Hello Bio, Inc. 304 Wall St., Princeton, NJ 08540 USA

T. 609-683-7500 F. 609-228-4994

customercare-usa@hellobio.com



DATASHEET

Recombinant human PEDF/Serpin-F1 (HEK expressed) protein

Product overview

Name Recombinant human PEDF/Serpin-F1 (HEK expressed) protein

Cat No HB7366 Species of origin human

Alternative names Recombinant Human Pigment Epithelium-Derived Factor, HEK, Pigment epithelium-derived factor,

PEDF, Serpin-F1, SerpinF1, EPC-1, EPC1, PIG35.

Purity >95%

Description HEK expressed recombinant human PEDF/Serpin-F1 protein

Solubility & Handling

Solubility overview To make a working stock solution, add deionized water to make a solution (0.5mg/mL) and allow the

lyophilized material to dissolve. Filter the product using an appropriate sterile filter before using it in cell

culture

Solutions should be made in sterile deionized water (not less than 100 μg/ml). This solution can then be further diluted with other aqueous solutions.

Following reconstitution, solutions may be stored at 4°C and are useable for around 2-7 days

and for future use store at -18°C.

• Freeze-thaw cycles should be prevented.

Shipping Conditions

Important

Stable for ambient temperature shipping. Follow storage instructions on receipt.

This product is for RESEARCH USE ONLY and is not intended for therapeutic or diagnostic use. Not

for human or veterinary use.

Chemical Data

UniProt ID P36955 Source HEK 293.

Appearance White lyophilized powder (filtered & freeze-dried)

Formulation Lyophilized from filtered (0.4µm) solution (0.5mg/ml) containing Tris (20mM) & NaCl pH 7.5 (20mM)

References

Pigment epithelium-derived factor (PEDF) is one of the most abundant proteins secreted by human adipocytes and induces insulin resistance and inflammatory signaling in muscle and fat cells

Famulla S *et al* (2011) Int J Obes (Lond) 35(6) **PubMedID**20938440

PEDF: a multifaceted neurotrophic factor

Tombran-Tink J *et al* (2003) Nat Rev Neurosci 4(8) **PubMedID** 12894238

PEDF and its roles in physiological and pathological conditions: implication in diabetic and hypoxia-induced angiogenic diseases

He X et al (2015) Clin Sci (Lond) 128(11)