

2023.6.1-7.31

DISCOUNT

**50%
OFF**

GMP Grade Raw Materials

IL-7

IL-15

IL-21

OKT3 antibody

Anti CD28 antibody

Promotional Product List

分子名	製品番号	サイズ	製品名	希望小売価格	キャンペーン価格
IL-15	GMP-L15H13	50ug	GMP Human IL-15 Protein DMF Filed	¥233,000	¥116,500
IL-7	GMP-L07H24	50ug	GMP Human IL-7 Protein DMF Filed	¥308,000	¥154,000
IL-21	GMP-L21H25	50ug	GMP Human IL-21 Protein DMF Filed	¥277,000	¥138,500
CD3	GMP-MC0323	500ug	GMP Monoclonal Anti-Human CD3 Antibody (OKT3) DMF Filed	¥266,000	¥133,000
		1mg		¥392,000	¥196,000
CD28	GMP-MC2824	500ug	GMP Monoclonal Anti-Human CD28 Antibody DMF Filed	¥293,000	¥146,500

Our GMP Quality Management System

Especially in the pre-clinical and clinical phases, we understand that the safety and quality of raw and ancillary materials are of the utmost importance. Our recombinant proteins are manufactured and tested in accordance to the relevant USFDA GMP (Good Manufacturing Practice) guidelines under an ISO9001:2015 and ISO 13485:2016 quality management system with animal-origin free (AOF) materials. To ensure our products do not impact your final product, we take a step further by implementing stricter quality controls that include mycoplasma, viral, and in vivo toxicity tests.



GMP Guidelines for Raw Materials



USP<92>

Growth Factors and Cytokines Used in Cell Therapy Manufacturing



USP<1043>

Ancillary Materials for Cells, Gene, and Tissue-Engineered Products



Ph. Eur Gen. Chapter 5.2.12

Raw Materials of Biological Origin for the Production of Cell & Gene Therapy Medicinal Products



ISO/TS 20399-1-2018

Biotechnology-Ancillary Materials Present during the Production of Cellular Therapeutic Products

The successful translation of your cell therapy products hinges upon material-selection decisions that impact manufacturing. Raw materials chosen in the early stages need to conform to all regulatory criteria when entering clinical trials and later stages. At ACROBiosystems, we ensure that our GMP products follow regulatory standards from internationally recognized bodies including ISO, US and European Pharmacopeias. Furthermore, we take pride in our comprehensive and strict quality management system to produce our products. Especially in cell therapy products where safety matters, we uphold our strict expectations to ensure the ancillary materials we provide to you do not impact your final product.

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企画開発課 TEL:03-3241-2573 FAX:03-3279-6397

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