

The role of hydrogen sulfide in homocysteine-induced cardiodynamic effects and oxidative stress markers in the isolated rat heart

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This study aimed to assess the role of H₂S in homocysteine-induced cardiodynamic effects in the isolated rat heart. The hearts were retrogradely perfused according to the Langendorff technique. The maximum and minimum rates of pressure in the left ventricle (dp/dt max, dp/dt min), systolic and diastolic left ventricular pressures (SLVP, DLVP), heart rate (HR), and coronary flow (CF) were measured. A spectrophotometrical method was used to measure the following oxidative stress markers: index of lipid peroxidation (thiobarbituric acid reactive substances, TBARS), nitrite level (NO₂⁻), superoxide anion radicals (O₂^{•-}), and hydrogen peroxide (H₂O₂) concentrations. The administration of 10 μmol/l DL-homocysteine (DL-Hcy) alone decreased dp/dt max, SLVP, and CF but did not change any oxidative stress parameters. The administration of 10 μmol/l DL-propargylglycine (DL-PAG) decreased all cardiodynamic parameters and increased the concentration of O₂^{•-}. The co-administration of DL-Hcy and DL-PAG induced a significant decrease in all estimated cardiodynamic parameters and decreased the concentration of NO₂⁻ and O₂^{•-} but increased the levels of TBARS and H₂O₂. Homocysteine shows a lower pro-oxidative effect in the presence of hydrogen sulfide (H₂S), which indicates a potential anti-oxidative capacity of H₂S.

Keywords: cardiodynamics, homocysteine, H₂S, Langendorff technique, oxidative stress

Introduction

Homocysteine (Hcy) is a sulfur-containing amino acid whose elevated values have a high correlation with the increased risk for cardiovascular diseases (CVDs). For this reason, hyperhomocysteinemia (HHcy) is suggested as a new and independent risk factor for developing these conditions (38, 39). HHcy occurs as a consequence of an impaired remethylation process or a deficiency in Hcy metabolic enzymes [cystathionine β-synthase (CBS), cystathionine γ-lyase (CSE), and 3-mercaptopyruvate sulfur transferase] (19). These three enzymes break down Hcy into hydrogen sulfide (H₂S) (33). Together with nitric oxide (•NO), carbon monoxide (CO), and methane, H₂S belongs to a group of gaseous messengers whose importance in different pathophysiological processes has rapidly increased over the years (11, 23, 24, 30, 34).

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In the cardiovascular system (CVS; the heart and aorta), H₂S is generated by CSE (45) and manifests effects opposite to those of Hcy (28). H₂S acts as a vasorelaxant (45) and an inhibitor of hypertrophy and fibrosis of vascular smooth muscle cells (17). Taking into consideration that HHcy can be caused by the inhibition of metabolic enzymes that produce H₂S from Hcy (19), the question arises whether the blockade of H₂S synthesis influence homocysteine-induced cardiac effects.

It is considered that hyperhomocysteinemia has adverse effects on the endothelium of blood vessels, mainly by increasing the production of free radicals, superoxide anion radicals (O₂^{•-}), and hydrogen peroxide (H₂O₂) (16, 22, 36, 39). The connection between the metabolism of homocysteine, free radicals, and H₂S in the CVS is still insufficiently investigated (31). However, much research has proven that H₂S is able to directly scavenge reactive oxygen species (ROS) and downregulate the ROS-producing enzymes, which indicates that H₂S is a potent vasodilator and has powerful anti-inflammatory, antioxidant, and antiapoptotic effects (2, 20, 21, 41–43, 45).

To reveal the mechanisms by which homocysteine influences myocardium and coronary circulation, we aimed to investigate the role of H₂S and oxidative stress in these effects of homocysteine.

Materials and Methods

Eight-week-old male Wistar albino rats with a body mass of 250 ± 30 g were kept in standard laboratory conditions (air temperature 23 ± 10 °C, relative humidity 50%), 12:12 dark–light cycle (the beginning of the light period was at 9 a.m.) and given water and food ad libitum. All research procedures were carried out in accordance with European Directive for welfare of laboratory animals (No. 86/609/EEC) and principles of good laboratory practice, approved by ethical committee of the Faculty of Medical Sciences (Reg. No. 01-994/4), University of Kragujevac, Serbia. The rats were divided into three experimental groups (12 animals per group). The hearts of the rats were excised and perfused according to the modified Langendorff technique at constant pressure conditions (Experimetria Ltd., Budapest, Hungary) as described previously (48). The animals were premedicated with heparin as an anticoagulant, and briefly, under ether anaesthesia, were sacrificed by cervical dislocation (Schedule 1 of the Animals/Scientific Procedures, Act 1986, UK). After emergency thoracotomy and rapid cardiac arrest by superfusion with ice-cold isotonic saline, the hearts were rapidly excised; the aortas were cannulated and retrogradely perfused at the constant pressure of 70 cm H₂O. The composition of the non-recirculating Krebs–Henseleit perfusate was as follows (in mmol/l): NaCl 118, KCl 4.7, CaCl₂ × 2H₂O 2.5, MgSO₄ × 7H₂O 1.7, NaHCO₃ 25, KH₂PO₄ 1.2, glucose 11, and pyruvate 2, equilibrated with 95% O₂ + 5% CO₂ and warmed to 37 °C (pH 7.4). Immediately after the normal heart rhythm returned, the sensor (transducer BS4 73-0184, Experimetria Ltd., Budapest, Hungary) was inserted through the newly damaged left atrium and mitral valve into the left ventricle for continuous monitoring of cardiac function.

To test coronary vascular reactivity, all hearts were challenged by short-term occlusions (5–30 s) followed by a bolus injection of 5 mmol/l adenosine [60 µl at a flow rate of 10 ml/min to elicit maximum coronary flow (CF)] during the stabilization period. Hearts were discarded if the flow did not increase by 100% over the control value for both tests (approximately 25% of the hearts). CF was measured using flowmetry. When the flow was

Table 1. Cardiodynamic parameters

	dp/dt max (mmHg/s)	dp/dt min (mmHg/s)	SLVP (mmHg)	DLVP (mmHg)	HR (bpm)	CF (ml/min)
(a) Before and after the application of DL-homocysteine						
Control (mean ± SE)	1246.5 ± 233.8	-475.2 ± 143.0	55.6 ± 11.2	8.4 ± 4.2	140.1 ± 32.9	5.2 ± 1.1
DL-Hcy (mean ± SE)	960.5 ± 258.0*	-415.8 ± 209.1	42.6 ± 9.6*	7.6 ± 5.1	134.0 ± 31.6	4.4 ± 1.1*
(b) Before and after the application of DL-propargylglycine						
Control (mean ± SE)	1964.5 ± 78.5	-1218.1 ± 87.2	61.3 ± 1.8	8.4 ± 0.7	244.9 ± 11.3	9.1 ± 0.3
DL-PAG (mean ± SE)	1831.5 ± 85.1*	-1052.5 ± 43.2*	58.9 ± 2.1*	7.9 ± 0.6*	232.9 ± 10.7*	8.5 ± 0.3*
(c) Before and after the co-application of DL-Hcy and DL-PAG						
Control (mean ± SE)	1615.7 ± 87.1	-926.1 ± 158.9	56.7 ± 2.7	5.1 ± 0.6	240.5 ± 3.5	8.2 ± 0.9
DL-Hcy + DL-PAG (mean ± SE)	1523.1 ± 99.9*	-773.1 ± 162.4*	53.6 ± 2.9*	4.2 ± 0.7*	228.3 ± 6.3*	8.0 ± 0.8*

The values are expressed as the mean ± SE.

*Statistical significance ($p < 0.05$)

Table II. The levels of pro-oxidants in coronary venous effluent

	O ₂ ^{•-} (nmol/ml)	H ₂ O ₂ (nmol/ml)	NO ₂ ⁻ (nmol/ml)	TBARS (μmol/ml)
(a) Before and after the application of DL-homocysteine				
Control (mean ± SE)	3.84 ± 0.83	0.48 ± 0.05	1.43 ± 0.24	0.75 ± 0.17
DL-Hcy (mean ± SE)	5.21 ± 3.84	0.45 ± 0.02	1.40 ± 0.20	0.86 ± 0.21
(b) Before and after the application of DL-propargylglycine				
Control (mean ± SE)	25.35 ± 0.86	35.76 ± 0.52	2.80 ± 0.41	23.46 ± 1.26
DL-PAG (mean ± SE)	26.30 ± 1.05*	34.95 ± 1.05	2.77 ± 0.36	24.82 ± 1.00
(c) Before and after the co-application of DL-Hcy and DL-PAG				
Control (mean ± SE)	81.46 ± 1.85	6.47 ± 0.45	18.11 ± 0.85	6.25 ± 0.83
DL-Hcy + DL-PAG (mean ± SE)	63.36 ± 1.89*	8.02 ± 0.49*	15.43 ± 0.74*	12.01 ± 0.96*

The values are expressed as the mean ± SE.

*Statistical significance ($p < 0.05$)

considered stable (three measurements with the same value), coronary effluent samples were collected.

After perfusion in the absence of any substance (control conditions), the hearts were perfused as follows:

Group 1: 10 μmol/l DL-homocysteine (DL-Hcy)

Group 2: 10 μmol/l DL-propargylglycine (DL-PAG) – irreversible CSE inhibitor

Group 3: 10 μmol/l DL-Hcy + 10 μmol/l DL-PAG

Groups were chosen to separate the independent influence of homocysteine (Group 1) or the endogenous CSE/H₂S pathway (Group 2) and to estimate the role of H₂S in the cardiac effects of homocysteine (Group 3). Each of the applied substances was perfused for a duration of 5 min. The substances were first dissolved in Krebs–Henseleit solution and, after control conditions, applied directly to the heart by perfusion pump.

By placing the sensor in the left ventricle, the following parameters of myocardial function were continuously registered:

- (1) Maximum rate of pressure development in the left ventricle (dp/dt max);
- (2) Minimum rate of pressure development in the left ventricle (dp/dt min);
- (3) Systolic left ventricular pressure (SLVP);
- (4) Diastolic left ventricular pressure (DLVP);
- (5) Heart rate (HR).

CF was measured using the flowmetric method and expressed in ml/min.

Coronary venous effluent (5 ml) was collected at the same time, at the end of control conditions and during the last minute of perfusion with any of the test compounds. Oxidative stress parameters [the index of lipid peroxidation (thiobarbituric acid reactive substances, TBARS), the superoxide anion radical (O₂^{•-}), H₂O₂, and nitrite (NO₂⁻)] were determined in coronary venous effluent samples using the spectrophotometric method (Specord S-600 Analytik Jena).

Determination of the index of lipid peroxidation (TBARS)

The degree of lipid peroxidation in the coronary venous effluent was estimated by measuring TBARS using 1% thiobarbituric acid in 0.05 NaOH incubated with the coronary effluent at 100 °C for 15 min and read at 530 nm. Krebs–Henseleit solution was used as a blank probe (27).

Determination of superoxide anion radical ($O_2^{\cdot-}$)

The level of the superoxide anion radical ($O_2^{\cdot-}$) was measured by nitro blue tetrazolium reaction in Tris-buffer with coronary venous effluent and read at 530 nm. Krebs–Henseleit solution was used as a blank probe (4).

Determination of hydrogen peroxide (H_2O_2)

The measurement of H_2O_2 was based on the oxidation of phenol red by H_2O_2 in a reaction catalyzed by horseradish peroxidase (HRPO) (29). The volume of 200 μ l of perfusate was precipitated with 800 μ l of fresh phenol red solution along with 10 μ l 1:20 HRPO (made ex tempore). An adequate volume of Krebs–Henseleit solution was used for a blank probe (instead of coronary venous effluent). The level of H_2O_2 was measured at 610 nm.

Determination of the nitrite level (NO_2^-)

The nitrite level (NO_2^-) was measured as an index of \cdot NO production using the Griess reagent. A total of 0.5 ml of perfusate was precipitated with 200 μ l of 30% sulfosalicylic acid, vortexed for 30 min, and centrifuged at $3,000 \times g$. Equal volumes of the supernatant and Griess reagent, containing 1% sulfanilamide in 5% phosphoric acid, 0.1% naphthalene ethylenediamine dihydrochloride were added, incubated for 10 min in the dark, and read at 543 nm. The nitrite levels were calculated using sodium nitrite as the standard (15).

All drugs were purchased from Sigma-Aldrich Chemie GmbH, Germany.

Statistical data processing

Statistical processes have been performed using the SPSS Statistics 19 program (IBM Corp., Armonk, USA). Variables are presented as mean values \pm standard deviation. We used non-parametric methods (Wilcoxon test) to calculate the significance of the differences between the data. A value of $p < 0.05$ is considered statistically significant. After statistical processing of the data, the results were shown in tables.

Results

The effects of DL-Hcy, DL-PAG, and DL-Hcy + DL-PAG on myocardial function parameters in the isolated rat heart

The administration of DL-Hcy on an isolated rat heart induced a significant decrease of the dp/dt max, SLVP and CF, compared to control conditions (Table Ia).

The administration of DL-PAG induced a significant decrease of all the estimated cardiodynamic parameters (Table Ib).

The effects of DL-Hcy and DL-PAG co-administration also induced a significant decrease of all the estimated cardiodynamic parameters (Table Ic).

The effects of DL-Hcy, DL-PAG, DL-Hcy + DL-PAG on oxidative stress markers in the isolated rat heart

Perfusion with DL-Hcy showed that there were no significant changes in the levels of all the estimated oxidative stress parameters in the coronary venous effluent (Table IIa).

The administration of DL-PAG induced a significant increase in the levels of $O_2^{\cdot-}$ in the coronary venous effluent, whereas other parameters of oxidative stress showed no statistically significant differences (Table IIb).

The effects of DL-Hcy and DL-PAG co-administration on oxidative stress parameters induced a significant decrease in the levels of NO_2 and $O_2^{\cdot-}$, and a significant increase in TBARS and H_2O_2 (Table IIc).

Discussion

One of the objectives of this study was to examine the direct effects of the basic form of homocysteine on the myocardium and the coronary circulation in the isolated rat heart. Although it has been well known that homocysteine has an increasingly significant role in the pathophysiology of CVDs (1, 12), the majority of investigations examined the influence of homocysteine on the vascular system (3, 25), while the effects on the myocardium and the coronary circulation are little known.

The results obtained show that homocysteine leads to a significant reduction of the following major cardiodynamic parameters: dp/dt max and SLVP, and also CF (Table Ia).

Related to our research, and with respect to the similarity with our experimental model, Kennedy et al. (18) studied the acute effects of homocysteine (30–100 μ mol) on systolic function and contractibility of myocardium; they found similar results in that there was a significant reduction of systolic function and contractive abilities (dp/dt max) in the isolated rat heart, but no changes in diastolic function were recorded, regardless of the applied dose.

Wan et al. (40) examined the effects of homocysteine in a cardiac ischemia model. The study showed that homocysteine reduced the contractible ability (dp/dt max), but without the effect on SLVP, DLVP and HR, which was very similar to our findings. Although our research was performed on healthy hearts, it is obvious that homocysteine in both normoxic and hypoxic conditions, either acutely or chronically administered, has a direct negative effect on cardiac function. One of the potential explanations for such results may be the fact that DL-Hcy, via still unknown mechanisms, showed cardiotoxic effects on cultured isolated cardiomyocytes (35).

On the other hand, it has been documented that gasotransmitters (1NO , CO, and H_2S) are increasingly important in various pathophysiological states in the heart. Using the same Langendorff apparatus, Bak et al. (5–7) have shown that CO could be a powerful tool for functional recovery of post-ischemic rat or mouse heart. H_2S is a biologically active gas whose role in the effects of homocysteine on the heart was investigated. In the conditions when there is enough methionine and homocysteine, homocysteine is subject to the process of transsulfuration when it converts into cystathionine with the help of the enzyme CBS in the presence of serine. Then, under the influence of the other enzyme CSE, cystathionine metabolizes to L-cysteine to finally generate H_2S , α -keto barbiturates and ammonium (32). It implies that this signal molecule may have an influence on the effects of homocysteine on the CVS.

The effects of endogenous and exogenous H_2S on the CVS have been intensively investigated, but the data obtained are still incoherent and without precise conclusions (32, 46). Much less is known about the effects of H_2S on the myocardium and the coronary circulation. We decided to inhibit the enzyme CSE, which is supposed to be an endogenous source of H_2S in the heart (13), by the administration of DL-PAG. The administration of

DL-PAG leads to a considerable reduction of all the examined cardiodynamic parameters, although the function and perfusion of the myocardium were weakened (Table Ib).

In comparison to our study, Carson et al. (9) studied the effects of H₂S on the myocardium and coronary perfusion in the same experimental model (Langendorff retrograde perfusion). This group showed that exogenously administered H₂S induced a dose-dependent reduction of dp/dt max and min like in the HR of the isolated rat heart, while paradoxically increasing the CF. In comparison to our study, these results could be due to different experimental approaches (exogenous uptake).

Although they are still unclear, the mechanisms by which H₂S influences the heart and coronary circulation probably include activation of K_{ATP} channels in both cardiomyocytes (47) and coronary smooth muscles (37). Despite the insufficient knowledge of the role of H₂S in the heart functioning, the coronary circulation and the discrepancy between the existing data, this gasotransmitter can obviously be synthesized in the heart, and together with NO and CO may be significant in the pathophysiology of the heart.

In our study, the potential role of H₂S in the effects of homocysteine on the myocardium and coronary vascular tonus was examined by the combined administration of homocysteine with a CSE inhibitor (DL-PAG). The administration of DL-PAG with DL-Hcy statistically significantly reduced the values of all cardiodynamic parameters (Table Ic).

In a study that used an experimental model similar to ours, the influence of H₂S on the infarction of myocardium in rats with hyperhomocysteinemia was examined (10). This research showed that in the case of development of hyperhomocysteinemia, myocardium expression and activity of CSE, and hence H₂S, were significantly reduced. On the other hand, the administration of H₂S during the period prior to ischemia improved functional recovery of the heart and reduced the bulk of infarction, which showed that H₂S could be significant in the ischemic preconditioning and reperfusion period.

The connection between homocysteine and oxidative stress has long been known. Like all thiols, homocysteine easily enters the processes of oxidation such as self-oxidation in plasma. During the oxidation of the sulfhydryl group, homocysteine generation of various ROS occurs, above all O₂^{•-}, H₂O₂, and OH^{•-}. These ROS then initiate lipid peroxidation of vascular endothelial cells and their morphofunctional disorder. The O₂^{•-} interacts with NO produced by the same cells, thereby generating even more toxic reactive nitrogen species – ONOO^{•-}. Thus, the damage to endothelial cells impairs the control of vascular tonus and leads to the endothelial dysfunction (16, 22, 36, 39).

One of the studies showed that homocysteine increased the production of pro-oxidation markers (TBARS) and reduced the activity of anti-oxidation enzymes of protection in the rat heart (26). The results of our study indicate that after the application of the basic homocysteine form (Table IIa), there were no significant changes in the values of all examined oxidative stress markers. In our case, the effects of acutely administered homocysteine were examined, whereas the previous research treated chronic effects. In addition, the dose of homocysteine in the present study was incomparably smaller to lead to possible damage. Additionally, it should be noted that our model was based on physiological conditions of cardiac function.

We wanted to examine a potential role of H₂S in the stimulation of oxidative stress. The majority of available data show that H₂S often reduces the production of ROS, i.e., it has an anti-oxidative character in the CVS (2, 20, 21, 41–43, 45). In our study, the administration of the inhibitor CSE/H₂S system (DL-PAG) induced the increased release of superoxide anion radicals in coronary venous effluent, while the levels of other parameters in the oxidative

stress did not show statistically significant change in comparison to the control conditions (Table IIb).

Our results are in accordance with those of Gao et al. (14), who examined the influence of exogenous (NaHS) and endogenous H₂S (DL-PAG) on the development of oxidative stress during ischemia and reperfusion of myocardium. They showed that the H₂S donor, NaSH, was related to the reduced lipid peroxidation, ROS production, and consequent decrease of the size of infarction. On the other hand, the blockage of synthesis of endogenous H₂S induced opposite effects. These findings show that H₂S could have a cardioprotective effect.

Based on these results, we can assume that H₂S, either exogenous or endogenous, seems to possess a certain anti-oxidative capacity that should be confirmed in further studies. Additionally, future research should be extended to use a wider range of administered doses to determine possible dose-dependence of the obtained effects.

It remains little known how significant H₂S is in the oxidative effects in the presence of homocysteine. According to the latest data, H₂S can reduce the endothelial stress induced by homocysteine (8, 10). Namely, endothelial cells are capable of secreting CBS and CSE enzymes, which then, while circulating, actively generate H₂S from homocysteine (8). In the presence of homocysteine, the H₂S synthesized in this way prevents oxidative damage and consequent dysfunction of endothelial cells (8, 10, 44).

Yan et al. (44) examined whether H₂S could influence the cytotoxicity and oxidative stress induced by homocysteine. Cultures of smooth aortic muscles were exposed to homocysteine in the presence or absence of an H₂S donor (NaHS) in increasing doses. The study showed that homocysteine in combination with small concentrations of H₂S reduced the production of H₂O₂, ONOO^{•-} and O₂^{•-}, and cell damage. The results of our study show that after the co-administration of DL-Hcy and DL-PAG, the levels of NO₂⁻ and O₂^{•-} statistically significantly decreased, while TBARS and H₂O₂ increased (Table IIc), so we cannot find a dominant trend in the dynamics of these parameters. In the case of the basic forms of homocysteine, H₂S does not yet have a precise pro- or anti-oxidative potential. However, in contrast to other studies (44), we examined the effects of endogenous H₂S.

Conclusion

The acute and direct effects of homocysteine on the isolated rat heart imply negative inotropic and lusitropic effects and a vasoconstrictor influence on the coronary endothelium. The inhibition of H₂S synthesis led to complete depression of cardiodynamics, which showed that these gasotransmitters can play a role in the contraction of cardiomyocytes and the control of coronary vascular tonus. Homocysteine and H₂S did not contribute to the development of oxidative stress in this study, which means that acute cardiodynamic effects are possibly not mediated by oxidative damage. It also seems that H₂S has certain anti-oxidative capacities, which should be definitely confirmed in further studies. It seems that homocysteine shows a lower pro-oxidative effect in the presence of H₂S in comparison to the absence of production of this signal molecule, which indicates a potential anti-oxidative capacity of the mentioned gasotransmitter.

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