

Recombinant Human E-Cadherin

Catalog Number: 8505-EC

DESCRIPTION	
Source	Human embryonic kidney cell, HEK293-derived Asp155-lle707, with a C-terminal 6-His tag
	Accession # P12830
N-terminal Sequence Analysis	Asp155
Structure / Form	Noncovalently-linked homodimer
Predicted Molecular Mass	61 kDa
SPECIFICATIONS	
SDS-PAGE	74-92 kDa, reducing conditions
Activity	Measured by the ability of the immobilized protein to support the adhesion of the MCF-7 human breast cancer cells. The ED_{50} for this effect is 0.2-1 μ g/mL
Endotoxin Level	<0.10 EU per 1 µg of the protein by the LAL method.
Purity	>95%, by SDS-PAGE with silver staining.
Formulation	Lyophilized from a 0.2 µm filtered solution in PBS. See Certificate of Analysis for details.
PREPARATION AND ST	TORAGE
Reconstitution	Reconstitute at 250 μg/mL in sterile PBS.
Shipping	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below.
Stability & Storage	Use a manual defrost freezer and avoid repeated freeze-thaw cycles. 12 months from date of receipt, -20 to -70 °C as supplied. 1 month, 2 to 8 °C under sterile conditions after reconstitution. 3 months, -20 to -70 °C under sterile conditions after reconstitution.

BACKGROUND

E-Cadherin/Cadherin-1, also known as Uvomorulin in the mouse and rat, is a 120 kDa member of the Cadherin family of cell surface glycoproteins that mediate cell adhesion (1). Human E-Cadherin shares 81% amino acid sequence identity with the rat and mouse proteins (2). It is a single-pass transmembrane protein that mediates calcium-dependent epithelial cell adhesion. E-Cadherin has five extracellular EC domains that form homophilic cis-clusters between adjacent epithelial cells and trans-clusters within the same cell. E-Cadherin clusters are critical components of adherens junctions between epithelial cells and act in the formation and maintenance of the epithelial cell barrier (3, 4). The intracellular domain of E-Cadherin binds to the Catenin cytoskeletal complex, which includes p120 Catenin, beta-Catenin, alpha-Catenin, and Vinculin. E-Cadherin expression is critical for epithelial tissue homeostasis. Decreased E-Cadherin is associated with physiological and pathological epithelial-to-mesenchymal transition and cell migration, and E-Cadherin loss contributes to cancer metastasis (5). The extracellular E-Cadherin N-terminal domain can be cleaved by several proteases and is released as a soluble factor that enhances cancer cell motility and EGFR-dependent survival and proliferation (6).

References:

- 1. Gumbiner, B.M. (2005) Nat. Rev. Mol. Cell Biol. 6:622.
- 2. Bussemakers, M.J. et al. (1993) Mol. Biol. Rep. 17:123.
- 3. Guillot, C. and T. Lecuit (2013) Science 340:1185.
- 4. Tian, X. et al. (2011) J. Biomed. Biotechnol. 2011:567305.
- 5. Stemmler, M.P. (2008) Mol. Biosyst. 4:835.
- 6. David, J.M. and A.K. Rajasekaran (2012) Cancer Res. 72:2917.



