Effects of Conjugated Linoleic Acid on Anaphylaxis and Allergic Pruritus

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The effects of conjugated linoleic acid (CLA) against anaphylaxis and allergic pruritus were investigated using a in vivo assay. Inhibitory effects of CLA were observed on the immediate (type 1) hypersensitivity reaction, with CLA significantly suppressing the decrease in blood pressure (BP) and blood flow (BF) induced by the hen egg-white lysozyme (HEL)-anaphylactic reaction in ddY mice. After oral administration, CLA showed antipruritic activity, with significant inhibition of scratching behavior induced by compound 48/80 (COM), a histamine-release agent. When painted onto the skin, CLA also inhibited COM, platelet-activating factor, and protease-induced scratching behavior, and COM-induced vasodilation of the skin. CLA offers promise as a drug for the treatment of allergic and inflammatory pruritus not only as an oral but also as a topical agent. The present findings demonstrate that CLA can be effective for the prevention and treatment of allergic disease with severe pruritus.

Key words conjugated linoleic acid (CLA); antipruritus; antiallergy; antianaphylaxis; IgE

Conjugated linoleic acid (CLA) is the fatty acid group of compounds with conjugated double bonds in the intermolecule of linoleic acid. CLA was discovered by Kepler et al. in 1966.1 It has been described as an anticarcinogenic agent in grilled ground beef.2 Recently, CLA has gained explosive popularity with athletes and body builders in the U.S.A. based on reports of body fat reduction and lean body mass enhancement attributed to it.3 CLA exerts diverse physiological actions against carcinogenesis,4 arteriosclerosis,5 diabetes,6 and cardiovascular disease.7 The immune system is also affected by CLA. CLA modulates certain aspects of immune defense, including increased lymphocyte8 blastogenesis, macrophage phagocytes,9 and lymphocyte proliferation.10 On the other hand, in the superfluous immune response, CLA increases the production of IgA and IgG, promoting the allergic response through their reduction, while reducing IgE by promoting the allergic response through a superfluous increase in lymphocytes.11 CLA was also found to stimulate moderate production12 of arachidonic acid and arachidonate-derived eicosanoid, including COX-2 activity or expression.13 CLA inhibits the release of cytokines,14 histamine,15 and prostaglandin-E2 (PGE2)16 from immune cells.

These reports led us to expect that CLA would inhibit the anaphylaxis (type I hypersensitivity reaction) and allergic pruritus by affecting chemical mediators. This paper describes the inhibitory effects of CLA against anaphylaxis using a blood pressure (BP)17) or blood flow (BF)18) monitoring assay system and against allergic pruritus using a scratch- ing (itch-associated response) measuring assay system.19

MATERIALS AND METHODS

Materials CLA (75% purity) was purchased from Natural Co., Ltd. Other agents were obtained as follows: compound 48/80 (COM), Sigma; platelet activating factor (PAF), Funakoshi Co., Ltd.; bradykinin (BK), Peptide Institute Inc.; disodium cromoglycate (DSCG), Cascade Biochem Ltd. Inc.; carboxymethyl cellulose sodium salt (CMC) and hen egg-white lysozyme (HEL), Wako Pure Chemical Industries, Ltd.; Macrogol 200, Sanyo Chemical Industries, Ltd. protease (PA), Worthington Biochemical Co.; serotonin hydrochloride (5-HT), Tokyo Kasei Kogyo Co., Ltd.; Freund’s complete adjuvant, DIFCO; Evans blue, Nacalai Tesque Inc.

Animals Male ddY mice (SPF grade), 4 weeks old, and male Wistar ST rats (SPF grade), 11 weeks old, were obtained from Japan SLC (Shizuoka, Japan) and housed at 24±2 °C and 60±5% relative humidity. Food and water were available ad libitum.

Administration of CLA CLA was emulsified in 0.5% CMC for oral administration or in Macrogol 200 for topical application on mice or rats.

Assay for Antianaphylactic Activity against HEL-Induced Anaphylaxis The antianaphylactic activity was investigated as previously reported using BP17) and BF18) monitoring. Immunization with HEL was performed as previously described.17,18) Male ddY mice 5 weeks of age were sensitized i.p. on day 0 with HEL 50 μg emulsified in Freund’s complete adjuvant. To provoke anaphylaxis, on day 9 each mouse was challenged i.v. with HEL 1 μg in saline 30 μl for BP monitoring or HEL 10 μg in saline 30 μl saline for BF monitoring. The change in systolic BP of the tail artery of anaesthetized mice was measured for 15 to 20 min after challenge (the time after challenge in which BP was the lowest) using a nondirect-type BP monitor (MK-1030, Muromachi Kikai Co., Ltd.). The change in BF of the tail venous microcirculation of the unanesthetized mouse was measured every 2 min for 30 min after challenge using a laser-Doppler blood flowmeter of the noncontact type (FLO-N1, Neuroscience). The baseline (normal) BP and BF were measured for 10 min before challenge. The results are expressed as the mean±S.E. of the percent of normal BP or BF of each mouse. As a control, CLA untreated mice were challenged with HEL alone. A dose of CLA 100 mg/kg was administered orally for 2 weeks (from 5 d before sensitization to the day of challenge).

Assay of Inhibitory Effects on Production of Anti-HEL–IgE The quantity of IgE in the serum of HEL-sensitized mice was investigated as previously reported,20) using a heterogenous passive cutaneous anaphylaxis (PCA) reaction. A dose of CLA 100 mg/kg was administered orally for 2 weeks (from 5 d before sensitization to the day of challenge).

Assay of Inhibitory Effects on Intradermal COM-Induced Vasodilation The inhibitory effects of CLA on in-
tradermal COM-induced vasodilation were investigated as previously reported\(^\text{20}\) in rat skin. COM (10 μg/ml) at a dose 50 μl was intradermally injected into the shaved backs of rats. After 30 min, the extent of extravasation of dye (0.5% Evans blue) by vasodilation was compared with the untreated (control) and CLA-painted skin. CLA (75% purity) 30 μl was applied topically 1 h before injection with COM.

**Assay of Antipruritic Activity against COM-Induced Scratching Behavior** The antipruritic activity was measured using a previously reported method examine the incidence of scratching behavior.\(^\text{19}\) COM 3 mg/kg was injected subcutaneously into the base of the neck on the back of mice to provoke scratching behavior. As a control, CLA-ununtreated mice were injected with COM alone to examine the incidence of scratching behavior without PAF and PA. CLA 200 mg/kg was administered orally 1 or 24 h before injection with COM. CLA 100 mg/kg/d was repeatedly administered orally once daily for 2 weeks before the provoked scratching behavior. CLA 0.625, 1.25, or 2.5 mg/30 μl/d was repeatedly applied topically on the COM-injected site once daily for 5 d. The incidences of scratching behavior on the whole body and the site injected with COM were counted for 20 min.

**Assay of Antipruritic Activity against PAF and PA-Induced Scratching Behavior** The antipruritic activity of CLA was measured after PAF (1 μg/kg s.c.) or PA (1 mg/kg s.c.) was injected into the base of the neck on the back of mice to provoke scratching behavior. As a control, CLA-ununtreated mice were injected with PAF or PA alone to determine the incidence of scratching behavior without PAF and PA. CLA 2.5 mg/30 μl was applied topically to the PAF- and PA-injected sites of mice once daily for 5 d. The incidences of scratching behavior at the PAF- and PA-injected sites of mice were counted for 20 min.

**Statistical Analysis** Each value represents the mean±S.E. The data were evaluated by Student’s \(t\)-test (\(n=3–7\)/group).

**RESULTS**

**Antianaphylactic Effects of CLA Using BP and BF Monitoring Methods** Figure 1 shows that CLA 100 mg/kg significantly inhibited the BP decrease induced by anaphylaxis when administered orally before challenge. In addition, CLA significantly inhibited the decrease in BF in anaphylaxis when administered orally before challenge with HEL (Fig. 2). CLA alone did not change either the BP or BF of the unchallenged mice (data not shown). These results suggest that CLA has an antianaphylactic effect. However, CLA did not decrease the concentration of anti-HEL–IgE in the serum of HEL-sensitized mice (data not shown).

**Inhibitory Effects of CLA on COM-Induced Vasodilation in Rat Skin** The diameter of Evans blue-dyed spots in untreated and CLA-treated rat skin was 9.48±0.3 mm and 8.71±0.5 mm, respectively. The diameter of blue-dyed spots reflects vasodilation (increase in vascular permeability) induced by allergic mediators. CLA significantly (\(p<0.05, n=3\) rats) inhibited COM-induced vasodilation in rat skin.

**Inhibitory Effects of CLA on Scratching Behavior in Mice** Figure 3A shows that CLA 200 mg/kg significantly inhibited COM-induced scratching behavior when administered orally 24 h before the evoked scratching behavior. However, it was not effective when administered orally 1 h before the evoked scratching behavior. Figure 3B shows that repetitive oral administration of CLA (100 mg/kg/d) for 2 weeks very significantly inhibited COM-induced scratching behavior. When administered topically, CLA inhibited COM-induced scratching behavior in a dose-dependent manner (Fig. 4). A dose of 2.5 mg/kg/d of CLA was significantly inhibited COM-induced scratching behavior. CLA mitigated PAF-induced (Fig. 5A) and PA-induced (Fig. 5B) scratching behavior. CLA alone did not affect scratching behavior in normal mice on all schedules or with all methods administration investigated.

**DISCUSSION**

We found that CLA has inhibitory effects against anaphyl-
CLA exhibited antipruritic activity (Fig. 4) and inhibited skin vasodilation induced by allergic mediators. These results suggest that CLA would be useful in treating allergic pruritus not only as an oral but also as a topical agent.

Fig. 4. Inhibitory Effect of CLA on COM-Induced Scratching Behavior
The bar labeled “control” refers to untreated mice injected with COM 3 mg/kg s.c. “Normal” shows the incidence of scratching behavior before injection with COM. CLA 0.625, 1.25, and 2.5 mg/d was repeatedly applied to the COM-injected site once daily for 5 d before injection with COM. The scratching behavior at the COM-injected site was counted for 20 min. Each value represents the mean ± S.E. of 5 mice. *p<0.05, **p<0.01 compared with control (Student’s t-test).

Fig. 5. Inhibitory Effect of CLA on PAF and Protease (PA)-Induced Scratching Behavior
PAF (A) 1 μg/kg s.c. or PA (B) 1 mg/kg s.c. was injected to provoke scratching. CLA 2.5 mg/d was repeatedly applied once daily for 5 d before injection with PAF or PA. The scratching behavior at the PAF- or PA-injected site was counted for 20 min. Each value represents the mean ± S.E. of 3 or 5 mice. *p<0.05 compared with control (Student’s t-test).

REFERENCES

In summary, the present findings demonstrate that CLA can be effective for the prevention and treatment of an anaphylaxis and an allergic disease with severe pruritus.