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Datasheet for ABIN6962824 CGRP ELISA Kit

1 Image

1 Publication



Overview

| Quantity: | 96 tests |
|--------------------------|--------------------------|
| Target: | CGRP (CALCA) |
| Reactivity: | Mouse |
| Method Type: | Sandwich ELISA |
| Detection Range: | 15.63 pg/mL - 1000 pg/mL |
| Minimum Detection Limit: | 15.63 pg/mL |
| Application: | ELISA |

Product Details

| between Mouse CGRP1 and analogues was observed. Sensitivity: 9.38 pg/mL Components: • Pre-coated, ready to use 96-well strip plate, flat buttom • Plate sealer for 96 wells | | |
|--|--------------------|---|
| Analytical Method:QuantitativeDetection Method:ColorimetricSpecificity:This kit recognizes Mouse CGRP1 in samples. No Significant cross-reactivity or interference between Mouse CGRP1 and analogues was observed.Sensitivity:9.38 pg/mLComponents:• Pre-coated, ready to use 96-well strip plate, flat buttom • Plate sealer for 96 wells | Purpose: | |
| Detection Method:ColorimetricSpecificity:This kit recognizes Mouse CGRP1 in samples. No Significant cross-reactivity or interference between Mouse CGRP1 and analogues was observed.Sensitivity:9.38 pg/mLComponents:• Pre-coated, ready to use 96-well strip plate, flat buttom • Plate sealer for 96 wells | Sample Type: | Cell Culture Supernatant, Plasma, Serum |
| Specificity: This kit recognizes Mouse CGRP1 in samples. No Significant cross-reactivity or interference between Mouse CGRP1 and analogues was observed. Sensitivity: 9.38 pg/mL Components: • Pre-coated, ready to use 96-well strip plate, flat buttom • Plate sealer for 96 wells | Analytical Method: | Quantitative |
| between Mouse CGRP1 and analogues was observed. Sensitivity: 9.38 pg/mL Components: • Pre-coated, ready to use 96-well strip plate, flat buttom • Plate sealer for 96 wells | Detection Method: | Colorimetric |
| Components: • Pre-coated, ready to use 96-well strip plate, flat buttom • Plate sealer for 96 wells | Specificity: | This kit recognizes Mouse CGRP1 in samples. No Significant cross-reactivity or interference between Mouse CGRP1 and analogues was observed. |
| Plate sealer for 96 wells | Sensitivity: | 9.38 pg/mL |
| Deference Otandard | Components: | Plate sealer for 96 wellsReference Standard |

Reference Standard & Sample Diluent

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- Biotinylated Detection Antibody (100 x concentrate)
- HRP Conjugate (100 x concentrate)
- Biotinylated Detection Antibody Diluent
- HRP Conjugate Diluent
- Substrate Reagent
- Stop Solution
- Wash Buffer (25 x concentrate)
- Instruction manual

Target Details

| Target: | CGRP (CALCA) |
|-------------------|---|
| Alternative Name: | Calcitonin Gene Related Peptide 1 (CALCA Products) |
| Background: | CGRP1 |
| Pathways: | Hormone Activity, cAMP Metabolic Process, Myometrial Relaxation and Contraction, Regulation of G-Protein Coupled Receptor Protein Signaling, Skeletal Muscle Fiber Development, Feeding Behaviour |

Application Details

| Sample Volume: | 100 μL |
|----------------------|--|
| Assay Time: | 3.5 h |
| Plate: | Pre-coated |
| Protocol: | 1. Add 100 μL standard or sample to each well. Incubate for 90 min at 37 °C. |
| | 2. Remove the liquid. Add 100 μ L Biotinylated Detection Antibody. Incubate for 1 hour at 37 °C. |
| | 3. Aspirate and wash 3 times. |
| | 4. Add 100 μL HRP Conjugate. Incubate for 30 min at 37 °C. |
| | 5. Aspirate and wash 5 times. |
| | 6. Add 90 μ L Substrate Reagent. Incubate for 15 min at 37 °C. |
| | 7. Add 50 μL Stop Solution. Read at 450 nm immediately. |
| | 8. Calculation of results. |
| Reagent Preparation: | 1. Bring all reagents to room temperature (18~25 °C) before use. Follow the Microplate reader manual for set-up and preheat it for 15 min before OD measurement. |
| | 2. Wash Buffer: Dilute 30 mL of Concentrated Wash Buffer with 720 mL of deionized or distilled |
| | water to prepare 750 mL of Wash Buffer.Note: if crystals have formed in the concentrate, |
| | water to prepare 750 mL of wash burren Note. If crystals have formed in the concentrate, warm it in a 40 °C water bath and mix it gently until the crystals have completely dissolved |
| | 3. Standard working solution: Centrifuge the standard at 10,000xg for 1 min. Add 1.0 mL of |
| | 5. Standard working Solution. Centinuge the Standard at 10,000xg for 11111. Add 1.0 The Of |

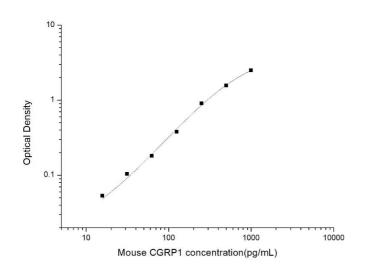
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| After it dissolves fully, mix it thoroughly with a pipette. This reconstitution produces a working solution of 1000 pg/mL. Them make serial dilutions as needed. The recommendia dilution gradient is as follows: 1000, 500, 250, 125, 625, 531 25, 1563, 0 pg/mL. Dilution method: Take 7 EP tubes, add 500 µL of Reference Standard & Sample Diluent to each tul Pipette 500 µL of the 1000 pg/mL working solution to the first tube and mix up to produce 500 pg/mL working solution. Pipette 500 µL of the solution from the former tube into the latter one according to these steps. The illustration below is for reference. Note the last is regarded as a blank. Don't pipette solution into it from the former tube. 4. Biotrinylated Detection Antibody working solution: Calculate the required amount before th experiment (100 µL/well). In preparation, sightly more than calculated should be prepare Centrifuge the stock tube before use, dilute the 100x Concentrated Biotrinylated Detection Antibody Diluent. 5. Concentrated HRP Conjugate working solution: Calculate the required amount before th experiment (100 µL/well). In preparation, sightly more than calculated should be prepare Dilute the 100x Concentrated HRP Conjugate to 1x working solution with Concentrated HRP Conjugate Diluent. 5. Sample Preparation: • It is recommended to use fresh samples without long storage, otherwise protein degrad and denaturationmay occur in these samples, leading to false results. Sample should therefore be stored for a short periodat 2-8 "Cor aliquoted at -20 "C (S1 months), Repeated freeze: thawcycles should be avoided Priot to assay, the frozen samples should be avoided Priot to assay, the frozen samples should be avoided Priot to assay, the frozen samples should be fore use or aliquoted at -20 "C (S1 months), Repeated freeze: thawcycles should be avoided Priot to assay, the frozen samples should be intherfo | | |
|--|---------------------|---|
| and denaturationmay occur in these samples, leading to false results. Samples should therefore be stored for a short periodat 2 - 8 °C or aliquoted at -20 °C (<1 month) or -80 °C (3 months). Repeated freeze-thawcycles should be avoided. Prior to assay, the frozen samples should be slowly thawed and centrifuged toremove precipitates. | | working solution of 1000 pg/mL. Then make serial dilutions as needed. The recommended dilution gradient is as follows: 1000, 500, 250, 125, 62.5, 31.25, 15.63, 0 pg/mL. Dilution method: Take 7 EP tubes, add 500 µLof Reference Standard & Sample Diluent to each tube. Pipette 500 µLof the 1000 pg/mL working solution to the first tube and mix up to produce a 500 pg/mL working solution. Pipette 500 µLof the solution from the former tube into the latter one according to these steps. The illustration below is for reference. Note: the last tube is regarded as a blank. Don't pipette solution into it from the former tube. 4. Biotinylated Detection Antibody working solution: Calculate the required amount before the experiment (100 µL/well). In preparation, slightly more than calculated should be prepared. Centrifuge the stock tube before use, dilute the 100x Concentrated Biotinylated Detection Antibody to 1xworking solution with Biotinylated Detection Antibody Diluent. 5. Concentrated HRP Conjugate working solution: Calculate the required amount before the experiment (100 µL/well). In preparation, slightly more than calculated should be prepared. Dilute the 100x Concentrated HRP Conjugate to 1x working solution with Concentrated HRP |
| experiment has to be determined.Samples should then be diluted with PBS (pH =7.0-7.2) Assay Precision: Intra-assay Precision (Precision within an assay): 3 samples with low, mid range and high le Mouse CGRP1 were tested 20 times on one plate, respectively. Inter-assay Precision (Precision between assays): 3 samples with low, mid range and high le Mouse CGRP1 were tested on 3 different plates, 20 replicates in each plate. Both intra-CV and inter-CV are < 10 %. | Sample Preparation: | therefore be stored for a short periodat 2 - 8 °C or aliquoted at -20 °C (≤1 month) or -80 °C (≤ 3 months). Repeated freeze-thawcycles should be avoided. Prior to assay, the frozen samples should be slowly thawed and centrifuged toremove precipitates. If the sample type is not specified in the instructions, a preliminary test is necessary to determinecompatibility with the kit. If a lysis buffer is used to prepare tissue homogenates or cell culture supernatant, there is a possibility of causing a deviation due to the introduced chemical substance. The |
| Mouse CGRP1 were tested 20 times on one plate, respectively. Inter-assay Precision (Precision between assays): 3 samples with low, mid range and high I Mouse CGRP1 were tested on 3 different plates, 20 replicates in each plate. Both intra-CV and inter-CV are < 10 %. | | are not in therange of the standard curve, the optimal sample dilution for the particular experiment has to be determined.Samples should then be diluted with PBS (pH =7.0-7.2). |
| Inter-assay Precision (Precision between assays): 3 samples with low, mid range and high I Mouse CGRP1 were tested on 3 different plates, 20 replicates in each plate. Both intra-CV and inter-CV are < 10 %. | Assay Precision: | Intra-assay Precision (Precision within an assay): 3 samples with low, mid range and high level |
| Mouse CGRP1 were tested on 3 different plates, 20 replicates in each plate. Both intra-CV and inter-CV are < 10 %. | | Mouse CGRP1 were tested 20 times on one plate, respectively. |
| Both intra-CV and inter-CV are < 10 %. | | Inter-assay Precision (Precision between assays): 3 samples with low, mid range and high level |
| Restrictions: For Research Use only Handling | | Mouse CGRP1 were tested on 3 different plates, 20 replicates in each plate. |
| Handling | | Both intra-CV and inter-CV are < 10 %. |
| - | Restrictions: | For Research Use only |
| | Handling | |
| Storage. 4 0,-20 0 | Storage: | 4 °C,-20 °C |

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| Storage Comment: | For unopened kit: All reagents should be stored according to the labels on the vials, so they are stable up to 6 months after receipt of the kit. The Reference Standard, Biotinylated Detection Antibody, HRP Conjugate and the 96-well stripe plate should be stored at -20 °C upon receipt while the other reagents should be stored at 4 °C. For used kit: When the kit is used, the remaining reagents need to be stored according to the above storage condition. Besides, please return the unused wells to the foil pouch containing the desiccant pack, and zip-seal the foil pouch. |
|-------------------|--|
| Expiry Date: | 6 months |
| Publications | |
| Product cited in: | Al-Hakeim, Najm, Al-Dujaili, Maes: "Major Depression in Children with Transfusion-Dependent Thalassemia Is Strongly Associated with the Combined Effects of Blood Transfusion Rate, Iron Overload, and Increased Pro-inflammatory Cytokines." in: Neurotoxicity research , Vol. 38, Issue 1, pp. 228-241, (2021) (PubMed). |
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| | Atwany, Hashemi, Jayakumar, Nagarkatti, Nagarkatti, Hassuneh: "Induction of CD4+CD25+ Regulatory T Cells from In Vitro Grown Human Mononuclear Cells by Sparteine Sulfate and Harpagoside." in: Biology , Vol. 9, Issue 8, (2020) (PubMed). |
| | Liu, Wang, Song, Sun, Hong, Pothukuchi, Xu: "Systematically transplanted human gingiva- derived mesenchymal stem cells regulate lipid metabolism and inflammation in hyperlipidemic mice with periodontitis." in: Experimental and therapeutic medicine , Vol. 19, Issue 1, pp. 672- 682, (2020) (PubMed). |
| | Zheng, Sun, Xu, Pan, Zhang, Fang, Fang, Cai: "Clinical characteristics of 34 COVID-19 patients admitted to intensive care unit in Hangzhou, China." in: Journal of Zhejiang University. Science. B , Vol. 21, Issue 5, pp. 378-387, (2020) (PubMed). |





ELISA

Image 1. Typical standard curve

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