Comparison of Some Biochemical Properties of Epidermis in Tumor Promotion-Susceptible and -Resistant Strains of Mice

Satoshi YAMAMOTO, Itsumi KIYOTO, Eriko AIZU, Nobuyuki SASAKAWA, Toshio NAKAKI and Ryuichi KATO
Department of Pharmacology, School of Medicine, Keio University, 35 Shinanomachi, Shinjuku-ku, Tokyo 160, Japan

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Abstract—It has been reported that CD-1 and SENCAR mice are susceptible and C57BL/6 mice are resistant to skin tumor promotion caused by phorbol esters. Specific binding of a phorbol ester to its epidermal receptor site, epidermal protein kinase C activity, and ornithine decarboxylase (ODC) induction in epidermis were compared between tumor promotion-susceptible and -resistant strains of mice. Specific binding of \[^{3}H\]12-O-tetradecanoylphorbol-13-acetate (TPA) to the particulate fraction of the epidermis of C57BL/6 mice gave a similar dissociation constant \((K_d)\) and a maximal number of binding sites \((B_{max})\) to those of CD-1 mice. Protein kinase C activity of the epidermal 105,000 g supernatant was not significantly different between C57BL/6 and CD-1 mice. Protein kinase C activity of the 105,000 g pellet, however, was significantly higher in C57BL/6 mice than in CD-1 mice. A topical application of TPA to the skin caused epidermal ODC induction in all of these strains of mice. At any doses of TPA, TPA-induced epidermal ODC activity of C57BL/6 mice was always higher than those of SENCAR and CD-1 mice. Maximal induction of epidermal ODC by TPA was also highest in C57BL/6 mice among these three strains of mice. These results indicate that the mechanism of the difference in susceptibility of C57BL/6, CD-1 and SENCAR mice to the tumor-promoting action of TPA resides in a step distal to or other than the protein kinase C activation and ODC induction.

Susceptibility to two-stage skin carcinogenesis by initiation and promotion is different according to the strains of mice (1). Some strains of mice are susceptible to the tumor-promoting action of phorbol esters such as 12-O-tetradecanoylphorbol-13-acetate (TPA), but some strains are far less susceptible (1). In this regard, C57BL/6 mice appear to be refractory to the tumor-promoting action of TPA (1–3). On the other hand, CD-1 and SENCAR mice are susceptible to TPA-caused tumor promotion (1–3). Little is known of the cellular basis for the difference in susceptibility. TPA-type promoters have been shown to alter many biochemical events in mouse skin including the induction of ornithine decarboxylase (ODC) activity (4, 5), which is believed to be one of the essential factors for tumor promotion. Binding of TPA to its cellular receptor results in the activation of protein kinase C, and protein kinase C-induced protein phosphorylation may mediate the tumor-promoting action of TPA (6, 7). In attempting to define the basis for the difference in susceptibility, we compared the receptor binding of \[^{3}H\]TPA, protein kinase C activity and ornithine decarboxylase induction in tumor promotion-susceptible and -resistant strains of mice.

Materials and Methods

In vivo experiments: Female C57BL/6, CD-1 and SENCAR mice, 7–8 weeks of age, were used. The dorsal hair of each mouse was shaved as described previously (8). Mice were topically treated with 200 \(\mu\)l of acetone containing the indicated amounts of TPA. Five hours after TPA application, the mice
were killed by cervical dislocation and the epidermis was separated by a brief heat treatment (9). The ODC activity of the soluble epidermal supernatant was determined by measuring the release of $^{14}\text{CO}_2$ from $[1-^{14}\text{C}]-\text{ornithine}$ as described previously (10).

$[^3\text{H}]\text{TPA}$ binding studies: Mouse epidermis was homogenized in 10 volumes (w/v) of 50 mM Tris-Cl buffer (pH 7.5) with a Polytron homogenizer. The homogenate was centrifuged at 10,000 $\times$ g for 30 min at 4°C, and the supernatant was centrifuged again at 105,000 $\times$ g for 60 min. The resultant pellet was resuspended in 50 mM Tris-Cl buffer (pH 7.5) and used for the binding assay. The samples (60 $\mu$g/ml protein; final concentration) were incubated with various concentrations (0.125–3 nM) of $[^3\text{H}]\text{TPA}$ in 50 mM Tris-Cl buffer (pH 7.5) supplemented with 4 mg/ml bovine serum albumin at 39°C for 20 min. For separating bound from free $[^3\text{H}]\text{TPA}$, a membrane filtration technique was performed with use of cold acetone just as described previously (11). Nonspecific binding was defined as the counts observed in the presence of 0.5 $\mu$M TPA. Specific binding was defined as the total binding minus the nonspecific binding.

Assay of protein kinase C activity: Epidermis was homogenized in 3 volumes (w/v) of 20 mM Tris-Cl buffer (pH 7.5) containing 2 mM EDTA, 2 mM EGTA, 0.25 M sucrose and 2 mM phenylmethylsulfonyl fluoride (PMSF) with a Polytron homogenizer, and the homogenate was centrifuged at 105,000 $\times$ g for 60 min. The resultant cytosol fraction was loaded onto a 0.8x3 cm column of DEAE cellulose (DE52, Whatman, Ltd., Maidstone, U.K.) pre-equilibrated with 20 mM Tris-Cl buffer (pH 7.5) containing 2 mM EDTA, 0.5 mM EGTA, 2 mM PMSF and 50 mM 2-mercaptoethanol. Columns were washed with 6 ml of the above buffer. Protein kinase C was eluted with 2 ml of the buffer containing 0.1 M NaCl. The 105,000 $\times$ g pellet was resuspended in the original volume of homogenizing buffer containing 0.3% (v/v) Triton X-100 and gently stirred at 4°C for 60 min. The suspension was centrifuged at 105,000 $\times$ g for 60 min at 4°C. The resulting supernatant was subjected to DEAE cellulose (DE52) column chromatography just as described above. In this case, however, the equilibration buffer contained 0.02% Triton X-100. Protein kinase C was eluted with the buffer containing 0.1 M NaCl. Protein kinase activity was determined by incubating 2.5 nmol $[^{32}\text{P}]\text{ATP}$, 30 $\mu$g histone, 500 nmol magnesium acetate, 6.25 nmol 2-mercaptoethanol and 50 $\mu$l of crude enzyme preparation in the presence or absence of 3.1 $\mu$g phosphatidylycerine, 300 nmol CaCl$_2$ and 3.75 pmol TPA in 20 mM Tris-Cl (pH 7.5) buffer in a total volume of 125 $\mu$l. The reaction was started by the addition of $[^{32}\text{P}]\text{ATP}$. Incubation was carried out at 37°C for 3 min; then the reaction was stopped by transferring a 50 $\mu$l aliquot of the incubation mixture onto 6.25-cm$^2$ pieces of Whatman cellulose phosphate paper (P81). The pieces of paper were washed 4 times with deionized water, twice with acetone, and once with petroleum-ether. The radioactivity on each piece of paper was determined by scintillation counting.

Materials: CD-1 mice were obtained from Charles River, Atsugi, Japan; C57BL/6 mice were from Clea, Tokyo, Japan; SENCAR mice were from Shizuoka Agricultural Cooperative Association for Laboratory Animals, Hamamatsu, Japan. TPA was purchased from Chemicals for Cancer Research, Inc., Chicago, IL; L-$\alpha$-phosphatidyl-L-serine, ATP, histone (type III-S), PMSF were from Sigma Chemical Co., St. Louis, MO; $[^3\text{H}]\text{TPA}$ (12.7 Ci/mmol) and $[^{32}\text{P}]\text{ATP}$ were from New England Nuclear, Boston, MA; DL-$[1-^{14}\text{C}]-\text{ornithine}$ (61 mCi/mmol) was from Amersham, Buckinghamshire, U.K.

Results

Epidermal protein kinase C activities of C57BL/6 and CD-1 mice were also compared. Since protein kinase C can be degraded by proteases to generate a Ca$^{2+}$ and phospholipid-independent kinase designated as M kinase (12), the enzyme preparation was prepared in the presence of the protease inhibitor PMSF. Protein kinase C activities of the 105,000 $\times$ g supernatant observed in the presence of Ca$^{2+}$ and phosphatidylinerine either with or without TPA were not significantly different between C57BL/6 and CD-1 mice (Table 1). On the other hand, protein kinase C activity of the 105,000 $\times$ g pellet of
C57BL/6 mice was significantly higher than that of CD-1 mice.

Specific binding of [3H]TPA to its receptor site in the epidermis was compared between C57BL/6 and CD-1 mice. Since it is known that phorbol ester binding remains intact after protease treatment (13-15), the epidermal particulate fraction was prepared in the absence of protease inhibitor. Linear increase in the specific binding was observed with the particulate fraction obtained either from C57BL/6 or CD-1 mice in the concentration range of 0 to 100 μg/ml protein (data not shown). Thus, the following experiments were performed with 60 μg/ml protein. In either strain of mice, the binding of [3H]TPA reached apparent equilibrium within 20 min at 39°C, and specific binding was apparently saturable (data not shown). As shown in Table 2, the dissociation constant (Kd) and the maximal number of binding sites (Bmax) were similar in both strains of mice.

We next determined whether epidermal responses of ODC induction to a topical application of TPA are different in C57BL/6, CD-1 and SENCAR mice (Fig. 1). Irrespective of the mouse strain, ODC was maximally induced at 5 to 7 hr after TPA application (data not shown). Therefore, epidermal ODC activity was determined at 5 hr following TPA application. At any doses of TPA, TPA-induced epidermal ODC activity of C57BL/6 mice was always higher than those of SENCAR and CD-1 mice. Maximal induction of epidermal ODC by TPA was also highest in C57BL/6 mice among these three strains of mice. The results were highly reproducible.

**Discussion**

The present results clearly show that no difference exists in the specific binding of [3H]TPA to its membrane receptor sites on the epidermis between the tumor promotion-susceptible CD-1 mice and the tumor pro-
motion-resistant C57BL/6 mice. Wheldon et al. also reported that the binding characteristics of [3H]phorbol-12,13-dibutyrate to the particulate fraction of mouse skin from tumor promotion-sensitive Swiss mice are indistinguishable from those of tumor promotion-resistant Balb/c mice (16). Protein kinase C activities detected in the epidermal 105,000 x g supernatant were also similar in both strains of mice. Garte et al. (17) reported that the cytosolic protein kinase C activity of C57BL/6 mice was not significantly different from that of SENCAR mice, consistent with our present results with C57BL/6 mice and CD-1 mice. Garte et al. (17), however, did not investigate the protein kinase C activities of the particulate fractions. As shown in our present study, protein kinase C activity of the epidermal 105,000 x g pellet was significantly higher in C57BL/6 mice than in CD-1 mice. Since it is plausible that the membrane receptor site for TPA is membrane-bound protein kinase C itself, the data of [3H]TPA binding and protein kinase C activity in the particulate fraction would correspond to each other. The differences in the procedures taken to make the particulate fraction for the binding experiments and the enzyme preparation or the assay conditions may cause the apparent dissociation in the experimental results. In any case, resistance of C57BL/6 mice to TPA-induced tumor promotion can be explained neither by the difference in [3H]TPA binding nor protein kinase C activity. At present, we cannot exclude the possibility that the level of the endogenous substrate protein is different between C57BL/6 mice and CD-1 mice. It has been well established that the protein kinase C activation by TPA is prerequisite for ODC induction (6, 7, 18), and the ODC induction is essential for the TPA-induced skin tumor promotion (4, 5). Response of epidermal ODC induction by a topical application of TPA was highest in C57BL/6 mice among these three strains of mice, indicating that the inducibility of ODC by TPA does not correlate with the susceptibility of mice to TPA-induced tumor promotion. These results suggest that the mechanism of the difference in susceptibility of C57BL/6, CD-1 and SENCAR mice to the tumor-promoting action of TPA resides in a step distal to or other than the protein kinase C activation and ODC induction.

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Fig. 1. Dose-response relationships of epidermal ODC induction by a topical application of TPA in C57BL/6, CD-1 and SENCAR mice. Mice were topically treated with the indicated amounts of TPA and killed for the determination of ODC activity 5 hr after the TPA application. Each value is the mean±S.E. of individual determinations from 5 mice. ▲, C57BL/6 mice; ●, CD-1 mice; ○, SENCAR mice.

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