

General Octreotide (OT) ELISA Kit | OT elisa kit

General Octreotide (OT) ELISA Kit

CATALOG #
AAA161553

REACTIVITY
General

PRODUCT SYNONYMS

Octreotide (OT); General Octreotide (OT) ELISA Kit; OT ELISA kit (General species); Sandostatin; OT elisa kit

PRODUCT SPECIFICATIONS

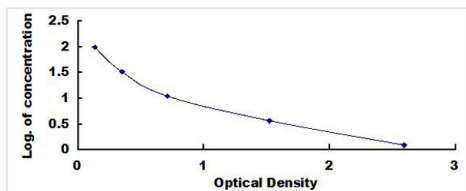
Reactivity	General
Specificity	This assay has high sensitivity and excellent specificity for detection of Octreotide (OT). No significant cross-reactivity or interference between Octreotide (OT) and analogues was observed. Reactive with General species OT/Octreotide
Assay Type	Competitive Inhibition
Samples	Serum, Plasma, Tissue homogenates, Cell lysates, Cell culture supernates and other biological fluids
Detection Method	Colormetric
Featured Series	CE kit
Detection Range	1.23-100ng/mL
Sensitivity	0.53ng/mL
Sample Size	50uL
Assay Time	2h
Function	Detects small molecule
Precision	Intra-assay Precision (Precision within an assay): 3 samples with low, middle and high level Octreotide (OT) were tested 20 times on one plate, respectively. Inter-assay Precision (Precision between assays): 3 samples with low, middle and high level Octreotide (OT) were tested on 3 different plates, 8 replicates in each plate. $CV(\%) = SD/mean \times 100$ Intra-Assay: CV<10%, Inter-Assay: CV<12%
Assay Procedure Summary	1. Prepare all reagents, samples and standards 2. Add 50ul standard or sample to each well. And then add 50ul prepared Detection Reagent A immediately. Shake and mix. Incubate 1 hour at 37 degree C 3. Aspirate and wash 3 times 4. Add 100ul prepared Detection Reagent B. Incubate 30 minutes at 37 degree C 5. Aspirate and wash 5 times 6. Add 90ul Substrate Solution. Incubate 10-20 minutes at 37 degree C 7. Add 50ul Stop Solution. Read at 450 nm immediately.
Test Principle	This assay employs the competitive inhibition enzyme immunoassay technique. A monoclonal antibody specific to octreotide has been pre-coated onto a microplate. A competitive inhibition reaction is launched between biotin labeled octreotide analogues and unlabeled antigen (Standards or samples) with the pre-coated antibody. After incubation the unbound conjugate is washed off. Next, avidin conjugated to Horseradish Peroxidase (HRP) is added to each microplate well and incubated. The amount of bound HRP conjugate is reverse proportional to the concentration of octreotide in the sample. After addition of the substrate solution, the intensity of color developed is reverse proportional to the concentration of octreotide in the sample.

Preparation and Storage

Store at 4 degree C for 1 month/-20 degree C for long-term (One year within shelf life)

The stability of kit is determined by the loss rate of activity. The loss rate of this kit is less than 5% within the expiration date under appropriate storage condition. To minimize extra influence on the performance, operation procedures and lab conditions, especially room temperature, air humidity, incubator temperature should be strictly controlled. It is also strongly suggested that the whole assay is performed by the same operator from the beginning to the end.

IMAGES



Standard Curve (Sample)

Typical Standard Curve for OT ELISA (Competitive)

For Research Use Only. Not for use in diagnostic or therapeutic procedures. Small volumes of product may become entrapped in the seal of the vial during shipment and storage; if necessary, briefly centrifuge the vial on a tabletop centrifuge to dislodge any liquid in the cap.

AAA Biotech | Tel: +1 (619) 604-9440 | Fax: +1 (619) 604-9441 | contact@aaabiotech.com

Online datasheet: <https://stage.aaabiotech.com/general-elisa-kits/octreotide-ot/161553>