

## Datasheet for ABIN2667430 Heregulin beta 1 Protein (AA 177-241)



Overview

| Overview                 |   |
|--------------------------|---|
| Quantity:                | 50 µg   |
| Target:                  | Heregulin beta 1  |
| Protein Characteristics: | AA 177-241  |
| Origin:                  | Human   |
| Source:                  | Escherichia coli (E. coli)  |
| Protein Type:            | Recombinant   |
| Biological Activity:     | Active  |
| Application:             | Flow Cytometry (FACS)   |
| Product Details          |   |
| Purity:                  | >98 % , as determined by Coomassie stained SDS-PAGE.  |
| Endotoxin Level:         | Less than 0.1 ng per µg of protein.   |
| Target Details           |   |
| Target:                  | Heregulin beta 1  |
| Alternative Name:        | Heregulin-Beta1 (Heregulin beta 1 Products)   |
| Background:              | Heregulin, also named neuregulin-1 type I, is a family of growth factors that act as ligands for    |
|                          | the epidermal growth factor receptors, including ERBB3 and ERBB4. Heregulins regulate               |
|                          | survival, proliferation, and differentiation of the cells during development and wound healing.     |
|                          |   |
|                          | Heregulin- $\beta$ 1 is one of the Heregulin isoforms derived from alternative splicing. Binding of |

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## Target Details

|                   | activation of ERBB2, subsequently transducing signaling cascades through the mitogen-                 |
|-------------------|---|
|                   | activated protein kinase (MAPK) and phosphatidylinositol-3 kinase (PI3K) pathways. It has been        |
|                   | suggested that heregulin- $\beta$ 1 participates in the tumorgenesis and metastasis of breast cancer, |
|                   | promoting proliferation, migration, and invasion of breast cancer cells.                              |
| Molecular Weight: | The 65 amino acid recombinant protein has a predicted molecular mass of approximately 7.5             |
|                   | kDa. The predicted N-terminal acid is Ser.  |

## Application Details

| Application Notes: | Optimal working dilution should be determined by the investigator.   |
|--------------------|--|
| Comment:           | Biological activity: ED50 $\leq$ 0.5 ng/ml, corresponding to a specific activity of $\geq$ 2.0 x 106 units/mg, as measured by its ability to stimulate proliferation of human MCF7 cells in a dose dependent manner.                                     |
| Restrictions:      | For Research Use only  |
| Handling           |  |
| Format:            | Lyophilized  |
| Reconstitution:    | For maximum results, quick spin vial prior to opening. Reconstitute in water to a concentration of 0.1-1.0 mg/mL. Do not vortex. It is recommended to further dilute in a buffer, such as 5 % Trehalose, and store working aliquots at -20 °C to -80 °C. |
| Buffer:            | Lyophilized, carrier-free.   |
| Handling Advice:   | Avoid repeated freeze/thaw cycles.   |
| Storage:           | -20 °C   |
| Ctorogo Componiti  | Unapproximation by stars $d$ at 20% or 70%   |

Storage Comment: Unopened vial can be stored at -20°C or -70°C.