ELISA PRODUCT INFORMATION & MANUAL

CXCL8/IL-8 ELISA Development Kit

NBP3-11755

Enzyme-linked Immunosorbent Assay for quantitative detection. For research use only.
Not for diagnostic or therapeutic procedures.

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CXCL8/IL-8 ELISA Development Kit

ELISA Development Kit for quantitative determination of native monkey CXCL8/IL-8 in solution, e.g. cell supernatant.

The kit includes		NBP3-11755 for 6 plates
Capture mAb:	MT8H6 (0.5 mg/ml)	300 µl
Detection mAb:	26E5, biotinylated (0.5 mg/ml)	50 µl
Streptavidin-HRP		80 µl
Recombinant human IL-8 (CXCL8) ELISA standard		1 vial
Standard reconstitution buffer A5		1 ml

To ensure total recovery of the stated quantity, vials have been overfilled.

Shipping and storage

Shipped at ambient temperature. All reagents should be stored at 4-8 °C upon receipt, except the standard which should be stored at -20 °C. Antibodies are supplied in sterile-filtered PBS with sodium azide (0.02%). Streptavidin-HRP is supplied in PBS with 0.002% Kathon CG. The expiry date indicates how long unopened products, stored according to instructions, are recommended for use.

General and Preparations

Specificity

The kit contains a matched pair of monoclonal antibodies (mAbs) specific for native and recombinant human CXCL8/IL-8 The mAbs cross-react with CXCL8/IL-8from non-human primates. The mAbs also cross-react with CXCL8/IL-8 from cow, thus the use of bovine serum in cell-cultures is not recommended.

Standard range

8-800 pg/ml

Calibration

No international standard exists for calibration.

Analysis of serum and plasma samples

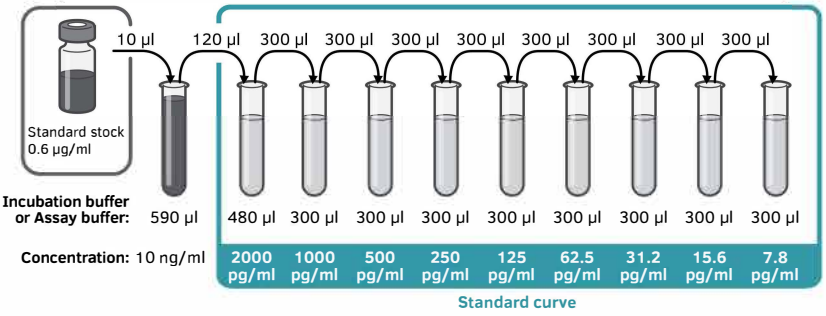
Analysis of serum/plasma requires the use of Assay buffer. The Assay buffer blocks heterophilic antibodies, commonly found in serum/plasma, from cross-linking the assay antibodies, thereby preventing false positive read-outs. The Assay buffer should be used for dilution of standard, samples, and detection antibody.

Reconstitution of ELISA standard

Reconstitute the ELISA standard to a stock solution of 0.6 µg/ml by adding 1 ml of the standard reconstitution buffer. Allow the standard to dissolve for 5 minutes and mix thoroughly. The standard should be kept in aliquots at -20 °C. Avoid repeated freeze-thaw cycles.

Preparation of standard curve

Prepare within 30 minutes of use. Volumes are sufficient for duplicates.



Protocol

Day 1

1. Add 100 µl/well of capture mAb MT8H6 diluted to 2 µg/ml in PBS, pH 7.4. Use high protein binding ELISA plates. Incubate overnight at 4-8 °C.

Day 2

2. Empty the plate and add 200 µl/well of PBS with 0.05% Tween 20 and 0.1% BSA (incubation buffer) to block the plate. Incubate for 1 hour at room temperature.

3. Wash the plate 5 times with PBS containing 0.05% Tween 20 (300 µl/well).

4. Add 100 µl/well of samples or standards diluted in incubation buffer or Assay buffer. Include assay background control, i.e. wells without standard. Incubate for 2 hours at room temperature.

5. Wash as above.

6. Add 100 µl/well of detection mAb 26E5-biotin diluted to 0.1 µg/ml in incubation buffer or Assay buffer. Incubate for 1 hour at room temperature.

7. Wash as above.

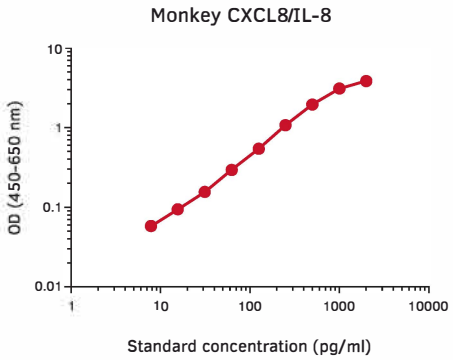
8. Add 100 µl/well of Streptavidin-HRP diluted 1:1000 in incubation buffer. Incubate for 1 hour at room temperature. Please note that sodium azide used in buffers will inhibit HRP activity.

9. Wash as above.

10. Add 100 µl/well of TMB substrate and incubate for 15 minutes.

11. Add 100 µl/well of 0.2 M H_2SO_4 to stop the reaction.

12. Measure the optical density in an ELISA reader at 450 nm within 15 min. Preferably use a reader capable of subtracting a reference wavelength of between 570 and 650 nm. Representative standard curve shown below.



Updated on 2022/02/23

Quality management system complies with the standards
ISO 9001:2015 & ISO 13485:2016.

The products are for research use only.