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In Vitro Antioxidant and Anti-inflammatory Activities of Cordyceps militaris Extract ○Yu Su SHIN¹, Chang Yeol YANG¹, Gwi Yeong JANG¹, Ah Young LEE², Ji Hyun KIM², Ji Hye YOON¹, Jehun CHOI¹, Eun Ju CHO², Takako YOKOZAWA³, Chan Hum PARK¹ (¹Rural Development Administration, Korea, ²Pusan National University, Korea, ³University of Toyama, Japan)

[Objectives] This study was conducted to *in vitro* antioxidant and anti-inflammatory activities of ethanolic extract from *Cordveens militaris* (CME).

[Materials and Methods] Antioxidant potential, total phenolic and flavonoid contents of CME were determined by Folin-Ciocalteu method and the aluminum chloride colorimetric method, respectively. Antioxidant activity of CME was measured by following some well-established methods for free radical scavenging such as 2,2-diphenyl-picrylhydrazyl hydrate and 1,2,2-azinobis-(3-ethylbenzothiazoline-6-sulfonicacid). Moreover, Anti-inflammatory activity of CME was determined by measuring the inhibition of nitric oxide (NO) production in lipopolysaccharide/interferon- γ -activated RAW 264.7 macrophage-like cells. In addition, cytotoxicity of CME against macrophages was determined by MTT assay.

[Results] Our results showed that total phenolic content was 19.7 mg gallic acid/g extract. Total flavonoid content was 5.0 mg Naringin/g. Its antioxidant activity was assessed by IC_{50} value and the values are 338.8 μ g/ml (DPPH radical scavenging), 35.4 μ g/ml (ABTS radical scavenging). In addition, CME attenuated NO production through the reduction of cellular inducible NO synthase protein expressions. Using MTT assay on indicate that CME showed no toxicity.

[Conclusion] These results provide important evidence that CME can potentially be used to antioxidant and anti-inflammatory agents.