SD [µg/ml]	CV [%]	Sample	n	Mean ± SD [µg/ml]	CV [%]
1	10	0.8 ± 0.1	10.8	1	13	1.7 ± 0.24	14.3
2	10	1.3 ± 0.1	8.7	2	14	5.0 ± 0.57	11.4
3	10	2.2 ± 0.1	6.3	3	14	10.6 ± 0.73	6.9
4	10	4.8 ± 0.2	4.0	4	13	3.0 ± 0.43	14.2
5	10	12.5 ± 0.6	4.6	5	14	5.6 ± 0.71	12.5
6	10	39.7 ± 2.2	5.6	6	14	10.0 ± 0.87	8.7

Lot-to-Lot			
	Sample	Mean ± SD [µg/ml]	CV [%]
Glutamate in urine (n=3)	1	13.3 ± 1.2	9.4
Glutamate in artificial matrix (n = 3)	2	5.0 ± 0.5	10.1

Recovery			
	Range [µg/ml]	Mean [%]	Range [%]
Urine	1.25 – 41.0	102	97 – 108

Linearity			
	Serial dilution up to	Mean [%]	Range [%]
Urine	1:64	105	94 – 113

9.2 Metrological Traceability

The values assigned to the standards and controls of the Glutamate ELISA are traceable to SI Units by weighing with quality-controlled analyte.

Standards and Controls	
	Uncertainty [%]
Glutamate	1.4

Glutamate ELISA	
Concentration [µg/ml]	Expanded Uncertainty [%] k = 2*
1.7	28.7
5	23.0
10.6	14.1

* This defines an interval about the measured result that will include the true value with a probability of 95%.

10. References/Literature

1. Bieger, W.P., *NeuroStress Guide*. 2011.
2. Bustillo, J.R., *Use of proton magnetic resonance spectroscopy in the treatment of psychiatric disorders: a critical update.* Dialogues Clin Neurosci, 2013. **15**(3): p. 329-37.
3. Duman, R.S., G. Sanacora, and J.H. Krystal, *Altered Connectivity in Depression: GABA and Glutamate Neurotransmitter Deficits and Reversal by Novel Treatments.* Neuron, 2019. **102**(1): p. 75-90.
4. Femenia, T., et al., *Dysfunctional hippocampal activity affects emotion and cognition in mood disorders.* Brain Res, 2012. **1476**: p. 58-70.
5. Flasnoecker, M., *Reise aus dem Stress - Körper, Geist und Psyche stärken*. 2015: W. Zuckschwerdt Verlag GmbH.
6. Frisardi, V., F. Panza, and A.A. Farooqui, *Late-life depression and Alzheimer's disease: the glutamatergic system inside of this mirror relationship.* Brain Res Rev, 2011. **67**(1-2): p. 344-55.
7. Gao, S.F. and A.M. Bao, *Corticotropin-releasing hormone, glutamate, and gamma-aminobutyric acid in depression.* Neuroscientist, 2011. **17**(1): p. 124-44.
8. Harris, R.E. and D.J. Clauw, *Imaging central neurochemical alterations in chronic pain with proton magnetic resonance spectroscopy.* Neurosci Lett, 2012. **520**(2): p. 192-6.
9. Kendell, S.F., J.H. Krystal, and G. Sanacora, *GABA and glutamate systems as therapeutic targets in depression and mood disorders.* Expert Opin Ther Targets, 2005. **9**(1): p. 153-68.
10. Krystal, J.H., et al., *Glutamate and GABA systems as targets for novel antidepressant and mood-stabilizing treatments.* Mol Psychiatry, 2002. **7 Suppl 1**: p. S71-80.
11. Lener, M.S., et al., *Glutamate and Gamma-Aminobutyric Acid Systems in the Pathophysiology of Major Depression and Antidepressant Response to Ketamine.* Biol Psychiatry, 2017. **81**(10): p. 886-897.
12. Sanacora, G., G. Treccani, and M. Popoli, *Towards a glutamate hypothesis of depression: an emerging frontier of neuropsychopharmacology for mood disorders.* Neuropharmacology, 2012. **62**(1): p. 63-77.
13. Strienz, J., *Nebennierenunterfunktion - Stress stört die Hormonbalance*. 3 ed. 2019: W. Zuckschwerdtverlag München.

For updated literature or any other information please contact your local supplier.

11. Changes

Version	Release Date	Chapter	Change
17.0-r	2024-05-28	4.1 9.1 9.2	- Hazard labelling updated according to SDS - Lot-to-Lot added - Chapter Metrological Traceability added
18.0-r	2024-09-30	9.1	- Lot-to-lot updated

SYMBOLS USED WITH DEMEDITEC ASSAYS

Symbol	English	Deutsch	Française	Espanol	Italiano
CE	European Conformity	CE-Konformitäts-kennzeichnung	Conforme aux normes européennes	Conformidad europea	Conformità europea
	Consult instructions for use	Gebrauchsanweisung beachten	Consulter les instructions d'utilisation	Consulte las Instrucciones	Consultare le istruzioni per l'uso
IVD	In vitro diagnostic device	In-vitro-Diagnostikum	utilisation Diagnostic in vitro	Diagnóstico in vitro	Per uso Diagnostica in vitro
RUO	For research use only	Nur für Forschungszwecke	Seulement dans le cadre de recherches	Sólo para uso en investigación	Solo a scopo di ricerca
REF	Catalogue number	Katalog-Nr.	Référence	Número de catálogo	No. di catalogo
LOT	Lot. No. / Batch code	Chargen-Nr.	No. de lot	Número de lote	Lotto no
	Contains sufficient for <n> tests/	Ausreichend für "n" Ansätze	Contenu suffisant pour "n" tests	Contenido suficiente para <n> ensayos	Contenuto sufficiente per "n" saggi
	Note warnings and precautions	Warnhinweise und Vorsichtsmaßnahmen beachten	Avertissements et mesures de précaution font attention	Tiene en cuenta advertencias y precauciones	Annoti avvisi e le precauzioni
	Storage Temperature	Lagerungstemperatur	Température de conservation	Temperatura de conservacion	Temperatura di conservazione
	Expiration Date	Mindesthaltbarkeits-datum	Date limite d'utilisation	Fecha de caducidad	Data di scadenza
	Legal Manufacturer	Hersteller	Fabricant	Fabricante	Fabbricante
Distributed by	Distributed by	Vertrieb durch	Distribution par	Distribución por	Distribuzione da parte di
V<x>	Version	Version	Version	Versión	Versione
	Single-use	Einmalverwendung	À usage unique	Uso único	Uso una volta