

Datasheet for ABIN5655568 IFNA21 CLIA Kit



Overview

| Quantity: | 96 tests |
|--------------------------|--------------------------|
| Target: | IFNA21 (IFNa21) |
| Reactivity: | Human |
| Method Type: | Sandwich ELISA |
| Detection Range: | 15.62 pg/mL - 1000 pg/mL |
| Minimum Detection Limit: | 15.62 pg/mL |
| Application: | ELISA |

Product Details

| Sample Type: | Plasma, Serum |
|--------------------|---|
| Analytical Method: | Quantitative |
| Detection Method: | Chemiluminescent |
| Specificity: | This assay has high sensitivity and excellent specificity for detection of Interferon Alpha 21 (IFNa21). No significant cross-reactivity or interference between Interferon Alpha 21 (IFNa21) and analogues was observed. |
| Sensitivity: | 0.5 pg/mL |
| Target Details | |
| Target: | IFNA21 (IFNa21) |
| Alternative Name: | Interferon Alpha 21 (IFNa21 Products) |

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| Background: | Gene Name: Interferon Alpha 21 |
|---------------------|---|
| | Gene Aliases: LeIF F, Interferon alpha-F |
| Gene ID: | 3452 |
| UniProt: | P01568 |
| Pathways: | JAK-STAT Signaling, Hepatitis C |
| Application Details | |
| Comment: | 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. |
| Assay Time: | 2 - 3 h |
| Plate: | Pre-coated |
| Protocol: | The microplate provided in this kit has been pre-coated with an antibody specific to Interferon |
| | Alpha 21 (IFNa21). Standards or samples are then added to the appropriate microplate wells |
| | with a biotin-conjugated antibody specific to Interferon Alpha 21 (IFNa21). Next, Avidin |
| | conjugated to Horseradish Peroxidase (HRP) is added to each microplate well and incubated. |
| | Then the mixture of substrate A and B is added to generate glow light emission kinetics. Upon |
| | plate development, the intensity of the emitted light is proportional to the Interferon Alpha 21 |
| | (IFNa21) level in the sample or standard., |
| Assay Precision: | Intra-assay Precision (Precision within an assay): 3 samples with low, middle and high level |
| | Interferon Alpha 21 (IFNa21) were tested 20 times on one plate, respectively |
| | Inter-assay Precision (Precision between assays): 3 samples with low, middle and high level |
| | Interferon Alpha 21 (IFNa21) were tested on 3 different plates, 8 replicates in each plate. CV(%) |
| | = SD/meanX100 |
| | Intra-Assay: CV<10% |
| | Inter-Assay: CV<12% |
| Restrictions: | For Research Use only |
| Handling | |
| Handling Advice: | Do not allow to contact skin or eyes. Calibrators, controls and specimen samples should be |

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| | assayed in duplicate. Once the procedure has been started, all steps should be completed without interruption. |
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| Storage: | 4 °C,-20 °C |
| Storage Comment: | -20°C. Bring all reagents to room temperature before beginning test. The kit may be stored at 4°C for immediate use within two days upon arrival. Reseal any unused strips with desiccant pack. Minimize freeze/thaw cycles. |
| Expiry Date: | 4-8 months |