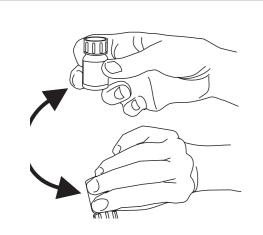


Reagents for Microdialysis Analyzer



Dissolve contents completely by gently turning the bottle or reagent casette upside-down a few times.

Preparation and stability of solution.

- 1. Unscrew the cap with the membrane from the reagent bottle. Remove and discard the rubber stopper.
- 2. Transfer the contents of the buffer bottle to the reagent bottle.
- 3. Fasten the cap with the membrane on the reagent bottle, without Rubber stopper.
- 4. Dissolve contents completely by gently turning the bottle upside-down at least ten times. Let the reagent stand and equilibrate in room temperature for at least 30 minutes prior to use. Turn the bottle a couple of times before placing the reagent in the Microdialysis Analyzer

Reconstituted reagent is stable for five days in the instrument and for two weeks when stored, protected from daylight, at +2 to +8°C

Ordering information	Ref. No.
Reagent kit Incl. Glucose 1x6mL, Lactate 1x6mL, Pyruvate 1x6m, Glycerol 1x6mL, Calibrator A 1x6 mL	P000011
L-P-G Reagent kit Incl. Glucose 1x6mL, Lactate 1x6mL, Pyruvate 1x6m, Calibrator A 1x6 mL	8010361
Reagent set A Incl. Glucose 1x6mL, Lactate 1x6mL, Pyruvate 1x6m, Glycerol 1x6mL, Calibrator A 1x6 mL	8002163
Reagent set B Incl. Glucose 1x6mL, Lactate 1x6mL, Pyruvate 1x6m, Glycerol 1x6mL, Glutamate 1x4m L, Calibrator A 1x6 mL	8002164
Reagent set C Incl. Glucose 1x6mL, Lactate 1x6mL, Pyruvate 1x6m, Calibrator A 1x6 mL	8002165



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Reagent kits & sets



Reagents for Microdialysis Analyzer

M Dialysis offers both Reagent kits and Reagents sets for the ISCUSflex Microdialysis Analyzer.

- Reagents sets are for routine use, comes in a casette and simplyfies usage with a casette code.
- Reagent kits are more for research, are loaded as single reagents in the analyzer.

This product sheet informs about the Calibrator A and the different Reagents that are available in the various Reagents sets and the kits.

Glucose

Colorimetric method for the quantitative determination of glucose in Microdialysates.

Measuring principle

Glucose is enzymatically oxidised by glucose oxidase (GOD). The hydrogen peroxide formed reacts with phenol and 4-amino-antipyrine. This reaction is catalyzed by peroxidase (POD) and yields the red-violet colored quinoneimine. The rate of formation is measured photometrically at 546 nm and is proportional to the glucose concentration.

Glucose + O_2 + $H_2O \xrightarrow{GOD}$ Gluconic acid + H_2O_2

 $2\,\mathrm{H_2O_2}$ + Aminoantipyrine + Phenol $\stackrel{\mathrm{pob}}{\longrightarrow}$ Quinoneimine + $4\,\mathrm{H_2O}$ Linear range: 0.1 - 25 mmol/L

Lactate

Colorimetric method for the quantitative determination of lactate in Microdialysates.

Measuring principle

Lactate is enzymatically oxidised by lactate oxidase. The hydrogen peroxide formed reacts with 4-chlorophenol and 4-amino-antipyrine. This reaction is catalyzed by peroxidase (POD) and yields the red-violet colored quinoneimine. The rate of formation is measured photometrically at 546 nm and is proportional to the lactate concentration.

Lactate + O_2 + $H_2O \longrightarrow Pyruvate + <math>H_2O_2$

 H_2O_2 + 4-Chlorophenol + 4-Aminoantipyrine $\stackrel{\tiny POD}{\longrightarrow}$

Quinoneimine + 2 H₂O + HCl

Default linear range: 0.1 - 12 mmol/L

Assay Conditions

Sample volume: 0.5 µL Reagent Volume:14.5 µL Wavelength: 546 nm Linear Range: 0.1-25 mmol/L

Assay Conditions

 $\begin{array}{lll} \text{Sample volume:} & 0.4\,\mu\text{L} & \text{or} \ 0.8\,\mu\text{L} \\ \text{Reagent Volume:} & 14.6\,\mu\text{L} & \text{or} \ 14.2\,\mu\text{L} \end{array}$

Wavelength: 546 nm

Linear Range: 0.1-12 or 0.02 - 2.5 mmol/L



Glutamate

Colorimetric method for the quantitative determination of Glutamate in Microdialysates.

Measuring principle

Glutamate is enzymatically oxidized by glutamate oxidase (GltOx). The hydrogen peroxide formed reacts with N-ethyl-N-(2-hydroxy-3-sulfopropyl)-m-toluidine and 4-amino-antipyrine. This reaction is catalyzed by peroxidase (POD) and yields the red-violet colored quinonediimine. The rate of formation is measured photometrically at 546 nm and is proportional to the glutamate.

Glutamate + $O_2 \xrightarrow{GITOX} 2$ -oxoglutarate + H_2O_2 H_2O_2 + 4-amino-antipyrine + $TOOS \xrightarrow{POD}$

Quinonediimine + 4 H_oO

Linear range: 1 - 150 µmol/L

Assay Conditions

Sample volume: 1.3 µL Reagent Volume: 7.7 µL Wavelength: 546 nm Linear Range: 1-150 µmol/L

Glycerol

Colorimetric method for the quantitative determination of glycerol in Microdialysates.

Measuring principle

Glycerol is phosphorylated by adenosine triphosphate (ATP) and glycerol kinase (GK) to glycerol-3-phosphate, which is subsequently oxidized in the presence of glycerol-3-phosphate oxidase (GPO). The hydrogen peroxide formed reacts with 3,5-dichloro-2-hydroxy-benzene sulphonic acid (DCHBS) and 4-amino-antipyrine. This reaction is catalyzed by peroxidase (POD) and yields the red-violet colored quinoneimine. The rate of formation is measured photometrically at 546 nm and is proportional to the glycerol concentration.

Glycerol + ATP Glycerol-3-phosphate + ADP

Glycerol-3-phosphate $+ O_2^{GPO} H_2 O_2 +$ dihydroxyacetone phosphate

 $H_2O_2 + DCHBS + 4-Aminoantipyrine \xrightarrow{POD} 2 H_2O + ACSB + HCI$

Default linear range: 10 - 1500 µmol/L

Assay Conditions

 $\begin{array}{ll} \text{Sample volume: } 0.4\,\mu\text{L} & \text{or } 2.0\,\mu\text{L} \\ \text{Reagent Volume: } 14.6\,\mu\text{L} & \text{or } 13.0\,\mu\text{L} \end{array}$

Wavelength: 546 nm

Linear Range: 10 - 1500 µmol/L or 2-500 µmol/L

Reagents for Microdialysis Analyzer



Pyruvate

Colorimetric method for the quantitative determination of pyruvate in Microdialysates.

Measuring principle

Pyruvate is enzymatically oxidized by pyruvate oxidase (PyrOx). The hydrogen peroxide formed reacts with N-ethyl-N-(2-hydroxy-3-sulfopropyl)-m-toluidine and 4-amino-antipyrine. This reaction is catalyzed by peroxidase (POD) and yields the red-violet colored quinonedimine. The rate of formation is measured photometrically at 546 nm and is proportional to the pyruvate concentration.

Pyruvate + inorganic phosphate + O_2 \xrightarrow{POD} acetylphosphate + CO_2 + H_2O_2 \xrightarrow{POD}

 $\rm H_2O_2 * 4\text{-}amino\text{-}antipyrine} * TOOS$

Quinonediimine + 4 H₂O

Default linear range: 10 - 300 µmol/L

Calibrator A

Calibrator for Microdialysis Analyser

For calibration of

P000023 Glucose Reagent P000024 Lactate Reagent P000025 Glycerol Reagent P000026 Urea Reagent P000063 Pyruvate Reagent P000064 Glutamate Reagent

Content

Analyte Concentration
Glucose 5.55 mmol/L
Lactate 2.5 mmol/L
Glycerol 475 µmol/L
Urea 13.3 mmol/L
Pyruvate 250 µmol/L
Glutamate 25 µmol/L

Assay Conditions

Sample volume: 0.5 μL or 2.0 μL Reagent Volume: 14.5 μL or 13.0 μL

Wavelength: 546 nm

Linear Range: 10 - 1500 µmol/L or 10 - 300 µmol/L