

# Polyclonal Antibody against Mouse FGF - 21

Catalog Number: 12180 Size: 100 µg Host: Rabbit

### Immunogen:

Recombinant full-length mouse FGF-21 expressed in E.Coli.

#### **Preparation:**

Rabbit specific IgG was purified by mouse FGF-21 affinity chromatography

# Specificity:

The antibody detects circular mouse FGF-21.

#### Formulation:

Solution in PBS

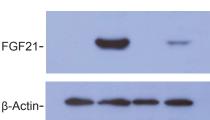
#### Storage:

Store at –20°C. For long-term storage, aliquot and freeze at -70°C. Avoid repeated freeze/defrost cycles.

## Application/Usage:

**ELISA** - This antibody can be used at 2  $\mu$ g/mL with the appropriate secondary reagents to detect mouse FGF-21. **Western blot** - This antibody can be used at  $0.2\mu$ g/mL $\sim$ 0.5 $\mu$ g/mL with the appropriate secondary reagents to detect mouse FGF-21.

Immunoprecipitation and immunocytochemistry are not tested.



Western blot analysis of FGF21 in 10ug non FGF21 expressing cell lysate (Lane 1,3) and FGF21 expressing cell lysate (Lane 2,4) using anti-FGF21 followed by goat anti-rabbit antibody. (The figure is from Prof. Cheah's lab, HKU.)

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#### Introduction

Fibroblast growth factor 21(FGF-21) is a novel protein that has been implicated in the regulation of lipid and glucose metabolism under fasting and ketotic conditions  $^{1,2}$ . In murine models, FGF-21 is predominantly expressed in liver, but it also expressed in adipose tissue and pancreatic  $\beta$  -cells  $^{3,4}$ . FGF-21 stimulates glucose uptake in adipocytes. It also protects animals from diet-induced obesity when overexpressed in transgenic mice and lowers blood glucose and triglyceride levels when administered to diabetic rodents  $^5$ . When administered daily for 6 weeks to diabetic rhesus monkeys, FGF-21 caused a dramatic decline in fasting plasma glucose, fructosamine, triglycerides, insulin, and glucagon  $^6$ . Furthermore, elevated plasma FGF-21 concentrations in humans appear to be related to the presence of hepatic and peripheral insulin resistance  $^7$ .

#### Reference:

- [1] Kharitonenkov A, Shiyanova TL, et al. (2005) J Clin Invest; 115: 1627–1635
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- [3] Nishimura T, Nakatake Y, et al. (2000) Biochim Biophys Acta; 1492: 203-206
- [4] Kurosu H, Choi M, et al. (2007) J Biol Chem; 282: 26687-26695
- [5] Kharitonenkov A, Shivanova TL, et al. (2005) *J. Clin. Invest.* 115: 1627–35.
- [6] Kharitonenkov A, Wroblewski VJ, et al. (2007) Endocrinology;148:774-81
- [7] Chavez AO, Molina-Carrion M, et al. (2009) Diabetes Care; 32:1542-6.