

Product datasheet for TP724110

OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

Human CD43 Protein, hFc Tag

Product data:

Product Type: Recombinant Proteins

Description: Human CD43 Protein, hFc Tag

Expression Host: HEK293

Tag: C-Human Fc

Predicted MW: The protein has a predicted molecular mass of 49.6 kDa after removal of the signal

peptide. The apparent molecular mass of CD43-hFc is approximately 55-130 kDa due to

glycosylation.

Purity: The purity of the protein is greater than 95% as determined by SDS-PAGE and Coomassie

blue staining.

Reconstitution Method: Lyophilized from sterile PBS, pH 7.4. Normally 5 % - 8% trehalose is added as protectants

before lyophilization.

Storage: Store at -20°C to -80°C for 12 months in lyophilized form. After reconstitution, if not intended

for use within a month, aliquot and store at -80°C (Avoid repeated freezing and thawing).

Lyophilized proteins are shipped at ambient temperature.

Stability: 12 months from date of despatch

Summary: This gene encodes a highly sialylated glycoprotein that functions in antigen-specific activation

of T cells, and is found on the surface of thymocytes, T lymphocytes, monocytes,

granulocytes, and some B lymphocytes. It contains a mucin-like extracellular domain, a transmembrane region and a carboxy-terminal intracellular region. The extracellular domain

has a high proportion of serine and threonine residues, allowing extensive O-glycosylation, and has one potential N-glycosylation site, while the carboxy-terminal region has potential phosphorylation sites that may mediate transduction of activation signals. Different glycoforms of this protein have been described. In stimulated immune cells, proteolytic cleavage of the extracellular domain occurs in some cell types, releasing a soluble

extracellular fragment. Defects in expression of this gene are associated with Wiskott-Aldrich

syndrome. [provided by RefSeq, Sep 2017]

