

Quercetin Inhibits $\alpha 3\beta 4$ Nicotinic Acetylcholine Receptor-Mediated Ion Currents Expressed in *Xenopus* Oocytes

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Quercetin mainly exists in the skin of colored fruits and vegetables as one of flavonoids. Recent studies show that quercetin, like other flavonoids, has diverse pharmacological actions. However, relatively little is known about quercetin effects in the regulations of ligand-gated ion channels. In the previous reports, we have shown that quercetin regulates subsets of homomeric ligand-gated ion channels such as glycine, 5-HT_{3A} and $\alpha 7$ nicotinic acetylcholine receptors. In the present study, we examined quercetin effects on heteromeric neuronal $\alpha 3\beta 4$ nicotinic acetylcholine receptor channel activity expressed in *Xenopus* oocytes after injection of cRNA encoding bovine neuronal $\alpha 3$ and $\beta 4$ subunits. Treatment with acetylcholine elicited an inward peak current (I_{ACh}) in oocytes expressing $\alpha 3\beta 4$ nicotinic acetylcholine receptor. Co-treatment with quercetin and acetylcholine inhibited I_{ACh} in oocytes expressing $\alpha 3\beta 4$ nicotinic acetylcholine receptors. The inhibition of I_{ACh} by quercetin was reversible and concentration-dependent. The half-inhibitory concentration (IC₅₀) of quercetin was $14.9 \pm 0.8 \mu\text{M}$ in oocytes expressing $\alpha 3\beta 4$ nicotinic acetylcholine receptor. The inhibition of I_{ACh} by quercetin was voltage-independent and non-competitive. These results indicate that quercetin might regulate $\alpha 3\beta 4$ nicotinic acetylcholine receptor and this regulation might be one of the pharmacological actions of quercetin in nervous systems.

Key Words: Flavonoids, Quercetin, $\alpha 3\beta 4$ nicotinic acetylcholine receptor, *Xenopus* oocyte

INTRODUCTION

Nicotinic acetylcholine receptors are members of the Cys-loop family of ligand-gated ion channels [1]. Currently, sixteen different nicotinic acetylcholine receptor subunits are known [2]. Muscle form of nicotinic acetylcholine receptor consists of $\alpha 1\beta 1\gamma \epsilon$. Activation of muscle form of nicotinic acetylcholine receptor initiates muscle contraction by inducing depolarization of neuromuscular junctions. Neuronal forms of nicotinic acetylcholine receptor consist of α ($\alpha 2-7$, $\alpha 9$, $\alpha 10$) and β ($\beta 2-4$) and their activations are mainly involved in rapid synaptic transmissions in central and peripheral nervous systems [2]. Neuronal nicotinic acetylcholine receptor contain $\alpha 2-6$ subunits that are usually expressed as heteropentamers in combination with $\beta 2-4$ subunits [3-6]. For example, the $\alpha 3$ and $\beta 4$ subunits can form heteromeric receptors [7]. Although many nicotinic acetylcholine receptor subunits are expressed in the central and peripheral nervous systems, the distributions

of $\alpha 3\beta 4$ nicotinic acetylcholine receptor are mainly restricted to several tissues such as adrenal chromaffin cells [8,9]. The $\alpha 3\beta 4$ nicotinic acetylcholine receptors play an important role for release of catecholamine releases [10].

Skin of colored fruits and vegetables contains a variety of flavonoids and quercetin is one of representative flavonoids (Fig. 1A) [11]. Quercetin exhibits multiple pharmacological activities in nervous and non-nervous systems [12-16]. However, the underlying cellular mechanisms of quercetin actions are relatively unknown, especially with regards to possible regulation of receptors or ionic channels involved in synaptic transmissions. Recently, we demonstrated that quercetin differentially regulates subsets of nicotinic acetylcholine receptor such as homomeric glycine, 5-HT_{3A}, $\alpha 7$ and $\alpha 9\alpha 10$ nicotinic acetylcholine receptors [17-20]. In the present study, we examined the effects of quercetin on heteromeric $\alpha 3\beta 4$ nicotine acetylcholine receptor channel activity and report here that quercetin inhibits heteromeric $\alpha 3\beta 4$ nicotine acetylcholine receptor channel activity with voltage-independent and non-competitive manner. These results indicate that quercetin might play a role for the regulation of nicotinic acetylcholine receptor channel activities.

Received December 27, 2010, Revised January 17, 2011,
Accepted January 18, 2011

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ABBREVIATIONS: 5-HT_{3A}, 5-hydroxytryptamine 3A; ACh, acetylcholine; Que, quercetin.

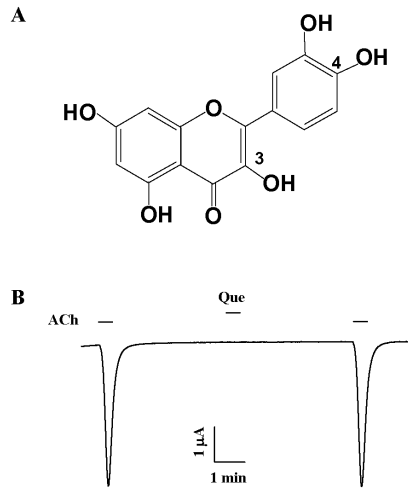


Fig. 1. Chemical structure of quercetin (A) and effect of quercetin (Que) in oocytes expressing $\alpha 3\beta 4$ nicotinic acetylcholine receptors. Quercetin itself had no effect on I_{ACh} in oocytes expressing $\alpha 3\beta 4$ nicotinic acetylcholine receptors (B).

METHODS

Materials

Bovine $\alpha 3$ and $\beta 4$ nicotine acetylcholine receptor subunit cDNAs were kindly provided by Dr. S. Sala (Universidad Miguel Hernández-Consejo Superior de Investigaciones Científicas, Spain). Quercetin (Fig. 1A) and all other reagents were purchased from Sigma-Aldrich (St. Louis, MO, USA).

Preparation of *Xenopus laevis* oocytes and microinjection

X. laevis frogs were purchased from Xenopus I (Ann Arbor, MI, USA). Animal care and handling were in accordance with the highest standards of Konkuk university guidelines. To isolate oocytes, frogs were anesthetized with an aerated solution of 3-amino benzoic acid ethyl ester, and the ovarian follicles were removed. The oocytes were separated with collagenase followed by agitation for 2 h in a Ca^{2+} -free medium containing 82.5 mM NaCl, 2 mM KCl, 1 mM $MgCl_2$, 5 mM HEPES, 2.5 mM sodium pyruvate, 100 units/ml penicillin, and 100 μ g/ml streptomycin. Stage V-VI oocytes were collected and stored in a ND96 medium (96 mM NaCl, 2 mM KCl, 1 mM $MgCl_2$, 1.8 mM $CaCl_2$, and 5 mM HEPES, pH 7.5) supplemented with 50 μ g/ml gentamicin. The solution containing the oocytes was maintained at 18°C with continuous gentle shaking and was replaced daily. Electrophysiological experiments were performed three to five days after oocyte isolation. For $\alpha 3\beta 4$ nicotine acetylcholine receptor experiments, $\alpha 3$ and $\beta 4$ nicotine acetylcholine receptor subunit-encoding cRNAs (40 nl) were co-injected into the animal or vegetal pole of each oocyte 1 day after isolation using a 10- μ l microdispenser (VWR Scientific, West Chester, PA, USA) fitted with a tapered glass pipette tip (15–20 μ m in diameter) [19].

Data recording

A custom-made Plexiglas net chamber was used for two-electrode voltage-clamp recordings, as previously reported [19]. A single oocyte was constantly superfused with a recording solution (96 mM NaCl, 2 mM KCl, 1.8 mM $CaCl_2$, and 10 mM HEPES, pH 7.5) in the absence or presence of glutamate or quercetin during recording. The microelectrodes were filled with 3 M KCl and had a resistance of 0.2–0.7 M Ω . Two-electrode voltage-clamp recordings were obtained at room temperature using an Oocyte Clamp (OC-725C, Warner Instrument) and were digitized using Digidata 1,200 A (Molecular Devices, Sunnyvale, CA, USA). Stimulation and data acquisition were controlled using pClamp 8 software (Molecular Devices). For most electrophysiological data, the oocytes were clamped at a holding potential of -80 mV. For current and voltage (I-V) relationship, voltage ramps were applied from -100 to $+60$ mV for 300-ms. In the different membrane-holding potential experiments, the oocytes were clamped at the indicated holding potentials. Linear leak and capacitance currents were corrected by means of the leak subtraction procedure.

Data analysis

To obtain the concentration-response curve for the effect of quercetin on the inward peak I_{ACh} mediated by the $\alpha 3\beta 4$ nicotinic acetylcholine receptor, the I_{ACh} peak was plotted at different concentrations of quercetin. Origin software (OriginLab Corp., Northampton, MA, USA) was used to fit the plot to the Hill equation: $I/I_{max} = 1/[1 + (IC_{50}/[A])^{nH}]$, where I_{max} was maximal current obtained from each ED₅₀ value of acetylcholine in wild-type receptors, IC_{50} was the concentration of quercetin required to decrease the response by 50%, $[A]$ was the concentration of quercetin, and nH was the Hill coefficient. All values were presented as mean \pm S.E.M. The differences between the means of control and treatment data were determined using the unpaired Student's t-test. A value of $p < 0.05$ was considered to be statistically significant.

RESULTS

Effect of quercetin on I_{ACh} in oocytes expressing heteromeric $\alpha 3\beta 4$ nicotinic acetylcholine receptors

The addition of acetylcholine to the bathing solution induced a large inward current in oocytes injected with $\alpha 3\beta 4$ nicotinic acetylcholine receptor, indicating that this nicotinic acetylcholine receptor was functionally expressed in this system (Fig. 1B). Quercetin itself had no effect in oocytes expressing $\alpha 3\beta 4$ nicotinic acetylcholine receptors at a holding potential of -80 mV (Fig. 1B). But co- and pre-treatment with quercetin and acetylcholine inhibited I_{ACh} in oocytes expressing $\alpha 3\beta 4$ nicotinic acetylcholine receptors (Fig. 2A and B, $n=9$ from three different frogs). The inhibition of I_{ACh} by quercetin in oocytes expressing $\alpha 3\beta 4$ nicotinic acetylcholine receptors was reversible with a negligible desensitization (Fig. 2A). Thus, these results suggest the possibility that quercetin regulates $\alpha 3\beta 4$ nicotinic acetylcholine receptor channel activity, although quercetin itself had no effect on $\alpha 3\beta 4$ nicotinic acetylcholine receptor channel activity.

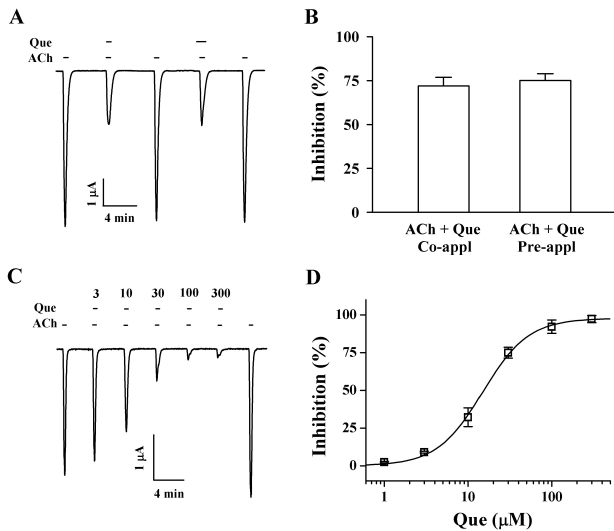


Fig. 2. Effect of quercetin (Que) on I_{ACh} in oocytes expressing $\alpha 3\beta 4$ nicotinic acetylcholine receptors. (A) Acetylcholine (ACh, 100 μ M) was first applied and then acetylcholine was co- or pre-applied with quercetin (Que, 30 μ M). Thus, co- and pre-application of quercetin with acetylcholine inhibited I_{ACh} . The resting membrane potential of oocytes was about -35 mV and oocytes were voltage-clamped at a holding potential of -80 mV prior to drug application. Traces are representative of six separate oocytes from three different frogs. (B) Co- or pre-application of quercetin did not affect differently on I_{ACh} . (C) I_{ACh} in oocytes expressing $\alpha 3\beta 4$ nicotinic acetylcholine receptors was elicited at -80 mV holding potential with indicated time in the presence of 100 μ M acetylcholine and then the indicated concentration of quercetin was co-applied with acetylcholine. (D) % Inhibition by quercetin of I_{ACh} was calculated from the average of the peak inward current elicited by acetylcholine alone before quercetin and the peak inward current elicited by acetylcholine alone after co-application of quercetin with acetylcholine. The continuous line shows the curve fitted according to the equation, $y/y_{max} = [Quercetin]/([Quercetin] + K_{1/2})$, where y_{max} , the maximum inhibition ($97.8 \pm 1.7\%$, mean \pm S.E.M.) and $K_{1/2}$ is the concentration for half-maximum inhibition (14.9 ± 0.8 μ M, mean \pm S.E.M.), and [Quercetin] is the concentration of quercetin. Each point represents the mean \pm S.E.M. ($n=9\sim 12$ from three different frogs).

Concentration-dependent effect of quercetin on I_{ACh} in oocytes expressing $\alpha 3\beta 4$ nicotinic acetylcholine receptor

Since pre-treatment of quercetin did not induce further inhibition on I_{ACh} in oocytes expressing $\alpha 3\beta 4$ nicotinic acetylcholine receptor compared to co-treatment, in next experiments we examined quercetin effects on I_{ACh} after co-treatment of quercetin with acetylcholine. In concentration-response experiments, co-treatment of quercetin with acetylcholine inhibited I_{ACh} in a concentration-dependent manner in oocytes expressing $\alpha 3\beta 4$ nicotinic acetylcholine receptor (Fig. 2C). The IC_{50} of I_{ACh} was 14.9 ± 0.8 μ M in oocytes expressing $\alpha 3\beta 4$ nicotinic acetylcholine receptors ($n=9\sim 12$ from three different frogs) (Fig. 2D).

Current-voltage relationship and voltage-independent inhibition in oocytes expressing $\alpha 3\beta 4$ nicotinic acetylcholine receptors by quercetin

As shown in Fig. 3, the current-voltage relationship induced by acetylcholine with voltage steps from -100 to

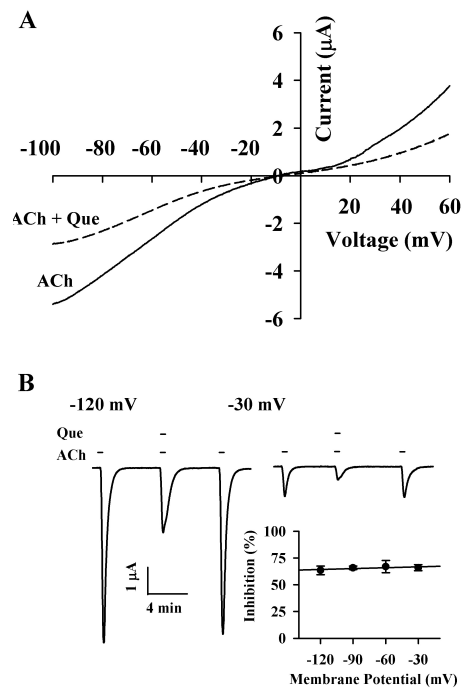


Fig. 3. Current-voltage relationship and voltage-independent inhibition by quercetin. (A) Current-voltage relationships of I_{ACh} inhibition by quercetin (Que) in $\alpha 3\beta 4$ nicotinic acetylcholine receptors. Representative current-voltage relationships were obtained using voltage ramps of -100 to $+60$ mV for 300 ms at a holding potential of -80 mV. Voltage steps were applied before and after application of 100 μ M acetylcholine in the absence or presence of 20 μ M quercetin. (B) Voltage-independent inhibition of I_{ACh} in the $\alpha 3\beta 4$ nicotinic acetylcholine receptors by quercetin. *Inset*; the values were obtained from the receptors in the absence or presence of 20 μ M quercetin at the indicated membrane holding potentials.

$+60$ mV showed a slight rectification at positive potentials in oocytes expressing $\alpha 3\beta 4$ nicotinic acetylcholine receptor. The reversal potential of $\alpha 3\beta 4$ nicotinic acetylcholine receptors was $V_r = -11.2 \pm 2.4$ mV (mean \pm S.E.M., $n=6$ from three different frogs). Co-treatment with quercetin and acetylcholine did not modify the reversal potential of $\alpha 3\beta 4$ nicotinic acetylcholine receptor with a reduction of I_{ACh} ($n=6$ from three different frogs). The inhibitory effect of quercetin on I_{ACh} in oocytes expressing $\alpha 3\beta 4$ nicotinic acetylcholine receptors was independent of the membrane holding potential (Fig. 3B). Thus, quercetin inhibited I_{ACh} by 65.4 ± 4.1 , 65.7 ± 1.7 , 66.8 ± 5.7 , and $65.8 \pm 2.8\%$ at -120 , -90 , -60 , and -30 mV membrane holding potential in oocytes expressing $\alpha 3\beta 4$ nicotinic acetylcholine receptor, respectively (Fig. 3B; $n=9\sim 12$, from three different frogs).

Noncompetitive inhibition of $\alpha 3\beta 4$ nicotinic acetylcholine receptors by quercetin

To study further the mechanism by which quercetin inhibits I_{ACh} in oocytes expressing $\alpha 3\beta 4$ nicotinic acetylcholine receptors, we analyzed the effect of 20 μ M quercetin on I_{ACh} evoked by different acetylcholine concentrations in oocytes expressing $\alpha 3\beta 4$ nicotinic acetylcholine receptors (Fig. 4). Co-application of quercetin with different concen-

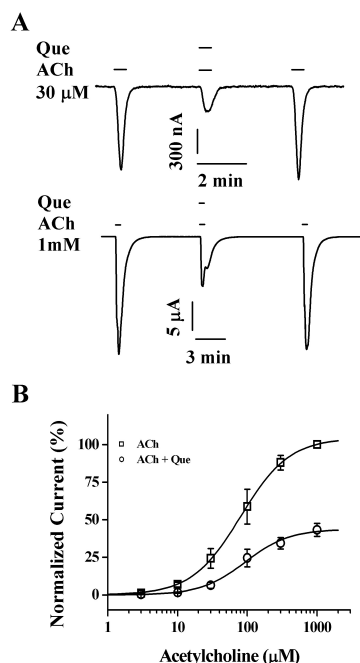


Fig. 4. Concentration-dependent effects of acetylcholine on quercetin-mediated inhibition of I_{ACh} . (A) The representative traces were obtained from $\alpha 3\beta 4$ nicotinic acetylcholine receptors expressed in oocytes. I_{ACh} of the upper and lower panels were elicited from concentration of 30 μ M ACh and 1 mM ACh at a holding potential of -80 mV, respectively. (B) Concentration-response relationships for ACh in the $\alpha 3\beta 4$ nicotinic acetylcholine receptors treated with ACh (3~1,000 μ M) alone or with ACh plus co-application of 20 μ M quercetin. The I_{ACh} of oocytes expressing the $\alpha 3\beta 4$ nicotinic acetylcholine receptors was measured using the indicated concentration of ACh in the absence (\square) or presence (\circ) of 20 μ M quercetin (Que). Oocytes were exposed to ACh alone or to ACh with quercetin. Oocytes were voltage-clamped at a holding potential of -80 mV. Each point represents the mean \pm S.E.M. ($n=9\sim 12$ /group).

trations of acetylcholine did not shift the dose-response curve of acetylcholine to the right (ED_{50} , from 81.5 ± 2.3 to 91.2 ± 6.5 μ M and Hill coefficient, from 1.25 to 1.37) in oocytes expressing $\alpha 3\beta 4$ nicotinic acetylcholine receptors, indicating that quercetin regulates $\alpha 3\beta 4$ nicotinic acetylcholine receptor channel activity with non-competitive manner ($n=9\sim 12$ from three different frogs) (Fig. 4).

DISCUSSION

In the present study, we demonstrated that (1) co- or pre-treatment with quercetin and acetylcholine inhibited I_{ACh} in oocytes expressing bovine $\alpha 3\beta 4$ nicotine acetylcholine receptor in reversible and concentration-dependent manner, (2) the inhibition of I_{ACh} by quercetin occurred in a non-competitive and voltage-independent manner in oocytes expressing $\alpha 3\beta 4$ nicotinic acetylcholine receptors, indicating that quercetin could be associated with the inhibitory regulator on I_{ACh} in oocytes expressing $\alpha 3\beta 4$ nicotinic acetylcholine receptors (Fig. 3 and 4).

From the present results, however, it is unclear precisely how quercetin acts to inhibit I_{ACh} in oocytes expressing $\alpha 3$

$\beta 4$ nicotinic acetylcholine receptor. One possible mechanism is that quercetin may act as open channel blocker of $\alpha 3\beta 4$ nicotinic acetylcholine receptors but this may be not the case because the inhibitory effect of quercetin on I_{ACh} in oocytes expressing $\alpha 3\beta 4$ nicotinic acetylcholine receptors was not voltage-dependent (Fig. 3). It is known that open channel blockers such as local anesthetics or hexamethonium are strongly voltage dependent, due to the charge that they carry in the transmembrane electrical field [21-23].

Another possibility is that quercetin may work as a competitive inhibitor by inhibiting acetylcholine binding to its binding site(s) in $\alpha 3\beta 4$ nicotinic acetylcholine receptors. In competition experiments, we observed that the presence of quercetin did not shift the concentration of acetylcholine in oocytes expressing $\alpha 3\beta 4$ nicotinic acetylcholine receptors without changing the Hill coefficient (Fig. 4). Thus, the non-competitive modulation of $\alpha 3\beta 4$ nicotinic acetylcholine receptor channel activity by quercetin shows that quercetin might have different binding or interaction site(s) as a non-competitive inhibitor at $\alpha 3\beta 4$ nicotinic acetylcholine receptors.

The third and last possibility is that quercetin might have its binding sites for the regulation of $\alpha 3\beta 4$ nicotinic acetylcholine receptor. In previous reports, we have demonstrated that the regulatory effects of quercetin on homomeric glycine, 5-HT₃ and $\alpha 7$ nicotinic acetylcholine receptor channel activities were attenuated or abolished after site-directed mutations of amino acid residues of pre-transmembrane domain of glycine and 5-HT₃ receptor or Ca²⁺-binding sites of $\alpha 7$ nicotinic acetylcholine receptor [17-19]. Based on present results and previous reports, it is likely that quercetin achieves its effect through direct interactions with $\alpha 3\beta 4$ nicotinic acetylcholine receptors. However, there might be some difficulties to construct mutant $\alpha 3\beta 4$ nicotinic acetylcholine receptors to confirm quercetin binding sites compared to the previous homomeric ligand-gated ion channels, since $\alpha 3\beta 4$ nicotinic acetylcholine receptors consist of heteromeric α and β subunits. In future, further investigations will be required to identify quercetin binding site(s) on $\alpha 3\beta 4$ nicotinic acetylcholine receptors.

Since subsets of nicotinic acetylcholine receptor channels play an important role for fast synaptic transmissions in postsynaptic sites, there are several reports on the regulations of nicotinic acetylcholine receptor channel activity using natural compounds such as flavonoids and polyphenol. Grønlien et al showed that genistein, one of flavonoids, inhibits $\alpha 7$ nicotinic acetylcholine receptor channel activity in oocytes expressing $\alpha 7$ nicotinic acetylcholine receptor [24]. Zhang et al showed that nobiletin, one of flavones inhibited catecholamine release by acetylcholine, whereas nobiletin stimulate catecholamine release via activation of Ca²⁺ channels or Na⁺/Ca²⁺ exchangers [25]. In addition, Shinohara et al demonstrated that resveratrol, one of grape polyphenol exhibited a profound effect by inhibiting on $\alpha 3\beta 4$ nicotinic acetylcholine receptor and polyphenols of *Rubus coreanum* also inhibited catecholamine release from adrenal medulla [26,27]. On the other hand, in the previous studies we found that quercetin enhanced $\alpha 7$ nicotinic acetylcholine receptor channel activity with extracellular Ca²⁺-independent manner, whereas quercetin inhibited $\alpha 9\alpha 10$ nicotinic acetylcholine receptor channel activity. In the present study, we found that quercetin inhibited I_{ACh} in oocytes expressing $\alpha 3\beta 4$ nicotinic acetylcholine receptors. Taking together of previous reports

and present results, these results indicate that quercetin might be a differential regulator of nicotinic acetylcholine receptors.

On the other hand, the previous reports have shown that various agents such as serotonin, strychnine, Ca^{2+} channel blockers, polyamines, steroids such as progesterone and hydrocortisone, ethanol, and metal ion like Zn^{2+} , regulate muscle or neuronal nicotinic acetylcholine receptors expressed in *Xenopus* oocytes [28-35]. Interestingly, the mechanism (i.e., voltage dependence or competition with acetylcholine for binding site) by which these substances regulate nicotinic acetylcholine receptors depend on the receptor subunit composition. The previous and present studies showed that quercetin inhibited heteromeric $\alpha 3\beta 4$ and $\alpha 9\alpha 10$ nicotinic acetylcholine receptor channel activities, whereas quercetin enhanced homomeric $\alpha 7$ nicotinic acetylcholine receptors. These results show the possibility that the differential effects of quercetin on subset of nicotinic acetylcholine receptor might be also due to different receptor subunit composition. In addition, since these nicotinic acetylcholine receptors may play an important role in modulating the neurotransmitter release or neuronal cell excitability induced by acetylcholine in pre-synaptic or post-synaptic site(s), quercetin-mediated differential regulations of subsets of nicotinic acetylcholine receptor channel activity might contribute to a diverse range of neuropharmacological effects of quercetin [12,13].

In summary, we found that quercetin inhibited I_{ACh} in oocytes expressing bovine neuronal $\alpha 3\beta 4$ nicotinic acetylcholine receptors. Since $\alpha 3\beta 4$ nicotinic acetylcholine receptors are closely involved in neurotransmitter release in adrenal chromaffin cells, these inhibitory effects of quercetin on I_{ACh} in oocytes expressing bovine neuronal $\alpha 3\beta 4$ nicotinic acetylcholine receptor might provide the single cellular basis for one of mechanisms for the pharmacological effects of quercetin.

ACKNOWLEDGEMENTS

This work was supported by the Basic Science Research Program through the National Research Foundation of Korea (NRF), funded by the Ministry of Education, Science and Technology (R01-2008-000-10448-0). Support was also provided by the Priority Research Centers Program through the National Research Foundation of Korea (NRF), funded by the Ministry of Education, Science and Technology (2009-0093824), and by BK21 to S.Y. Nah.

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