

Recombinant Human SPINK1 (C-6His)

Catalog No: C542

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| Description | Recombinant Human Serine Protease Inhibitor Kazal-Type 1 is produced by our Mammalian expression system and the target gene encoding Asp24-Cys79 is expressed with a 6His tag at the C-terminus. |
| Source | Human cells |
| Alternative name | Pancreatic Secretory Trypsin Inhibitor; Serine Protease Inhibitor Kazal-Type 1; Tumor-Associated Trypsin Inhibitor; TATI; SPINK1; PSTI |
| Accession No. | P00995 |
| Predicted Molecular Weight | 7.28kDa |
| Apparent Molecular Weight | 13kDa, reducing conditions. |
| Quality Control | Purity: greater than 95% as determined by reducing SDS-PAGE. Endotoxin: less than 0.1 ng/μg (1 EU/μg) as determined by LAL test. |
| Formulation | Supplied as a 0.2 μm filtered solution of 20mM MES, 150mM NaCl, 2mM CaCl ₂ , 1mM DTT, 0.05% Brij35, 10% Glycerol, pH 6.0. |
| Shipping | The product is shipped on dry ice pack. Upon receipt, store it immediately at the temperature listed below. |
| Storage | Store at < -20°C, stable for 6 months after receipt. Please minimize freeze-thaw cycles. |
| Background | Serine Protease Inhibitor Kazal-Type 1 (SPINK1) is a trypsin inhibitor that prevent the trypsin-catalyzed premature activation of zymogens within the pancreas. Defects in SPINK1 are a cause of pancreatitis (PCTT). A disease characterized by the presence of calculi in pancreatic ducts. It causes severe abdominal pain attacks. Defects in SPINK1 are the cause of susceptibility to tropical calcific pancreatitis (TCP). Recombinant SPINK1 protein (rSPINK1) stimulated cell proliferation in benign RWPE as well as cancerous prostate cells. The research result indicated that the potential of SPINK1 as an extracellular therapeutic target in prostate cancer. In contrast, knockdown of SPINK1 in 22RV1 cells inhibited cell proliferation, cell invasion, and tumor growth in xenograft assays. |

SDS-PAGE

