The inotropic effect of isoniazid on isolated atria of guinea-pigs

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Abstract

The incidence of heart failure is increasing every year, while there is no ideal inotropic agent yet available. Therefore the search for an ideal inotropic agent is still needed. Isoniazid (INH) has been widely used as an anti-tuberculosis agent that has similar structure to the $K^+$ channel blocker 4-aminopyridine. Considering that $K^+$ channel blocker may prolong depolarization leading to a more influx of $Ca^{2+}$ into the cell and increase cardiac contractility, the effect of INH on cardiac contractility were investigated. In guinea-pig isolated atria preparations, INH produced concentration-dependent increase contractility. The maximal concentration of INH in increasing force of contractility is 100 mM. This is approximately an equieffective concentration compared to 1 uM of adrenaline induced cardiac contractility. The difference is that INH has no effect on the rate of contraction. A non-selective beta-adrenoceptor antagonist propranolol (1 uM) inhibits the inotropic action of adrenaline (1 uM) but with no effect on INH (100 mM) induced cardiac contractility. It is speculated that INH possesses a positive inotropic effect. This effect is not associated with the activation of cardiac beta-adrenoceptor by catecholamine as a consequence of monoamine oxidase inhibition, but may be through the prolongation of depolarization of the myocardial cells by blocking of $K^+$ channels.

Keywords: Isoniazid, 4-aminopyridine, $K^+$ channels blocker, positive inotropic
of these agents are safe and effective in long term trial. Therefore, looking for an effective inotropic agents is needed in conquering heart failure.

Isoniazide (INH) is an oral anti-tuberculosis which has been use world-wide since 1945. The structure of INH very similar to that of 4-aminopyridine, a potassium channel blocking agent (Fig.1). It has been well established that by blocking of potassium channel, 4-aminopyridine is able to prolong cell depolarization and excitation. In cardiac cell, prolonged depolarization may allow more calcium influx into the cell leading to increase contractility.

In this study, the effect of INH in increasing cardiac contractility was investigated using isolated guinea-pig atria.

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<th>Isoniazid</th>
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_Figure 1. The chemical formulae of isoniazid and 4-aminopyridine (K⁺ - channel blocker)_

METHODS

Guinea-pigs (300-350 g) of either sex were killed and the hearts rapidly removed. The atria were dissected free and suspended between a tissue hook and an isotonic transducer in an organ bath containing 10 ml fresh physiological salt solution (PSS), maintained at 37 degree Celcius and bubbled with a mixture of 95% O2/5% CO2.

The atria were placed under basal tension of 1 g wt. The spontaneous atrial contraction were measured with a strain gauge transducer for recording the rate and force of atrial beating. The isolated atria were allowed to equilibrate for a period of 20 min before any experimental procedures were commenced, during each time the bathing solution was regularly exchanged. The responsiveness of the tissue was then tested by the addition to the organ bath of adrenaline (1 uM). Responses to addition of adrenaline was allowed to develop fully, after which the agonist was removed by repeatedly exchanging the PSS in the organ bath.

Concentration-response relationships for the positive inotropic effect of isoniazide (INH) was established by its cumulative addition to the organ bath until obtaining the maximal response.

To examine the effect of beta-adrenoceptor blocking agent on the positive inotropic response of INH, the atria was exposed two times by either INH or adrenaline, 30 min apart. Propranolol (1 uM) was added to the atrial bathing solution 15 min before commencing to construct the second exposure to the test agonist; it then remaining present throughout the rest of the experiment.

Data were expressed as mean and standard error of the means (+ s.e.m), and were analyzed by paired, two-tailed Student’s t-test. In all cases, level of probability (P) of less than 0.05 was taken to indicate a statistical significance.

RESULTS

Positive inotropic response of isoniazide (INH)

The mean resting forces and contraction rate of isolated guinea-pigs atria were 0.5 g/weight (s.e.m. = 0.07, n = 5) and 235 beats/min (s.e.m. = 16, n = 5), respectively.

INH (1-100 mM) produced concentration-dependent increases of contractility (Fig 2a,b) without influencing the rate of beating. The mean increase in force of contraction to 10, 30, 100 and 300 mM of INH were 6 ± 2,5; 26 ± 2; 69 ± 3 and 72 ± 4%, respectively. The response being maximal at a concentration of 100 mM.

Effect of propranolol on INH-induced positive inotropic response

Concentration of INH (100 mM) and adrenaline (1 uM) which produced similar increases in force of contraction were selected.
Figure 2. The effect of isoniazid (INH) and adrenaline on cardiac contractility.
(a) INH (10-300 mM) produced concentration dependent increase of contractility
(b) The maximal effect of INH was achieved at 100 mM which is equipotent to adrenaline 1 uM
The beta-adrenoceptor antagonist propranolol (1 \( \mu \)M) when introduced 15 min before the second exposure of guinea-pig atria had no significant effect on INH (100 mM) induced contractility (P>0.05, unpaired t-test). The mean value of percent increase in force of contraction to 100 mM INH alone and in the present of propranolol were 64 ± 2.5 and 61 ± 3\%, respectively. However, propranolol significantly reduced the adrenaline (1 \( \mu \)M) induced contractility (P<0.05, unpaired t-test). The mean value of percent increase in force of contraction to adrenaline (1 \( \mu \)M) given alone and in the present of propranolol were 72 ± 4 and 32 ± 2.5\%, respectively (Fig. 3).

DISCUSSION

Myocardial cell membrane exhibits selective permeability to ions, positively charged such as Na*, K*, Ca** or negatively charged such as Cl-, creating an electrical potential across the cell membrane. The ions move across the cell membrane through ion channels.

An action potential responsible for initiating each "heart contraction" is due to a sudden increase in membrane conductance to Na*, this allows rapid influx of Na* into the cell which in turn is responsible for activating voltage-dependent calcium channels to allow Ca** to enter the cell, triggers release of large amount of Ca** from sarcoplasmic reticulum. Ca** binding by troponin results in contractile activity of cardiac fibers. Thereby, there is a rapid repolarization owing to the activation of an outward K* current and cessation of the inward Ca** current.

It has been shown that blockade of outward K+ conductance with potassium channels blocking agent such as 4-aminopyridine enables the influx of Ca** to reinforce the applied depolarization and to become regenerative. In cardiac muscle, 4-amino-pyridine has been shown to increase contractility through this mechanism.

In view of the structural similarity of INH and 4-aminopyridine, INH may have K* channel-blocking action and thus prolong depolarization in excitable cells. In this study, as expected, INH clearly produced a concentration dependent increase of cardiac contractility, or in other word, INH possesses a positive inotropic effect. Interestingly, there was no alteration of heart beat occurred in any given doses of INH. The underlying mechanism is still unknown.

Previous studies have shown that a derivative of INH, iproniazid, inhibited monoamine oxydase (MAO) and

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**Figure 3.** The effect of propranolol (1 \( \mu \)M) on the INH (100 mM, \( n = 5 \)) and adrenaline (1 \( \mu \)M, \( n = 5 \)) induced contractility. (*) indicated significant differences from corresponding control (p<0.05)
lead to an increase of catecholamine release. It has also been reported that euphoria or convulsion in patients with tuberculosis who receive INH are due to the increase level of catecholamine as a consequence of MAO-inhibition. Therefore, it is possible that the positive inotropic effect of INH is due to the activation of the beta-adrenoceptor within the heart by catecholamine.

In order to investigate this, the positive inotropic effect of INH was tested by the presence of non-selective beta-adrenoceptor antagonist propranolol. The effect of propranolol was also compared to adrenaline (1 uM), which is approximately equieffective concentration to INH (100 mM) to induce cardiac contractility.

In the presence of propranolol, INH (100 mM) still produced a significant positive inotropic effect. In contrast, propranolol did inhibit the positive inotropic effect of adrenaline (1 uM). This results clearly demonstrate that INH increases cardiac contractility which is not due to activation of cardiac beta-adrenoceptor by increase level of catecholamine. This is in accord to the finding that INH has no effect on heart rate.

In conclusion, INH possesses a positive inotropic effect, which is not through the activation of beta-adrenoceptor in the heart, but may be through the prolongation of depolarization of the myocardial cells by blocking of K+ channels. Further experiments are needed to confirm this hypothesis.

REFERENCES