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<th>Inhibitory effect of β-hydroxybutyric acid on L-type Ca(^{2+}) current under β-adrenergic stimulation in guinea pig Cardiac ventricular myocytes</th>
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**INTRODUCTION**

Ketoacidosis is a life-threatening complication of both type1 and type2 diabetes\(^1\)\(^-\)\(^3\). However, the mechanism of ketoacidosis affecting cardiac function has not been fully understood. Ketone bodies include β-hydroxybutyric acid (BHB), acetoacetic acid, and acetone. In diabetic ketoacidosis, the ratio of BHB:acetone is elevated from normal (1 : 1) to values as high as (10 : 1)\(^9\). The serum BHB is 0.1 mM or less in normal humans, but could increase to a level higher than 25 mM during ketoacidosis\(^5\). Recently, hydroxy-carboxylic acid receptor 2 (HCA\(_2\)) was identified as a receptor for BHB\(^8\). This receptor was formally called GPR109A, which was initially reported as a receptor for nicotinic acid in adipocytes\(^7\)\(^-\)\(^9\), and was coupled with Gi-GTP-binding-protein\(^9\). GPR109A expression was found not only in adipocytes but also in guinea pig ventricular myocytes\(^10\).

To assess the mechanism of ketone body affecting cardiac function, we attempted to reveal that BHB stimulates Gi-coupled signal transduction in the heart. To do so, we investigated the effect of BHB on L-type Ca\(^{2+}\) current pre-augmented by β-adrenergic agonist, isoproterenol under the whole-cell voltage clamp in guinea pig ventricular myocytes. The HCA\(_2\)-mediated signal transduction may be associated with ketoacidosis-induced cardiac suppression.

**MATERIALS AND METHODS**

All experiments were performed with the approval of the Animal Research Committee of Fukushima Medical University.

The method of cell isolation and the whole-cell-voltage clamp were described previously\(^11\). Briefly
the heart was isolated from guinea pig under pentobarbital-anesthesia. Single ventricular myocytes were isolated by perfusing the heart with collagenase (5 mg/50 ml; Wako Pure Chemical Industries, Ltd., Tokyo, Japan) and protease (1 mg/50 ml; Sigma, St. Louis, MO, USA) on Langendorff apparatus. Tyrode solution contained (in mM) 140 NaCl, 5.4 KCl, 1.8 CaCl₂, 1 MgCl₂, 0.33 Na₂HPO₄, 5.5 glucose, 5 HEPES (pH 7.4 with NaOH). The pipette solution contained (in mM) 30 CsCl, 100 CsOH, 3 MgCl₂, 100 aspartic acid, 5 MgATP, 20 BAPTA, 20 HEPES (pH 7.2 with aspartic acid). Membrane currents were recorded by the whole-cell voltage clamp with a patch-clamp amplifier (CEZ2400, Nihon Kohden, Tokyo, Japan). The temperature of the bath solution was kept at 35±0.5°C with a water jacket connected to a thermostat incubator (HAAKE, E15, Berlin, Germany). Patch pipette resistance was 2-4 MΩ when filled with the pipette solution. Recording signals were filtered at 2.5-kHz bandwidth, stored online and analyzed with pCLAMP Version 9 (Axon Instruments, Union City, CA, USA).

D-(−)-beta-hydroxybutyrate was purchased from MP Biomedicals, LLC (California, USA). BHB was directly dissolved in Tyrode solution. Pertussis toxin (Wako Pure Chemical Industries, Ltd., Tokyo, Japan) at 5 μg was activated by mixing with 50 μl dithiothreitol (100 μM) and 450 μl pipette solution and incubated at 37°C for 15-20 min. Then, 4.5 ml pipette solution was added, so that the final pertussis toxin (PTX) concentration of in the pipette solution was 1 μg/ml. Each experiment was repeated 4 to 5 times. The data were expressed as the means±S.E.M. Statistical significance between two groups was evaluated using Welch’s t-test.

RESULTS

Effect of BHB on L-type Ca²⁺ current pre-augmented by isoproterenol

First we found that BHB alone did not affect the L-type Ca²⁺ current in single ventricular myocytes of guinea pig (data not shown). Then, we tested BHB on the L-type Ca²⁺ current pre-augmented by isoproterenol. A family of depolarizing square voltage pulses were given from the holding potential of −40 mV to +40 mV for 400 ms duration (Fig. 1A inset). Isoproterenol at 0.1 μM in the extracellular Tyrode solution increased L-type Ca²⁺ current (Fig. 1A and 1B). When 10 mM BHB was added to isoproterenol, the Ca²⁺ current was gradually decreased (Fig. 1C). Finally, 1 μM nifedipine was added to completely inhibit the Ca²⁺ current (Fig. 1D). In some cells, positive background currents increased as shown in Fig. 1C and Fig. 1D. These currents were most likely isoproterenol-induced Cl⁻ currents. We measured the magnitude of the Ca²⁺ current between the peak and at the inactivated level at the end of 400 ms depolarizing pulse at each voltage in order to avoid contamination of the isoproterenol-induced Cl⁻ current. The I-V curves of the Ca²⁺ current were plotted (Fig. 1E). Since the peak potential of the Ca²⁺ current varied between −20 mV and 10 mV among different cells, we employed the value at 10 mV for further analysis.

A concentration-response curve of BHB was obtained (Fig. 2). The peak value of Ca²⁺ current in isoproterenol was set at 100%. BHB at 10 mM significantly inhibited the Ca²⁺ current to 48.9±4.4% (n=5, p<0.001) of the control in isoproterenol. The inhibitory effect of BHB was concentration dependent, and the IC₅₀ value of BHB was approximately 1.1 mM. Since the IC₅₀ value of BHB to HCA₂ receptor was reported to be 0.7 mM, our value of 1.1 mM is close enough to that of HCA₂. Therefore, our result suggested that BHB reduced Ca²⁺ current pre-augmented by isoproterenol possibly via HCA₂ receptor in guinea pig ventricular myocytes.

Effect of nicotinic acid on Ca²⁺ current pre-augmented by isoproterenol

If the above result was mediated by HCA₂ receptor, nicotinic acid, another ligand reported for HCA₂, should behave similarly to BHB. Therefore, we tested nicotinic acid instead of BHB. When nicotinic acid at 30 μM was added in the external solution, it inhibited the Ca²⁺ current to 73% of isoproterenol-treated control in this cell (Fig. 3). The I-V curves were superimposed in Fig. 3E. In 5 different cells, nicotinic acid inhibited the Ca²⁺ current to 67±6% of the isoproterenol-treated control. Since the effects of nicotinic acid and BHB were similar, this suggests that the effects may be mediated by HCA₂ receptor.

Effect of pertussis toxin

HCA₂ receptor is coupled with Gi-GTP-binding protein. Pertussis toxin (PTX) is a specific inhibitor of Gi. If the effects of BHB and nicotinic acid on the Ca²⁺ current were mediated by Gi, those effects should be suppressed by PTX. As shown in Fig. 4A, in the presence of activated PTX in the
pipette solution, 10 mM BHB did not significantly decrease the pre-augmented Ca\(^{2+}\) current. Similar results were obtained with 30 μM nicotinic acid (Fig. 4B). Nicotinic acid decreased the isoproterenol-augmented Ca\(^{2+}\) current only to 90.8±6.3% (n=4) in the presence of PTX (Fig. 4B). These results further suggest that the effects of BHB and nicotinic acid were mediated by Gi-protein.

**DISCUSSION**

In this study, we obtained the following results. (1) BHB inhibited the L-type Ca\(^{2+}\) current pre-augmented by isoproterenol, and this effect was BHB-concentration dependent. (2) Nicotinic acid also inhibited the pre-augmented Ca\(^{2+}\) current in a similar manner to BHB. (3) Pertussis toxin significantly inhibited the effects of BHB and nicotinic acid. (4) Estimated IC\(_{50}\) value of 1.1 mM BHB was close to a reported value of 0.7 mM for HCA\(_2\) receptor. From these results, we suggest that BHB and nicotinic acid activate Gi-mediated signal transduction via HCA\(_2\) in guinea pig cardiac ventricular cells.

L-type Ca\(^{2+}\) channels in cardiac myocytes are regulated dually; stimulation mediated by Gs-coupled receptors and inhibition via Gi-coupled receptors. The former is represented by β1-adrenergic receptor and the latter by muscarinic M2.
Fig. 2. Concentration-inhibition curve of BHB on the L-type Ca\(^{2+}\) current pre-augmented by isoproterenol (Iso).

The current magnitude was measured at 10 mV. Data are mean ± S.E.M of 5 to 7 cells.
Fig. 3. Inhibitory effect of nicotinic acid (NA) on the Ca$^{2+}$ current pre-augmented with isoproterenol. A: Control Ca$^{2+}$ currents before isoproterenol. Voltage pulses are the same as those shown in Fig. 1. B: Ca$^{2+}$ currents augmented by 0.1 $\mu$M isoproterenol (Iso). C: Ca$^{2+}$ currents in the presence of 30 $\mu$M nicotinic acid (NA) added to Iso. D: Nifedipine (Nif) at 1 $\mu$M was added to completely inhibit the Ca$^{2+}$ current. A-D were obtained from the same cell. E: Current-voltage relations obtained from the data in Fig. 3A-D. Control (X) ; 0.1 $\mu$M isoproterenol (■) ; isoproterenol + nicotinic acid (30 $\mu$M) (▲) ; isoproterenol + nicotinic acid + nifedipine (1 $\mu$M) (●).

Fig. 4. Effects of BHB (A) and nicotinic acid (NA) (B) in the presence of pertussis toxin (PTX) in the pipette solution. Control Ca$^{2+}$ currents before isoproterenol was set as 100%. The current magnitude was measured at 10 mV. BHB : n=5 # ; p < 0.05 vs. Iso 0.1 $\mu$M NS ; no significance. NA : n=4 # ; p < 0.05 vs. Iso 0.1 $\mu$M.
KETONE BODY ACTIVATES HCA₂ RECEPTOR IN GUINEA PIG CARDIAC MYOCYTES

Fig. 5. A schematic representation of the effects of BHB and nicotinic acid (NA) on L-type Ca²⁺ channel. Isoproterenol (Iso) binds to β₁ adrenergic receptor and activates adenylate cyclase (AC) via activation of Gs. HCA₂ receptor activates Gi and suppresses AC. cAMP activates protein kinase A (PKA) and phosphorylates Ca²⁺ channel. Phosphorylation of Ca²⁺ channel enhances the Ca²⁺ current in cardiac myocytes. Pertussis toxin (PTX) inhibits Gi.

bonate decreases ketone bodies including BHB³¹. More aggressive administration of sodium bicarbonate may be revalidated in severe ketoacidosis with high levels of catecholamine and BHB in the blood.

Limitations of the present study include the following. Whether our results obtained from guinea pig cardiac myocytes are applicable to human heart should be evaluated. It is not so common to see ketoacidosis so advanced to accompany cardiac dysfunction.

In conclusion, BHB stimulates Gi signaling pathway possibly via HCA₂ receptors and decrease the Ca²⁺ current pre-augmented by adrenergic β stimulation in guinea pig cardiac ventricular myocytes. This BHB effect may be responsible for suppression of cardiac function in ketoacidosis. Further investigation of the BHB effect on cardiac function is necessary.

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