



The *In Vitro* Immunomodulatory and Anti-Tumor Activity of Thymosin α 1-Thymopentin Fusion Peptide

Juan LI¹, Deqing SUN¹, Rongmei WANG^{1*} & Fengshan WANG^{2,3*}

¹ Department of Pharmacy, The Second Hospital of Shandong University, Jinan 250033, PR China

² Key Laboratory of Chemical Biology of Natural Products (Ministry of Education), Institute of Biochemical and Biotechnological Drugs, School of Pharmaceutical Sciences, Shandong University, Jinan 250012, China

³ National Glycoengineering Research Center, Shandong University, Jinan 250012, China

SUMMARY. In this study, the *in vitro* immunomodulatory and anti-tumor activity of thymosin α 1-thymopentin (T α 1-TP5) fusion peptide was evaluated. Flow cytometry was used to detect CD69 expression in lymphocytes and thymocyte apoptosis induced by dexamethasone (Dex). Reverse transcription-PCR (RT-PCR) was used to detect the effect of T α 1-TP5 on the mRNA level of interferon γ (IFN- γ) and interleukin-2 (IL-2) in lymphocytes. 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assay was used to detect the cytotoxicity of natural killer (NK) cells and the anti-tumor activity of T α 1-TP5. The results showed that T α 1-TP5 significantly enhanced CD69 expression than thymopentin (TP5) and thymosin α 1 (T α 1). T α 1-TP5 exhibited higher activity than TP5 in inducing IFN- γ and IL-2 expression. T α 1-TP5 showed stronger activity than TP5 and T α 1 in increasing NK cells cytotoxicity and alleviating thymocytes apoptosis. Besides, T α 1-TP5 had a better tumor growth inhibitory effect than TP5. T α 1-TP5 can possibly be developed as a new immunomodulatory agent.

RESUMEN. En este estudio se evaluó la actividad inmunomoduladora y anti-tumor *in vitro* del péptido de fusión timosina α 1-timopentina (T α 1-TP5). La citometría de flujo se utilizó para detectar la expresión de CD69 en los linfocitos y la apoptosis de timocitos inducida por dexametasona (Dex). Se usó PCR con transcripción inversa (RT-PCR) para detectar el efecto de T α 1-TP5 en el nivel de mRNA del γ interferón (IFN- γ) y la interleucina-2 (IL-2) en los linfocitos. El ensayo con 3-(4,5-dimetiltiazol-2-il)-2,5-difeniltetrazolio (MTT) se utilizó para detectar la citotoxicidad de las células asesinas naturales (NK) y la actividad anti-tumor de T α 1-TP5. Los resultados mostraron que T α 1-TP5 mejora de forma significativa la expresión de CD69 de la timopentina (TP5) y α 1 timosina (T α 1). T α 1-TP5 exhibe mayor actividad que TP5 en la inducción de la expresión de IFN- γ y IL-2. T α 1-TP5 mostró mayor actividad que TP5 y T α 1 en el aumento de la citotoxicidad de células NK y la disminución de la apoptosis de timocitos. Además, T α 1-TP5 tenía un mejor efecto inhibitor del crecimiento tumoral que TP5. Posiblemente T α 1-TP5 pueda ser desarrollado como un nuevo agente inmunomodulador.

KEY WORDS: anti-tumor activity, immunomodulatory effect, thymopentin, thymosin α 1, thymosin α 1-thymopentin fusion peptide.

* Authors to whom correspondence should be addressed. E-mails: rongmeiwang@hotmail.com (Rongmei Wang), wangfengshansdu@hotmail.com (Fengshan Wang)