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Produktinformation



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Lieferung & Zahlungsart

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Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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Datasheet

Arg⁸-Vasopressin ELISA Kit

Catalog Number: KA7281

Regulation Status: For research use only (RUO)

Product Description: Arg⁸-Vasopressin ELISA kit is a colorimetric competitive enzyme immunoassay kit with results overnight + 1 hour.

Calibration Range: 4.10 to 1,000 pg/mL

Limit of Detection: 2.84 pg/mL

Absorbance (nm): 450

Suitable Sample : serum, plasma, and tissue culture media

Detection Method : Colorimetric

Assay type: Quantitative

Regulation Status : For research use only (RUO)

Protocols: See our web site at <http://www.abnova.com/support/protocols.asp> or product page for detailed protocols

Storage Instruction: Store all components at 4°C.
Store standard at -20°C.
Avoid repeated freezing and thawing.

Entrez GeneID: 551

Gene Symbol: AVP

Gene Alias: ADH, ARVP, AVP-NP11, AVRP, VP

Gene Summary: This gene encodes a precursor protein consisting of arginine vasopressin and two associated proteins, neurophysin II and a glycopeptide, copeptin. Arginine vasopressin is a posterior pituitary hormone which is synthesized in the supraoptic nucleus and paraventricular nucleus of the hypothalamus. Along with its carrier protein, neurophysin II, it is packaged into neurosecretory vesicles and transported axonally to the nerve endings in the neurohypophysis where it is either stored or secreted into the bloodstream. The precursor is thought to be activated while it is being transported

along the axon to the posterior pituitary. Arginine vasopressin acts as a growth factor by enhancing pH regulation through acid-base transport systems. It has a direct antidiuretic action on the kidney, and also causes vasoconstriction of the peripheral vessels. This hormone can contract smooth muscle during parturition and lactation. It is also involved in cognition, tolerance, adaptation and complex sexual and maternal behaviour, as well as in the regulation of water excretion and cardiovascular functions. Mutations in this gene cause autosomal dominant neurohypophyseal diabetes insipidus (ADNDI). [provided by RefSeq]