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Datasheet for ABIN2345135

OxiSelect™ In Vitro Nitric Oxide (Nitrite / Nitrate) Assay Kit (Fluorometric)

Overview

Quantity:	100 tests
Reactivity:	Others
Application:	Biochemical Assay (BCA)

Product Details

Purpose:	Conversion of nitrate to nitrite by nitrate reductase (A), followed by the reaction with the fluorescent probe DAN (2,3-diaminonaphthalene) (B).
Brand:	OxiSelect™
Sample Type:	Serum, Plasma, Saliva, Urine
Detection Method:	Fluorometric
Sensitivity:	500 nM
Characteristics:	OxiSelect™ Nitric Oxide (Nitrite/Nitrate) Assay Kit is a simple, fluorometric assay that quantitatively measures NO in various samples by NO ⁻² /NO ⁻³ determination. First, the nitrate (NO ⁻³) in the sample is converted to nitrite (NO ⁻²) by nitrate reductase enzyme (Figure 2). Next, total nitrite is detected with the fluorescent probe DAN (2,3-diaminonaphthalene). Each kit provides sufficient reagents to perform up to 100 assays using a 96-well microtiter plate format, including blanks, standards and unknown samples. The kit is suitable for serum, plasma, urine, saliva, lysates, and media (see Preparation of Samples) with detection sensitivity limit of ~ 500 nM (in 50 µL sample volume). Figure 1: Oxidation of L-arginine to nitrate.
Components:	<ol style="list-style-type: none"> 1. Nitrite Standard : One 500 µL vial of 14 mM sodium nitrite. 2. Nitrate Standard : One 500 µL vial of 14 mM sodium nitrate. 3 3. Nitrite Assay Buffer : Three 1.4 mL vials.

Product Details

4. Total Nitrate/Nitrite Assay Buffer : Three 1.4 mL vials.
5. Enzyme Cofactor : One 20 µL amber vial.
6. 100X DAN Fluorometric Probe : One 25 µL amber vial.
7. 2X Fluorometric Probe Diluent : One 1.5 mL vial.
8. Stop Solution : One 1.1 mL vial.

Material not included:

1. Deionized or distilled water
2. PBS for sample dilution
3. 10 kDa MWCO ultrafilter (for high protein content samples)
4. Standard 96-well fluorescence black microtiter plate
5. Microtiter plate shaker
6. 10 µL to 1000 µL adjustable single channel micropipettes with disposable tips
7. 50 µL to 300 µL adjustable multichannel micropipette with disposable tips
8. Multichannel micropipette reservoir
9. Fluorescence microplate reader capable of reading excitation in the 360-365 nm range and emission in the 430-450 nm range.

Target Details

Background:

Reactive nitrogen intermediates (RNI) and reactive oxygen intermediates (ROI) work together to damage cells and have been implicated in the pathogenesis of several disease states. RNI are a family of molecules derived from nitric oxide (NO) and superoxide anion (O^{-2}), produced via nitric oxide synthase (NOS) and NADPH oxidase, respectively. Nitric oxide is an established mediator in vascular diseases, diabetes, renal ischemia, atherosclerosis, inflammatory diseases, and cancer. However, because of its extremely short half life, direct quantitation of NO production remains challenging and unsuitable for most detection systems. Therefore, the common method for measurement of total NO has become the sum of its final oxidized products, nitrite (NO^{-2}) and nitrate (NO^{-3}) (Figure 1).

Application Details

Application Notes:

Optimal working dilution should be determined by the investigator.

Comment:

- Measure NO in various samples by NO_2^-/NO_3^- -determination
- Suitable for serum, plasma, urine, saliva, lysates, and media

Reagent Preparation:

- Nitrite and Nitrate Standards should be thawed/maintained at 4 °C during assay preparation. Both are stable for 2 weeks at 4 °C. For longer term storage, each should be aliquoted and frozen at -80 °C to avoid multiple freeze/thaws.
- Nitrite Assay Buffer and Total Nitrate/Nitrite Assay Buffer should be thawed/maintained at 4 °C during assay preparation. Both are stable for 1 week at 4 °C. Any unused material should

be aliquoted and frozen at -80 °C to avoid multiple freeze/thaws. Note: This component is provided in multiple tubes to minimize multiple freeze/thaws.

- Enzyme Cofactor should be thawed/maintained at 4 °C during assay preparation. The stock solution is stable for 1 week at 4 °C. Immediately before use, dilute the Enzyme Cofactor 1:100 with deionized water and mix well. Do not store diluted solutions. Any unused stock material should be aliquoted and frozen at -80 °C to avoid multiple freeze/thaws.
- 2X Fluorometric Probe Diluent should be thawed/maintained at room temperature during assay preparation. The stock solution is stable for 1 week at room temperature. Dilute the 2X 4 Fluorometric Probe Diluent with deionized water (1:1) and mix well. Any unused stock material should be aliquoted and frozen at -80 °C to avoid multiple freeze/thaws.
- 100X DAN Fluorometric Probe should be thawed/maintained at room temperature during assay preparation. The stock solution is stable for 1 day at room temperature. Immediately before use, dilute the 100X DAN Fluorometric Probe 1:100 with 1X Fluorometric Probe Diluent (see above) and mix well. Do not store diluted solutions. Any unused stock material should be aliquoted and frozen at -80 °C to avoid multiple freeze/thaws.
- Stop Solution should be thawed/maintained at room temperature during assay preparation. The stock solution is stable for 1 week at room temperature. Any unused stock material should be aliquoted and frozen at -80 °C to avoid multiple freeze/thaws. Preparation of Nitrite or Nitrate Standards
- Thaw the nitrite or nitrate standards at 4 °C and mix well. Freshly prepare a dilution series of standard in the concentration range of 0 µM - 140 µM by diluting the standard stock solution (provided at 14 mM) in deionized water (Table 1) or desired sample buffer. 14 mM Nitrite or DI Water or Nitrite or Nitrate Standard Nitrate Standard Desired Standard Tubes (µL) Buffer (µL) (µM) 1 10 990 140 2 500 of Tube #1 500 70 3 500 of Tube #2 500 35 4 500 of Tube #3 500 17.5 5 500 of Tube #4 500 8.75 6 500 of Tube #5 500 4.38 7 500 of Tube #6 500 2.19 8 500 of Tube #7 500 1.09 9 500 of Tube #8 500 0.547 10 0 500 0 Table 1. Preparation of Nitrite or Nitrate Standards

Sample Preparation:

- Plasma: Collect plasma with an anticoagulant such as heparin, citrate or EDTA and mix by inversion. Samples must then be filtered through a 10 kDa MWCO ultrafilter before assaying (to reduce protein interference and turbidity). Use immediately or store at -80 °C. Note: Citrate or EDTA plasma is recommended. Heparin is known to interfere with the assay when samples contain > 1 IU/mL, however, ultrafiltering will remove average MW heparin (15 kDa and larger). If low MW heparin is used, a 3 kDa MWCO ultrafilter should be used.
- Serum: Collect serum with no anticoagulant. Samples must then be filtered through a 10 kDa MWCO ultrafilter before assaying (to reduce protein interference and turbidity). Use immediately or store at -80 °C. 5
- Urine: Samples must be filtered through a 10 kDa MWCO ultrafilter before assaying (to reduce protein interference and turbidity). Urine must be diluted before assaying (typically 1:10 or greater in PBS). Use immediately or store at -80 °C.
- Saliva: Samples must then be filtered through a 10 kDa MWCO ultrafilter before assaying (to reduce protein interference and turbidity). Use immediately or store at -80 °C.
- Culture Media: Media known to contain high levels of nitrite/nitrate (e.g. RPMI) should be avoided. Remove particulates/debris in samples by centrifugation or filtration. To account for

any background caused by media components, nitrite/nitrate standards should be prepared in the same media. Samples containing phenol red do not interfere with the assay.

- Cell and Tissue Lysates: Remove particulates/debris in samples by centrifugation or filtration. Samples must then be filtered through a 10 kDa MWCO ultrafilter before assaying (to reduce protein interference and turbidity). Use immediately or store at -80 °C. Potential Interference from Sample Components To accurately determine the degree of interference, nitrate and nitrite standard curves should be prepared in water vs. sample buffer. Dilution of the buffer, and ultimately samples, may be required to completely eliminate interference. However, even with some interference, accurate quantitation can be achieved by running standards in the same buffer as samples, although kit sensitivity may be compromised.

Assay Procedure:

Note: This kit may be used for measurement of nitrite only, or for total nitrite plus nitrate through nitrate reduction. If desired, nitrate only levels in samples may be calculated by subtracting the nitrite only level from the total nitrite plus nitrate.

I. Measurement of Nitrite only Note: Each nitrite standard and sample should be assayed in duplicate or triplicate. A freshly prepared standard curve should be used each time the assay is performed.

1. Add 50 µL of the nitrite standards, samples, or blanks to the 96-well fluorescence microtiter plate.
2. Maintain the Nitrite Assay Buffer and Cofactor at 4 °C. According to Table 2 (below), prepare the desired volume of Reaction Mixture (based on the number of tests) in the following sequence: a. In a tube, add the appropriate volume of Nitrite Assay Buffer. b. Next, add the corresponding volume of diluted, Enzyme Cofactor Solution (see Preparation of Reagents). Note: Enzyme Cofactor is diluted 1:100 in deionized water immediately before use. c. Mix well and immediately use. 6 # of Tests in Nitrite Assay Diluted Final Volume of 96-well Plate
Buffer Cofactor Reaction Mixture (50 µL/test) (mL) Solution (mL) (mL) 100 4 1 5 50 2 0.5 2.5 25 1 0.25 1.25 Table
3. Preparation of Reaction Mixture
4. Transfer 50 µL of the above Reaction Mixture to each well (already containing 50 µL of nitrite standard or sample).
5. Cover the plate wells to protect the reaction from light.
6. Incubate at room temperature for 1 hour on an orbital shaker.
7. Add 20 µL of diluted, DAN Fluorometric Probe to each well (see Preparation of Reagents). Note: DAN Fluorometric Probe is diluted 1:100 in 1X Fluorometric Probe Diluent immediately before use.
8. Incubate the plate at room temperature for 10 minutes on an orbital shaker, protected from light.
9. Immediately add 10 µL of Stop Solution to each well.
10. Incubate the plate at room temperature for 10 minutes on an orbital shaker, protected from light.
11. Read the plate with a fluorescence microplate reader equipped for excitation in the 360-365 nm range and for emission in the 430-450 nm range.

12. Calculate the concentration of nitrite within samples by comparing the sample absorbance to the standard curve. Negative controls (without nitrite) should be subtracted.

II. Measurement of Total Nitrate/Nitrite via Nitrate Reduction Note: Each nitrate standard and sample should be assayed in duplicate or triplicate. A freshly prepared standard curve should be used each time the assay is performed.

1. Add 50 μ L of the nitrate standards, samples, or blanks to the 96-well fluorescence microtiter plate.

2. Maintain the Total Nitrate/Nitrite Assay Buffer and Cofactor at 4 °C. According to Table 3 (below), prepare the desired volume of Reaction Mixture (based on the # of tests) in the following sequence: a. In a tube, add the appropriate volume of Total Nitrate/Nitrite Assay Buffer. b. Next, add the corresponding volume of diluted, Enzyme Cofactor Solution (see Preparation of Reagents). Note: Enzyme Cofactor is diluted 1:100 in deionized water immediately before use. c. Mix well and immediately use. 7 # of Tests in Total Diluted Final Volume of 96-well Plate Nitrate/Nitrite Cofactor Reaction Mixture (50 μ L/test) Assay Buffer Solution (mL) (mL) (mL) 100 4 1 5 50 2 0.5 2.5 25 1 0.25 1.25 Table

3. Preparation of Reaction Mixture

4. Transfer 50 μ L of the above Reaction Mixture to each well (already containing 50 μ L of nitrate standard or sample).

5. Cover the plate wells to protect the reaction from light.

6. Incubate at room temperature for 1 hour on an orbital shaker.

7. Add 20 μ L of diluted, DAN Fluorometric Probe to each well (see Preparation of Reagents). Note: DAN Fluorometric Probe is diluted 1:100 in 1X Fluorometric Probe Diluent immediately before use.

8. Incubate the plate at room temperature for 10 minutes on an orbital shaker, protected from light.

9. Immediately add 10 μ L of Stop Solution to each well.

10. Incubate the plate at room temperature for 10 minutes on an orbital shaker, protected from light.

11. Read the plate with a fluorescence microplate reader equipped for excitation in the 360-365 nm range and for emission in the 430-450 nm range.

12. Calculate the concentration of nitrate within samples by comparing the sample absorbance to the standard curve. Negative controls (without nitrate) should be subtracted. 8

Restrictions: For Research Use only

Handling

Storage: -80 °C

Storage Comment: Store the entire kit at -80°C. Avoid multiple freeze/thaws by aliquoting. The Enzyme Cofactor and 100X DAN Fluorometric Probe are light sensitive and should be maintained in amber tubes.