

Protective effect of nano-capsule of naturally occurring caffeic acid and its analogs against allergic inflammation on normal human skin keratinocytes

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Introduction:

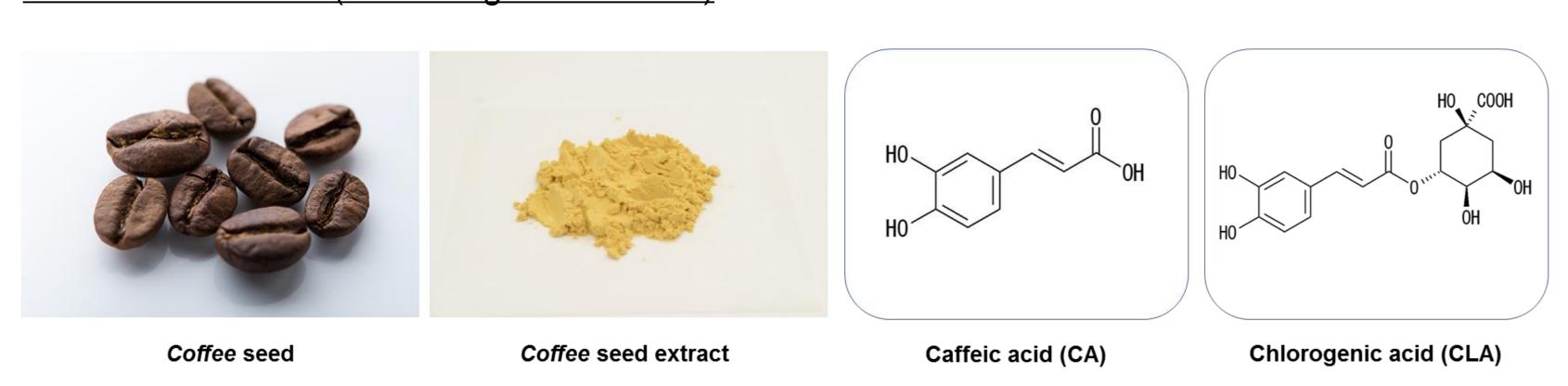
Atopic dermatitis (AD) is also known as atopic eczema, is an inflammatory chronically relapsing skin disease. ^[1] Epidermal keratinocytes release various inflammatory mediators, such as chemokines and adhesion molecules by Tumor necrosis factor –alpha (TNF-α) and Interferongamma (IFN-γ) stimulations ^{[2] [3]}. These inflammatory factors such as chemokines (IL-8, TARC, and MDC), and the adhesion molecule (ICAM-1), and MMP-9 are involved in the development of inflammatory skin diseases including AD. Common treatments for AD are steroid therapy and immuno suppressive drugs, but these show side-effects, especially with continuous application. ^[4] Therefore is needed a new natural effective agent and delivery system for anti-inflammatory cosmetics and pharmaceuticals.

Coffea Robusta seed extract (CRS) and Caffea Arabica seed extract (CAS) contain many amounts of Caffeic acid (CA) and Chlorogenic acid (CLA). We have been reported that we developed a nano-capsule (NC-CS) containing naturally CA and its analogs, and evaluated the skin permeation in epidermal human skin model and activities of anti-photoaging.

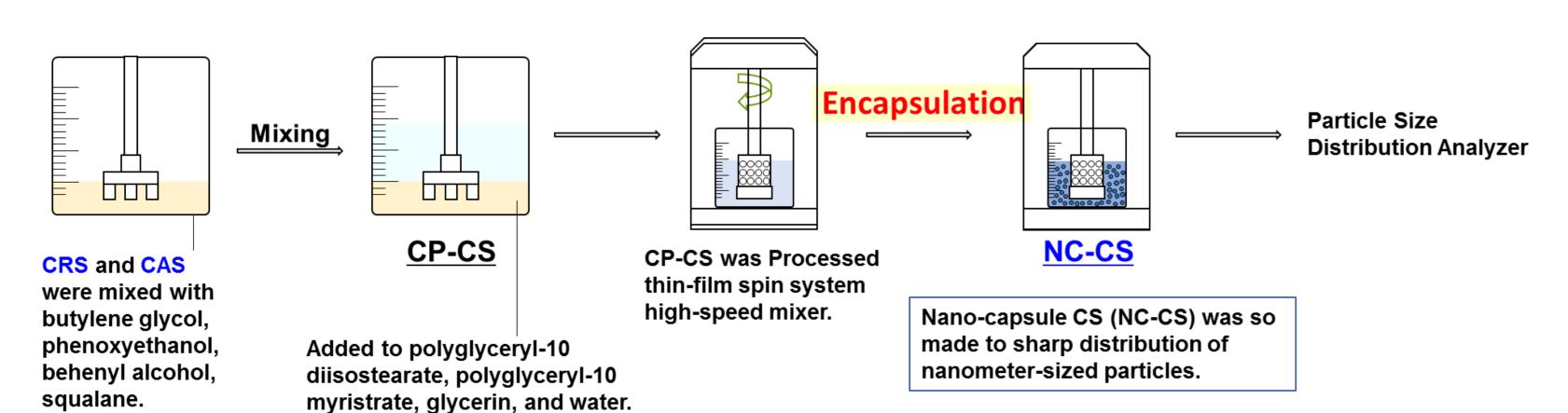
In this study, we evaluated the inhibitory effect of NC-CS of TNF- α /IFN- γ -induced anti-inflammatory in the normal human keratinocytes, as a further effect of skin care.

Materials & Methods:

Coffee seed extract (containing CA and CLA)



Preparation and morphological analysis of nano-capsules (NC-CS)

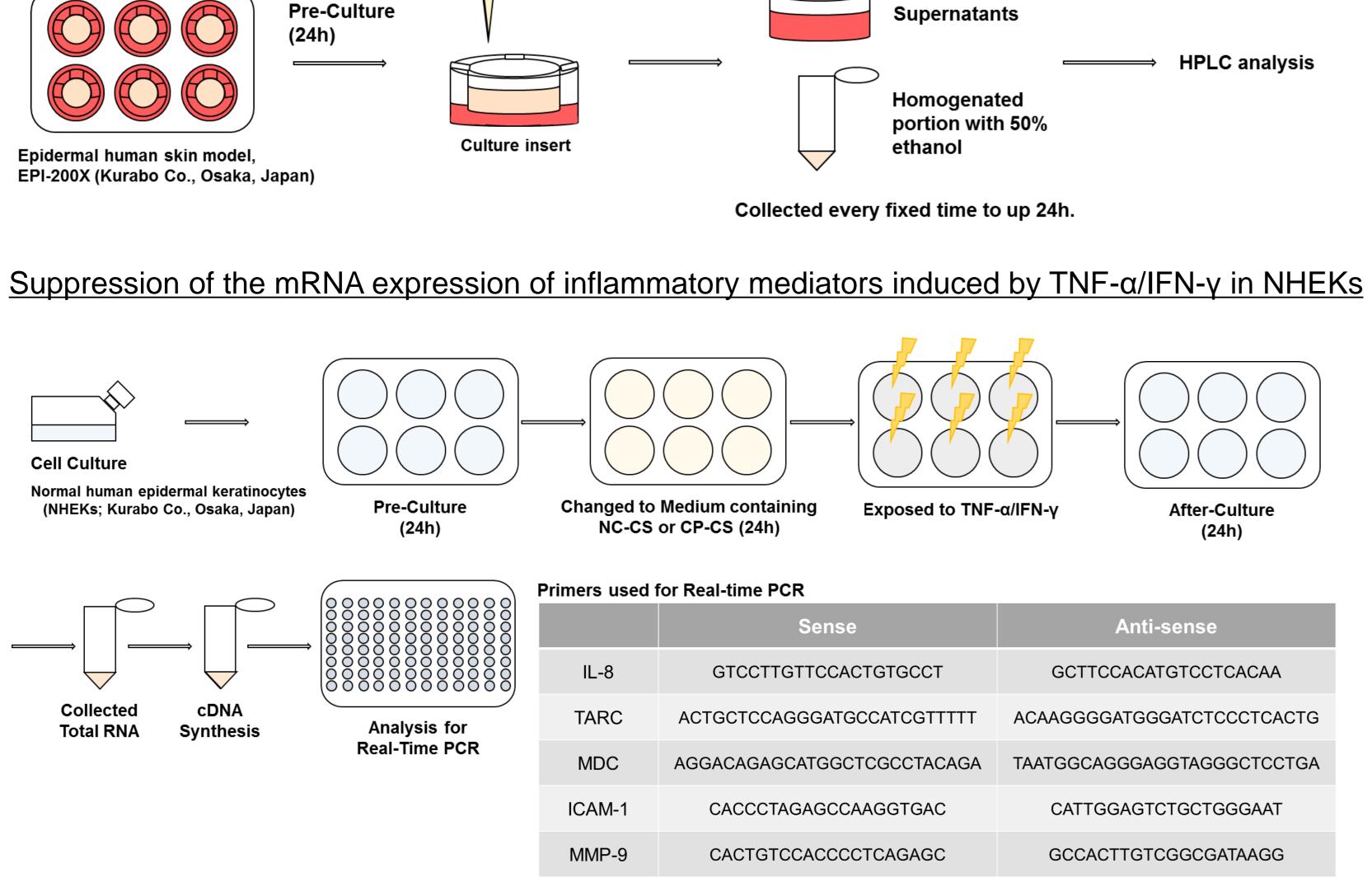


Skin permeation and accumulation assay with reconstructed human skin model

Added to

(50µL)

NC-CS or CP-CS



Results & Discussion:

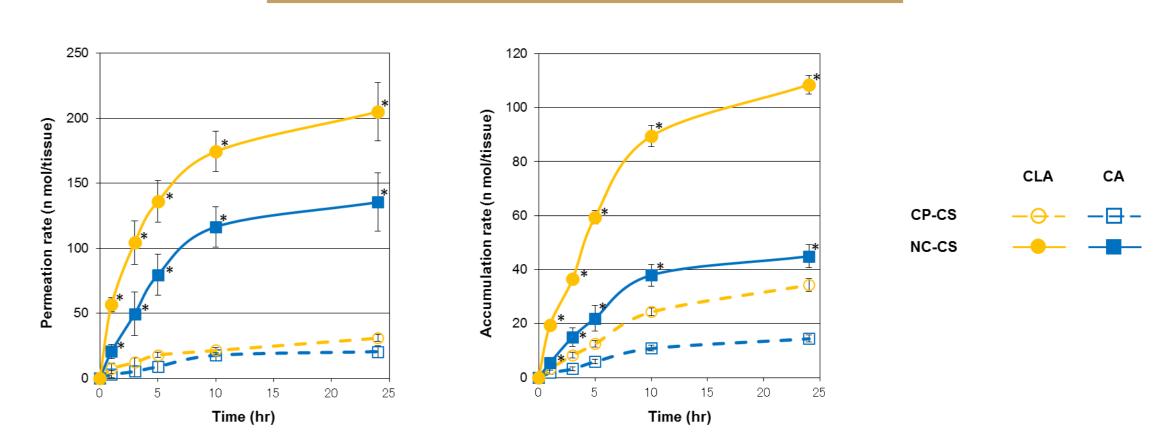


Fig.-1. Skin permeation with NC-CS on EPI-200X.

Each value represents the mean \pm SD of three experiments, and values containing asterisks were significantly different (*P<0.05) from CP-CS group.

We demonstrated a novel nano-capsule NC-CS showed higher skin permeation than non-capsule complex(Fig.-1). This results indicate that NC-CS was an effective agent for the supply of CA and CLA with transdermal activity in the skin.

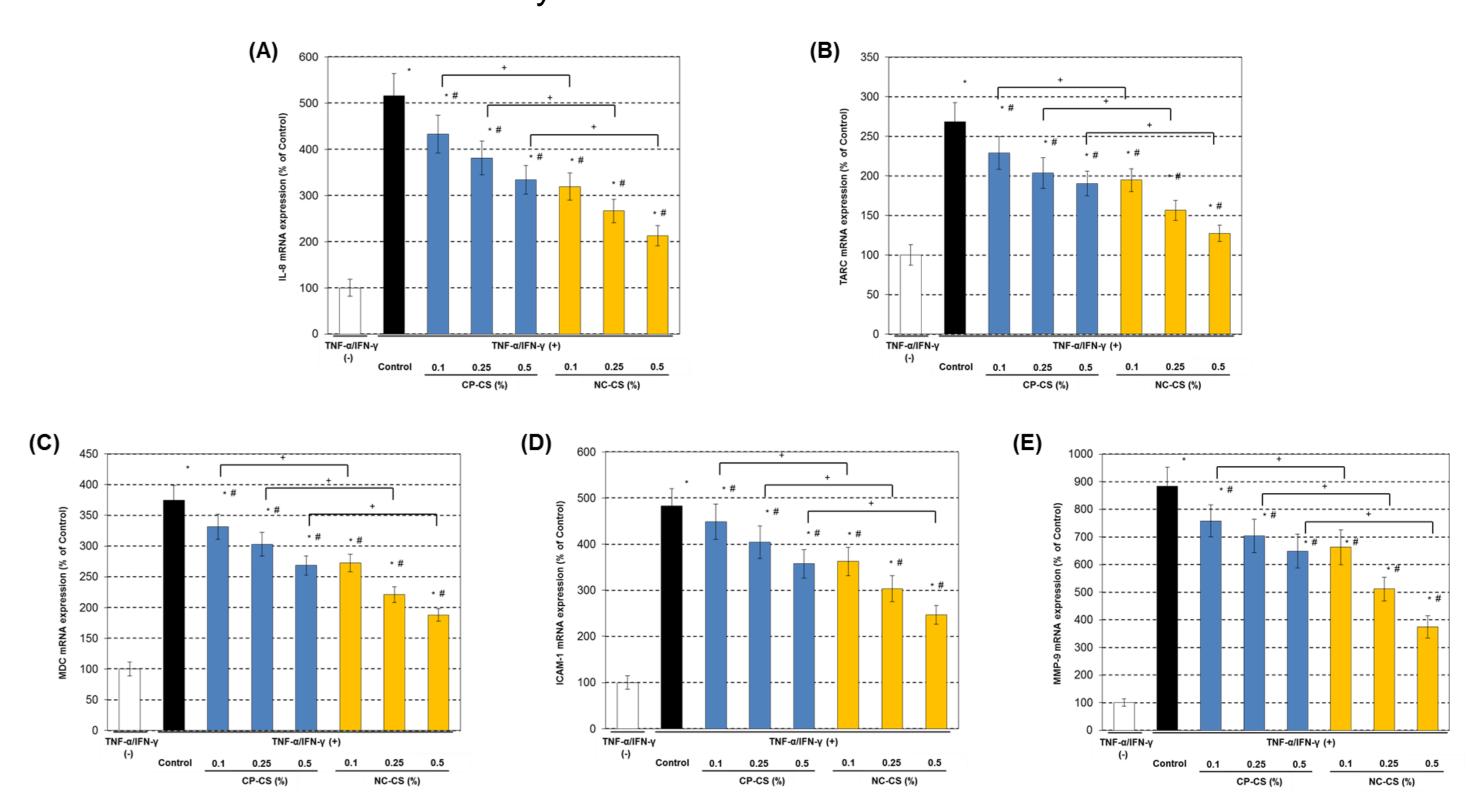


Fig.-2. Suppressive effect of NC-CS on inflammatory mediators induced by TNF-α/IFN-γ. (A)IL-8, (B)TARC, (C)MDC, (D)ICAM-1, (E)MMP-9

Delivery system of NC-CS in the skin

Each value represents the mean \pm S.E. of three experiments. Values were significantly different from the non-irradiated group, TNF- α /IFN- γ (-), at p < 0.05 (*). Values were significantly different from the irradiated group, TNF- α /IFN- γ (+), at p < 0.05 (#). Values were significantly different from the CP-CS group at p < 0.05 (+).

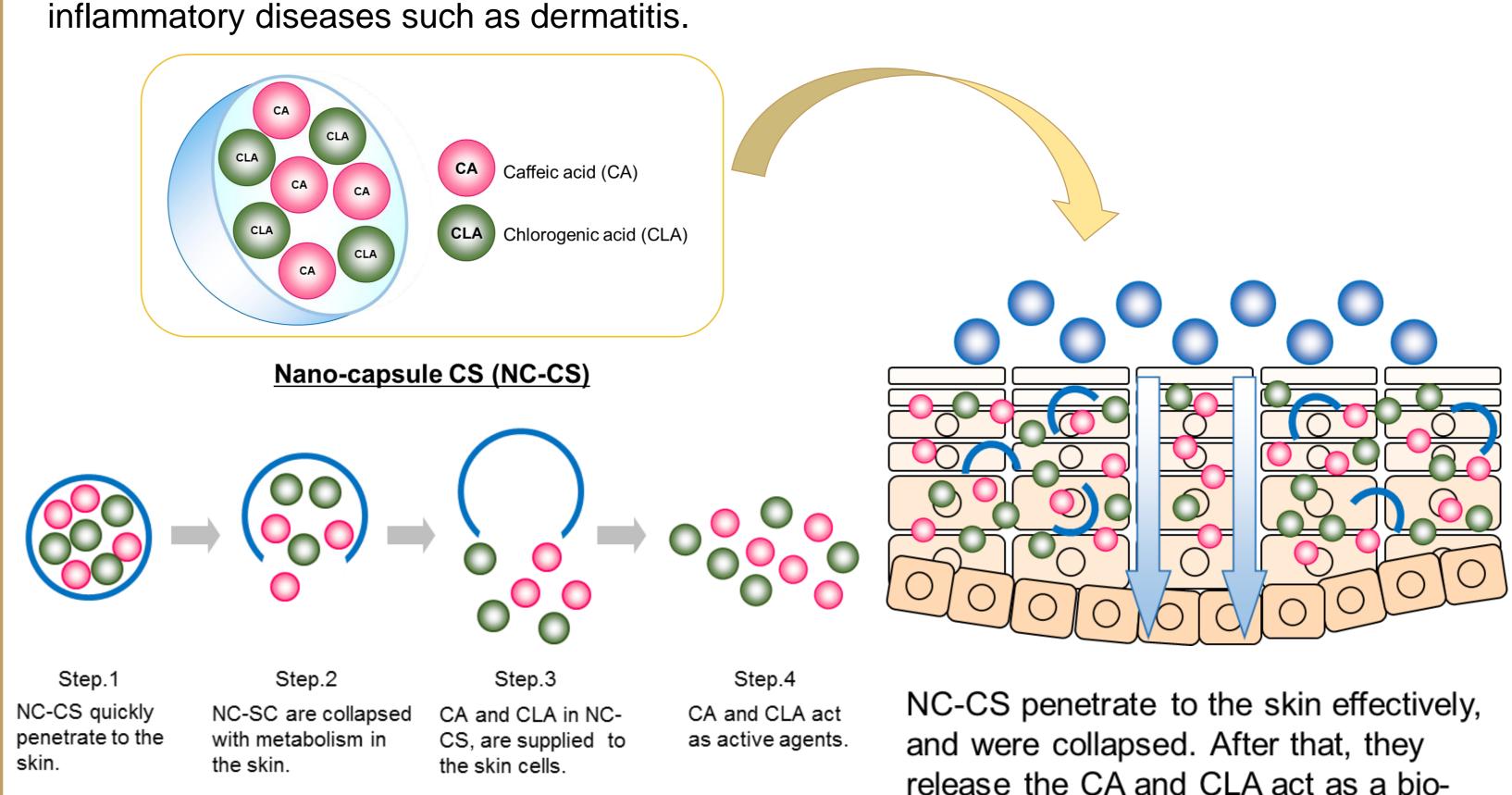
NC-CS suppressed the increases of the mRNA expression of IL-8, TARC, MDC, and ICAM-1, and MMP-9 induced with TNF- α /IFN- γ in a dose-dependent manner(Fig.-2). Furthermore, NC-CS suppressed effect stronger than CP-CS.

These results indicate that NC-CS containing CRS and CAS was an effective anti-inflammatory approaching to pro-inflammatory chemokines and adhesion molecule agent for skin.

Conclusions:

- 1. These results indicate that NC-CS was an effective agent for the supply of caffeic acid and its
- analogs with transdermal activity in the skin, and effective anti-inflammatory agent for skin care.

 2. By this study, we developed a novel naturally occurring effective nano-capsule agent, NC-CS and delivery system for anti-inflammatory cosmetics and pharmaceuticals.
- 3. In the future, we will expect NC-CS is an useful agent as a potential therapy for allergic inflammatory diseases such as dermatitis.



active agent.

References:

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