# Elabscience Biotechnology Co., Ltd.





# Recombinant Human FGF23(Arg179GIn) Protein(His Tag)

Catalog Number: PDMH100467

Note: Centrifuge before opening to ensure complete recovery of vial contents.

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|----|----|-------|------|----|
| JU |    | VIII. | 34.7 |    |

Species Human

Source Mammalian-derived Human FGF23 proteins Tyr25-Ile251(Arg179Gln), with an C-

terminal His

Mol\_Mass 24.8 kDa Accession Q9GZV9

**Bio-activity** Not validated for activity

#### **Properties**

**Purity** > 95% as determined by reducing SDS-PAGE.

**Endotoxin** < 1.0 EU/mg of the protein as determined by the LAL method

**Storage** Generally, lyophilized proteins are stable for up to 12 months when stored at -20 to -80

°C. Reconstituted protein solution can be stored at 4-8°C for 2-7 days. Aliquots of

reconstituted samples are stable at < -20°C for 3 months.

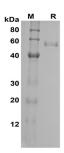
ShippingThis product is provided as lyophilized powder which is shipped with ice packs.FormulationLyophilized from a 0.2 μm filtered solution in PBS with 5% Trehalose and 5%

Mannitol.

**Reconstitution** It is recommended that sterile water be added to the vial to prepare a stock solution of

0.5 mg/mL. Concentration is measured by UV-Vis

### Data



SDS-PAGE analysis of Human FGF23 proteins, 2µg/lane of Recombinant Human FGF23 proteins was resolved with SDS-PAGE under reducing conditions, showing bands at 55

KD

## Background

This gene encodes a member of the fibroblast growth factor family of proteins, which possess broad mitogenic and cell survival activities and are involved in a variety of biological processes. The product of this gene regulates phosphate homeostasis and transport in the kidney. The full-length, functional protein may be deactivated via cleavage into N-terminal and C-terminal chains. Mutation of this cleavage site causes autosomal dominant hypophosphatemic rickets (ADHR). Mutations in this gene are also associated with hyperphosphatemic familial tumoral calcinosis (HFTC).

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