

# ELISA Kit for Fibroblast Growth Factor 2, Basic (FGF2)

Catalog No: tcue6687



## Available Sizes

**Size:** 96T



## Specifications

**Research Area:**

Cytokine;Tumor immunity;Infection immunity;

**Species Reactivity:**

Sus scrofa; Porcine (Pig)

**Sample Type:**

serum, plasma, tissue homogenates, cell lysates, cell culture supernates and other biological fluids

**Sensitivity:**

The minimum detectable dose of this kit is typically less than 4.69pg/mL

**Detection Range:**

12.35-1000pg/mL

**Assay Time:**

2h

**Detection Method:**

Enzyme-linked immunosorbent assay for Antigen Detection.

**Tested Application:**

ELISA

**Relevance:**

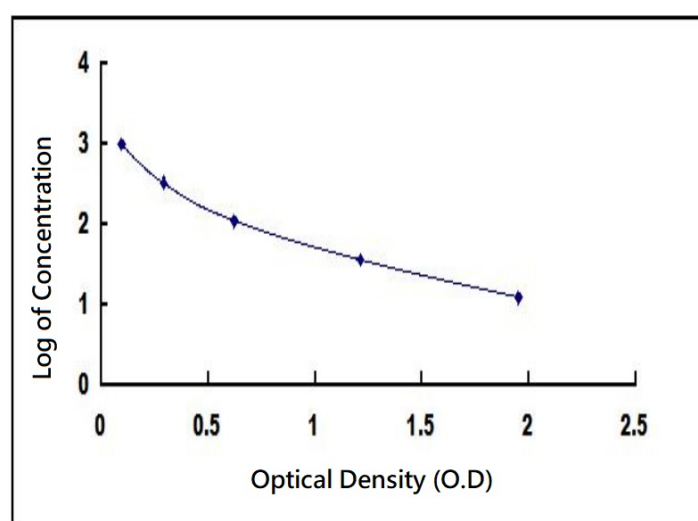
This assay has high sensitivity and excellent specificity for detection of Fibroblast Growth Factor 2, Basic (FGF2). No significant cross-reactivity or interference between Fibroblast Growth Factor 2, Basic (FGF2) and analogues was observed.

## Notes

CV(%) = SD/meanX100 Intra-Assay: CV<10% Inter-Assay: CV<12%

## Test Principle

This assay employs the competitive inhibition enzyme immunoassay technique. A monoclonal antibody specific to Fibroblast Growth Factor 2, Basic (FGF2) has been pre-coated onto a microplate. A competitive inhibition reaction is launched between biotin labeled Fibroblast Growth Factor 2, Basic (FGF2) and unlabeled Fibroblast Growth Factor 2, Basic (FGF2) (Standards or samples) with the pre-coated antibody specific to Fibroblast Growth Factor 2, Basic (FGF2). 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 Fibroblast Growth Factor 2, Basic (FGF2) in the sample. After addition of the substrate solution, the intensity of color developed is reverse proportional to the concentration of Fibroblast Growth Factor 2, Basic (FGF2) in the sample.



All products are for RESEARCH USE ONLY. Not for diagnostic & therapeutic purposes!