

Catalog Number: 10034-EV

DESCRIPTION	
Source	<i>Spodoptera frugiperda, Sf</i> 21 (baculovirus)-derived <i>r. sanguineus</i> Evasin-3 protein Leu21-Arg86. with a C-terminal 6-His tag Accession # P0C8E8
N-terminal Sequence Analysis	Leu21
Predicted Molecular Mass	7.8 kDa

SPECIFICATIONS	
SDS-PAGE	8-17 kDa, reducing conditions
Activity	Measured by its ability to inhibit CXCL8-induced chemotaxis of BaF3 mouse pro-B cells transfected with human CXCR2. The ED ₅₀ for this effect is 0.15-0.9 μg/mL.
Endotoxin Level	<0.10 EU per 1 μ g of the protein by the LAL method.
Purity	>95%, by SDS-PAGE visualized with Silver Staining and quantitative densitometry by Coomassie® Blue Staining.
Formulation	Lyophilized from a 0.2 µm filtered solution in PBS. See Certificate of Analysis for details.

PREPARATION AND STORAGE		
Reconstitution	Reconstitute at 250 µg/mL in PBS.	
Shipping	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below.	
Stability & Storage	 12 months from date of receipt, ≤ -20 °C as supplied. 1 month, 2 to 8 °C under sterile conditions after reconstitution. 3 months. ≤ -20 °C under sterile conditions after reconstitution. 	



BACKGROUND

Evasin-3 is a highly selective chemokine-binding protein isolated from tick saliva. The cDNA of tick Evasin-3 encodes an 86 amino acid (aa) precursor, which includes a 20 aa signal peptide and a 66 aa mature protein (1). Ticks are blood sucking parasites that secrete a wide variety of immunomodulatory proteins to evade the host immune response. The saliva isolated from Ticks has shown to contain chemokine neutralization activity. These proteins have been identified as chemokine binding proteins (CHPBs) that were termed as Evasins (1,2,3). Evasin-3 belongs to a new class of chemokine binding proteins in that it shows high affinity binding to a very limited set of chemokines, including CXCL8 and CXCL1. Since it is very small molecule Evasin-3 may be therapeutically useful as novel anti-inflammatory agent.

References:

- 1. Frauenschuh, A. et al. (2007) J. Biol. Chem. 282:27250.
- 2. Deruaz, M. et al. (2008) J. Exp. Med. 205:2019.
- 3. Deruaz, M. et al. (2013) FEBS J. 280:4876.

Rev. 11/14/2018 Page 1 of 1



Global bio-techne.com info@bio-techne.com techsupport@bio-techne.com TEL +1 612 379 2956 USA TEL 800 343 7475 **Canada** TEL 855 668 8722 **China** TEL +86 (21) 52380373 **Europe | Middle East | Africa** TEL +44 (0)1235 529449