biotechne

RDSYSTEMS

DESCRIPTION	
Source	<i>E. coli</i> -derived human IL-2 protein Ala21-Thr153 (C145S) with and without an N-terminal Met Accession # P60568.1 Proprietary point mutations
N-terminal Sequence Analysis	Ala21 & Met-Ala21
Predicted Molecular Mass	15 kDa

SPECIFICATIONS	
SDS-PAGE	12-14 kDa, under reducing conditions.
Activity	Measured in a cell proliferation assay using NK-92 human natural killer lymphoma cells. The ED ₅₀ for this effect is 0.0500-0.500 ng/mL.
Endotoxin Level	<0.20 EU per 1 µg 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 Sodium Acetate with Trehalose. See Certificate of Analysis for details.

PREPARATION AND STORAGE		
Reconstitution	Reconstitute the 10 µg size at 100 µg/mL in water. Reconstitute all the other sizes at 500 µg/mL in water.	
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

CD122 directed IL-2 has a 7-fold increased binding affinity to the human IL-2 beta receptor while exhibiting significantly reduced affinity for the IL-2 alpha receptor. According to research by Levin (2012), this configuration results in more effective expansion of effector cells and less Treg (1). Our findings show that CD122 directed IL-2 improves immune cell expansion for workflows requiring significant amounts of IL-2. One concern with improving T cell expansion is the possibility of increasing the number of terminally differentiated and exhausted T cells with reduced anti-tumor activity. However, our studies have shown that CD122 directed IL-2 significantly improves T cell expansion without changing T cell phenotype or the expression of exhaustion markers.

References:

1. Levin, A., Bates, D., Ring, A. et al. (2012) Nature. 484:529.