Protective role of hydrogen-rich water on aspirin-induced gastric mucosal damage in rats

Jing-Yao Zhang, Qi-Fei Wu, Yong Wan, Si-Dong Song, Jia Xu, Xin-Sen Xu, Hu-Lin Chang, Ming-Hui Tai, Ya-Feng Dong, Chang Liu

Abstract

AIM: To investigate the role of the hydrogen-rich water (HRW) in the prevention of aspirin-induced gastric mucosal injury in rats.

METHODS: Forty male rats were allocated into four groups: normal control group, HRW group, aspirin group, and HRW plus aspirin group. The protective efficacy was tested by determining the gastric mucosal damage score. Malondialdehyde (MDA), superoxide dismutase (SOD), myeloperoxidase (MPO), interleukin (IL)-06 and tumor necrosis factor (TNF)-α in gastric tissues were evaluated. The serum levels of IL-1β and TNF-α were also detected. Histopathology of gastric tissues and localization of Cyclooxygenase 2 (COX-2) were detected using hematoxylin and eosin staining and immunohistochemistry, respectively.

RESULTS: Pretreatment with HRW obviously reduced aspirin-induced gastric damage scores (4.04 ± 0.492 vs 2.10 ± 0.437, P < 0.05). The oxidative stress levels of MDA and MPO in the gastric tissues increased significantly in the aspirin-treated group compared with the HRW group (2.43 ± 0.145 vs 1.79 ± 0.116 nmol/mg prot, P < 0.05 and 2.53 ± 0.238 vs 1.40 ± 0.208 U/g tissue, P < 0.05, respectively). HRW could obviously elevated the SOD levels in the gastric tissues (37.94 ± 8.44 vs 59.55 ± 9.02 nmol/mg prot, P < 0.05). Pretreatment with HRW significantly reduced IL-06 and TNF-α in the gastric tissues (46.65 ± 5.50 vs 32.15 ± 4.83 pg/mg, P < 0.05 and 1305.08 ± 101.23 vs 855.96 ± 93.22 pg/mg, P < 0.05), and IL-1β and TNF-α in the serum (505.38 ± 32.97 vs 343.37 ± 25.09 pg/mL, P < 0.05 and 264.53 ± 28.63 vs 114.96 ± 21.79 pg/mL, P < 0.05) compared to treatment with aspirin alone. HRW could significantly decrease the COX-2 expression in the gastric tissues (staining score: 8.4 ± 2.1 vs 2.9 ± 1.5, P < 0.05).

CONCLUSION: HRW pretreatment alleviated the aspirin-induced gastric lesions by inhibiting the oxidative stress, inflammatory reaction and reducing the COX-2 in the gastric tissues.

Key words: Hydrogen; Aspirin; Gastric lesion; Oxidative stress; Cytokines; Cyclooxygenase 2

Core tip: Aspirin is one of the most widely used medicines, but can cause adverse side effects of gastric tissues.
Hydrogen has been found to have powerful antioxidant and anti-inflammatory effects. We investigated the protective role of hydrogen-rich water on aspirin-induced gastric mucosal damage in rats. We found that hydrogen could alleviate the aspirin-induced gastric lesions by inhibiting the oxidative stress, inflammatory reaction and reducing the Cyclooxygenase 2 in the gastric tissues. This may provide a potential therapy for the adverse effects of aspirin.


INTRODUCTION

Non-steroidal anti-inflammatory drugs (NSAIDs), including the anti-inflammatory and analgesic agents, are the most common prescription medicines for the treatment of many diseases[1]. Aspirin, in particular, is widely used in cardiovascular disorders and even in the treatment of cancer because of its multiple actions[11,12]. However, there are adverse effects even it is used at a low and safe dose, especially the side effects in the digestive tract, such as the dyspeptic symptoms, gastrointestinal erosions, peptic ulcers, overt bleeding or perforation[13,14]. Hence, attempts have been made to develop a risk-free dose, or coated and buffered aspirin to mitigate the injury. However, little progress has been made[15].

Research has revealed that various factors, such as endogenous prostaglandin (PG), neutrophil-dependent microvascular injury, oxygen-derived free radicals, inflammatory cytokines[16-18], are associated with the aspirin-induced gastric mucosal damage. Tumor necrosis factor (TNF)-α is a proinflammatory cytokine and can augment the neutrophil-derived superoxide generation, leading to oxygen radical-mediated tissue injury. Pretreatment of TNF-α inhibitors have been reported to suppress the gastric mucosal injury[19,20]. Moreover, the tissue enzymatic activity, such as dismutase (SOD), myeloperoxidase (MPO) and malondialdehyde (MDA), are also considered to be related to aspirin-induced gastric mucosal injury[21,22]. Among the multiple mechanisms, the function of cyclooxygenase (COX) is one of the biggest concerns for the scientists. COX-1 plays a definite protective role in the NSAIDs-induced gastric injury[23]. Nevertheless, the COX-2 plays a perplexing role in this pathological process. First, COX-2 can be induced by the damaging agents such as luminal acid and aspirin in the gastric mucosa, and inhibition of COX-2 can alleviate the damage[24,25]. It involves the maintenance of gastric mucosal integrity by preventing exogenous injury and by promoting gastric mucosal healing[26]. However, there are also studies reporting that COX-2 inhibitors exacerbate gut injury and attenuate the tissue’s ability to respond to mild damaging agents[27,28]. The possible mechanism is that COX-2 expression may be a compensatory response to increase the levels of gastroprotective PG in the process of gastric injury[29].

Hydrogen therapy is a new method which has gained much appreciation recently and shown efficacy in many diseases. Hydrogen has anti-oxidant, anti-inflammatory, anti-apoptotic, anti-allergy, and anti-cancer effects. Several methods, including inhalation, drinking hydrogen-rich water (HRW) and injection of hydrogen-saturated saline, have been developed and proved to be valid and reliable[30,31]. Oral intake of HRW is an effective and convenient way to deliver the molecular hydrogen, which is more suitable for application. Some researches showed that oral intake of HRW could protect cardiac allografts from inflammation-associated deterioration, kidney allografts from chronic rejection and so on[32-34]. Although the research of hydrogen as a medical therapy has been extensively investigated, its effect on the aspirin-induced gastric mucosal injury has not been reported.

The aim of this study was to assess the protective role of HRW on aspirin-induced gastric mucosal damage in rats mainly through measurement of oxidative stress indicators and cytokines, including levels of MDA, SOD, MPO, interleukin (IL)-6 and TNF-α in the gastric tissues, IL-1β and TNF-α in the serum, and expression of COX-2 in gastric mucosa.

MATERIALS AND METHODS

Experimental animals and preparation of hydrogen-rich water

The study was conducted using male Sprague Dawley rats (200-250 g) (Animal Feeding Center of Xi’an Jiaotong University Health Science Center, Xi’an, China). All rats were housed (5 per cage) in clear, pathogen-free polycarbonate cages in the animal care facility, and were fed a standard animal diet and water ad libitum under controlled temperature with 12-h light-dark cycles. They were cared in accordance with the guidelines of the Ethical Committee, Xi’an Jiaotong University Health Science Center. The HRW was produced by Naturally Plus Japan International Co, Ltd (Japan), which was stored under atmospheric pressure at 4 ℃ in an aluminum bag with no dead volume. The gas chromatography was used to confirm the content of hydrogen by the method described by Oh sawa et al[35]. The hydrogen concentration of the HRW we used in this study was 0.63-0.82 mmol/L.

Study design

Forty rats were divided into four groups randomly, each consisting of 10 animals, with different pre-treatments for 7 d, as follows: (1) normal control group received oral saline; (2) HRW control group received oral HRW (changed every 3 h, at about 80 mL/d); (3) aspirin group took oral saline; and (4) HRW plus aspirin group received HRW (80 mL/d per rat). On the 8th day, gastric injury was induced by administration of aspirin (400 mg/kg).
Extensive erosions over entire mucosa
Slight edema and congestion
Edema, congestion, and bleeding
Many small and a few large erosions

The supernatant was used for the determination of cytokines. The levels of cytokines (IL-6 and TNF-α) in the gastric tissue samples and IL-1β and TNF-α in the blood samples were evaluated using the ELISA kit reagent from Dakewe Biotech Co. (Shenzhen, China) according to the manufacturer’s instructions. A standard curve was run on each assay plate using recombinants of the respective cytokines in serial dilutions.

Immunohistochemistry
To establish the immunolocalization of COX-2, a mouse polyclonal antibody (Beijing Biosynthesis Biotechnology Co., Ltd, Beijing, China), was used at a working dilution of 1:20. The antibody was applied directly to sections, and slides were incubated overnight at 4 °C in a humidified chamber. Immune complexes were subsequently treated with the secondary antibody (containing anti-rabbit and anti-mouse immunoglobulins) and detected after streptavidin peroxidase treatment for 20 min at room temperature. After rinsing sections with three changes of PBS, immunoreactivity was visualized with diaminobenzidine (DAB)-hydrogen peroxide (20 min). Sections were gently rinsed in distilled water, counterstained with HE, and photomicrographs were taken under a microscope (Olympus Optical Co., Tokyo, Japan).

RESULTS

Histopathological changes
Histopathological examination revealed aspirin-induced severe congestion and multiple hemorrhagic erosions in the stomach tissue, particularly in mucus-secreting cells, characterized by gastric pit damage and vacuolization of the glandular portion. Pretreatment with HRW considerably attenuated, but did not completely prevent the severity of these histopathological changes, while some erosion in sub-glandular and epithelial necks was evident (Figure 1A). In the aspirin group, the mean gastric mucosal damage score was 4.04 ± 0.492, while HRT pretreat-
treatment with HRW could also significantly decrease the elevation of IL-6 and TNF-α in the gastric tissue (46.65 ± 5.50 vs 32.15 ± 4.83 pg/mg, P < 0.05 and 1305.08 ± 101.23 vs 855.96 ± 93.22 pg/mg, P < 0.05), (Figure 2).

Effects of HRW on cytokine levels in the peripheral blood

The levels of IL-1β and TNF-α in the peripheral blood were markedly increased in the aspirin-treated group compared with the normal control and HRW control groups (P < 0.01). The increase in IL-1β and TNF-α concentration in the gastric mucosa elicited by aspirin were significantly suppressed by HRW pretreatment with HRW could also significantly decrease the elevation of IL-6 and TNF-α in the gastric tissue (46.65 ± 5.50 vs 32.15 ± 4.83 pg/mg, P < 0.05 and 1305.08 ± 101.23 vs 855.96 ± 93.22 pg/mg, P < 0.05), (Figure 2).

**Figure 1** Histopathological examination of stomach sections. A: Hematoxylin-eosin stained results showed severe degenerative changes in glandular region, epithelial folds and connective septa in aspirin-induced mucosal tissue (1, × 100 and 2, × 200), while hydrogen-rich water (HRW) pretreatment displayed slight changes (3, × 100 and 4, × 200); B: The gastric mucosal damage score in four groups. The mean scores are significantly higher in the aspirin group and HRW plus aspirin group (HRW + Asp) when compared with the normal control group and HRW alone group (P < 0.01). Pretreatment with HRW could significantly decrease the damage score in HRW + aspirin group (Asp) group when compared with aspirin group (P < 0.05).
Immunohistochemical analysis of COX-2 in gastric mucosal tissues

Immunohistochemical analysis was performed to ascertain the localization of COX-2 in gastric mucosal tissues. Rats treated with aspirin displayed significant immunoreactivity for COX-2 around the glandular regions, mucoid cells and neck gland cells of gastric tissues (staining score: 8.4 ± 2.1). In contrast, the group of rats pretreated with HRW presented lower COX-2 immunoreactivities (staining score: 2.9 ± 1.5) (P < 0.05) (Figure 4).

DISCUSSION

Aspirin is widely used as an anti-inflammatory and analgesic drug, but it often induces gastrointestinal adverse effects, which limits its clinical use. Studies focusing on...
reduction of the aspirin-induced gastric mucosal damages is urgently needed. Here, we examined the effect of HRW on the aspirin-induced gastric mucosal injury. Although hydrogen is the lightest gas in nature, it has an enormous anti-oxidation and anti-inflammatory capacity. In fact, there are few studies investigating the preventive or therapeutic effects of gas on gastric mucosal damages. In one study, hydrogen sulfide was found to protect against aspirin-induced gastric injury via reducing oxidative stress. In another study, Liu et al. found that hydrogen treatment successfully ameliorated stress-associated gastric ulceration via its anti-oxidant, anti-inflammatory and anti-apoptotic effects. Our research shows that hydrogen can reduce aspirin-induced gastric injury by reducing MDA, MPO, IL-06 and TNF-α levels and increasing SOD activity in gastric tissues. Simultaneously, hydrogen could decrease the elevation of IL-1β and TNF-α in the serum compared to treatment with aspirin alone. We also found that hydrogen can decrease COX-2 expression in the gastric tissues, which may be the key mechanism of hydrogen action because of its importance in the aspirin-induced gastric injury. The present study demonstrated that hydrogen may prevent against aspirin-induced gastric mucosal injury, mainly depending on the modulation of anti-inflammatory cytokines, anti-oxidative stress and activation of COX-2.

In the past few years, the research of hydrogen therapy attracted wide attention from scientists and physicians. Hydrogen molecules have been proven to act as an important physiological regulatory factor to cells.
and organs by anti-oxidant, anti-inflammatory, anti-apoptotic and other protective effects, which can be applied in the treatment of various diseases\textsuperscript{[24]}. Compared with other studies on hydrogen therapy, a distinctive experimental design in our research was the delivery of hydrogen molecule. We designed and used the random oral intake of HRW to deliver the hydrogen molecule, which was similar to the human physiological status. This special and convenient delivery method implies that the HRW can be used as a kind of drink, which can immensely expand its applications.

According to a previous study, redox imbalance plays a major pathogenic role in aspirin gastropathy\textsuperscript{[13]}. The over oxidative stress in the gastric mucosa can increase the production of oxygen radicals or decrease the capability of antioxidant defenses\textsuperscript{[16]}. We found in the present study that hydrogen can significantly reduce the Aspirin-induced elevation of MDA and MPO, the most typical markers of free radical species-related injury. In addition, hydrogen also reversed the aspirin-induced elevation of SOD, which is responsible for converting superoxide radicals to molecular oxygen and hydrogen peroxide within cytoplasm and mitochondria\textsuperscript{[30]}. The cytokines, such as TNF-\(\alpha\), IL-1\(\beta\) and IL-06, induced by lymphocytes and macrophages that infiltrate the gastric mucosa, are associated with the tissue injury\textsuperscript{[40]}. They are responsible for the neutrophil adherence in the microcirculation of gastric mucosa, and the release from activated macrophages with parallel accumulation of neutrophils within the gastric pits\textsuperscript{[41]}. In the present study, we observed that pretreatment with hydrogen resulted in significantly decreased TNF-\(\alpha\), IL-1\(\beta\) and IL-06 production which could prevent subsequent neutrophils infiltration and alleviate injury.

On a deeper level, we focused on COX-2, which was the rate-limiting enzyme to regulate the synthesis of PG. This is a key factor in response to stress. It was reported that aspirin can rapidly up-regulate COX-2 mRNA expression in rat gastric mucosa, probably as a compensatory response to inhibition of COX-2 activity and gastric PG synthesis\textsuperscript{[42,43]}. COX-2 products play a relevant role in the maintenance of gastric mucosal integrity by preventing exogenous injury to the stomach and accelerating gastric mucosal healing\textsuperscript{[44]}. Interestingly, HRW pre-treatment counteracted the increased expression of COX-2 induced by aspirin treatment, which promote mucosal healing. According to the description in the introduction section that the COX-2 plays a controversial role in the aspirin-induced gastric injury, we speculate that the COX-2 can act as the double-edged sword. First, COX-2 is induced by cytokines and then promotes the release of cytokines to exacerbate inflammation\textsuperscript{[28]}; meanwhile, as a compensatory mechanism, the expression of the COX-2 can accelerate the prostaglandin production to protect the gastric mucosa\textsuperscript{[39]}. Thus, when the noxious stimuli are strong enough to cover the protective role of the COX-2, high expression of COX-2 is a harmful indicator in the pathological process of aspirin-induced gastric injury. In this study, a high dosage of 400 mg/kg of aspirin was used for the rats and the high expression of COX-2 was induced by the cytokines. The hydrogen can suppress its expression to protect the gastric mucosa. Although the different molecular mechanism between hydrogen and COX-2 is not clear, COX-2 as a definite target of hydrogen can be confirmed. A clue of “hydrogen → mitigating oxidative stress → alleviating inflammatory reaction → suppressing COX-2 expression → protecting aspirin-induced gastric injury” can be used to summarize the research.

In conclusion, the present study shows that: (1) hydrogen is able to prevent aspirin-induced injury to the rat gastric mucosa through a mechanism, which is in part contributed by the anti-oxidant and anti-inflammatory activities; (2) aspirin causes up-regulation of COX-2 expression in the gastric mucosa; and (3) hydrogen significantly counteracts aspirin-induced up-regulation of COX-2 expression, probably as a consequence of the reduction in the extent of aspirin-induced injury. Therefore, hydrogen therapy might be safe and effective in the prevention of the stomach injury due to NSAIDs. Future clinical studies are necessary to determine whether it can be applied as preventive or therapeutic agents as shown in this preclinical study.

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**COMMENTS**

**Background**

Aspirin is widely used as an anti-inflammatory and analgesic drug, but it often induces gastrointestinal adverse effects which include dyspeptic symptoms, gastrointestinal erosions, peptic ulcers, overt bleeding or perforation. Hence, efforts have been made to develop the risk-free dose or coated and buffered aspirin to mitigate the injury, but little progress has been made. This limits the clinical use of aspirin, and studies aimed to reduce the aspirin-induced gastric mucosal damages are urgently needed.

**Research frontiers**

Hydrogen therapy is a new medical method which has gained much attention recently. It has been shown that hydrogen has anti-oxidant, anti-inflammatory, anti-apoptotic, anti-allergy, and anti-cancer effects. Several methods used to deliver hydrogen, including inhalation, drinking hydrogen-rich water (HRW) and injection with hydrogen-saturated saline, have been proved to be valid and reliable.

**Innovations and breakthroughs**

Compared with other researches on hydrogen therapy, this study is characterized by a distinctive experimental design in the delivery of hydrogen molecule. The authors used the random oral intake of HRW to deliver hydrogen molecule, which was similar to human physiological status. This special and convenient delivery method implies that the HRW can be used as a kind of drink, which can immensely expand its applications. The authors also found that the hydrogen did have a protective role in the aspirin-induced gastric injury, which is in part contributed by the anti-oxidant and anti-inflammation activities. Hydrogen can significantly counteract aspirin-induced up-regulation of Cyclooxygenase 2 (COX-2) expression, probably as a consequence of the reduction in the extent of aspirin-induced injury.

**Applications**

Hydrogen therapy is safe and effective in prevention of the stomach injury derived from non-steroidal anti-inflammatory drugs administration. The special and convenient delivery method used in this study implies that HRW can be
Hydrogen is the lightest gas in the nature, but it has huge anti-oxidant, anti-inflammatory, anti-apoptotic, anti-allergy, and anti-cancer effects. HRW is produced by pressing the hydrogen gas into the water by a specific device under high pressure.

**Peer review**

This study demonstrates the protective effects of hydrogen therapy (hydrogen rich water) on acute aspirin gastropathy in rats, which are probably dependent on the modulation of COX-2, cytokines and antioxidant activities. This is an interesting paper, with a new clinical application of a new therapy, in an experimental model of gastropathy. The results are well presented and provide sufficient experimental evidence to support the conclusions.

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**Zhang JY et al. Hydrogen protects against aspirin-induced gastric injury**

**Terminology**

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