Consumption of vitamin B6 reduces colonic damage and protein expression of HSP70 and HO-1, the anti-tumor targets, in rats exposed to 1,2-dimethylhydrazine

TOMOKO KAYASHIMA1,4, KENTA TANAKA2, YUKAKO OKAZAKI3, KIMINORI MATSUBARA1, NORIYUKI YANAKA2 and NORIHISA KATO2

1Graduate School of Education, Hiroshima University, Higashi-Hiroshima 739-8524; 2Graduate School of Biosphere Science, Hiroshima University, Higashi-Hiroshima 739-8528; 3Faculty of Human Life Sciences, Fuji Women’s University, Ishikari 061-3204, Japan

Received April 29, 2011; Accepted July 15, 2011

DOI: 10.3892/ol.2011.370

Abstract. Mounting evidence indicates that vitamin B6 is a protective factor for colon cancer. Elevations in colonic damage, cell proliferation and heat shock proteins (HSPs, molecular chaperones) have been suggested to be associated with colon carcinogenesis. This study was performed to examine the effect of dietary levels of vitamin B6 (1, 7 or 35 mg pyridoxine HCl/kg diet) for 22 weeks on colonic damage, epithelial cell proliferation and expression of HSPs in rats exposed to 1,2-dimethylhydrazine (DMH). Supplemental vitamin B6 with a low vitamin B6 diet (1 mg pyridoxine HCl/kg diet) significantly reduced fecal activity of intestinal alkaline phosphatase (an index of intestinal damage) and the colonic epithelium PCNA labeling index (a marker of cell proliferation). Analysis using ELISA indicated that supplemental vitamin B6 significantly lowered protein levels of colonic HSP70 and heme oxygenase-1, HSP32 (HO-1). However, real-time RT-PCR analysis revealed that the mRNA levels of these HSPs were not decreased by supplemental vitamin B6, suggesting that the lowering effect of vitamin B6 on the colon protein expression of the HSPs is mediated by mechanisms not involving altered gene expression. This study provided evidence that dietary supplemental vitamin B6 suppresses colon damage, epithelial cell proliferation and protein expression of HSP70 and HO-1, the targets for anti-tumor agents, in rats exposed to DMH.

Introduction

In previous studies, we provided the first evidence for the anti-colon tumor effect of dietary vitamin B6 in azoxymethane (AOM)-treated mice (1,2). In accordance with this finding, accumulating epidemiological evidence indicates an inverse association between vitamin B6 status and colon cancer (3-5). It has been suggested that the anti-tumor effect of vitamin B6 is mediated by lowering cell proliferation, inflammation, angiogenesis, oxidative stress and DNA damage (6-10). However, the underlying mechanism has yet to be clarified.

Colon damage and loss of epithelial cells was considered to be a risk factor for colon carcinogenesis (11,12). Epidemiological studies indicated that the association between vitamin B6 status and colon cancer is particularly significant in males consuming alcohol (3). Consumption of alcohol has been reported to cause colonic epithelium damage, which has been considered to promote colon carcinogenesis (13). Evidence suggests that vitamin B6 has a protective role against cytotoxicity in cell culture studies (14,15), although the mechanism remains unclear. Thus, the protective effect of vitamin B6 on colonic damage caused by a carcinogen should be determined.

In general, tissue damage caused by stress exposure (heat shock, oxidative stress and hypoxia) is associated with a higher expression of heat shock proteins (HSPs, molecular chaperones) (16-18). HSPs are overexpressed in a variety of human cancers, and are involved in tumor cell proliferation, differentiation, invasion, metastasis, death and recognition by the immune system (16-18). Conversely, HSPs are well known for their cytoprotective factor in response to cell injury. Accordingly, it is of interest to examine whether dietary vitamin B6 modulates the expression of colonic HSPs. The present study was performed to examine the effect of dietary vitamin B6 on colonic damage and expression of HSPs in 1,2-dimethylhydrazine (DMH)-treated rats.

Materials and methods

Animals. Male Sprague Dawley rats (4 weeks of age) were purchased from the Hiroshima Laboratory Animal Center.
Materials. The activity of intestinal-derived alkaline phosphatase in feces was determined according to the method described elsewhere (20). p-Nitrophenyl phosphate was used as the substrate, and the absorbance of the reaction product p-nitrophenol was determined spectrophotometrically at 405 nm. Intestinal alkaline phosphatase activity was inhibited using 60 mM L-phenylalanine, which acts as a specific non-competitive inhibitor of the intestinal isozyme. The difference between total activity (non-inhibited) and the activity following inhibition with L-phenylalanine is the activity of the intestinal isozyme. One unit of the activity was defined as μmol p-nitrophenol/min, and the fecal activity was expressed as units/3 days of feces collected. The colonic epithelium PCNA labeling index was determined according to the method described elsewhere (21).

Protein and mRNA analyses. Colon protein levels of HSP70 and heme oxygenase-1 (HO-1) were determined by ELISA kits (Stressgen, MI, USA). Protein concentration was determined by the Bradford method with a standard of bovine serum albumin (22). HSP70, HO-1 and β-actin gene primers were purchased from Qiagen (QT00187397, QT00175994 and QT00193473, respectively; Helden, Germany). A single melt curve was observed for each primer set in all real-time RT-PCR reactions. Real-time RT-PCR was conducted using an iQ5 real-time PCR detection system (Bio-Rad Laboratories, CA, USA). Gene expressions were quantitated using Bio-Rad iQ5 optical system software (Bio-Rad Laboratories).

Statistical analysis. Statistical analysis was conducted by one-way analysis of variation (ANOVA) and Scheffe’s multiple range test (Excel Statistics 2006 for Windows, Social Survey Research Information Co., Ltd., Tokyo, Japan). P<0.05 was considered to be statistically significant.

Results

Growth was unaffected by dietary treatment (data not shown). Food intake (for 3 days) at 8, 15 and 22 weeks was also not affected by the dietary level of vitamin B6. Two rats in the 1 mg vitamin B6/kg group had an adenoma in the colon, and were not used for the further analysis of colon parameters. Fecal activity of intestinal alkaline phosphatase was significantly lower in the 7 and 35 mg PN HCl/kg diet groups (-58 and -64%, respectively) as compared to the 1 mg PN HCl/kg group (P<0.05, Fig. 1A). The activity of intestinal microbial-derived alkaline phosphatase was unaffected by dietary vitamin B6 (data not shown). The PCNA labeling index of the colonic epithelium was also significantly lower in the 7 and 35 mg PN HCl/kg diet groups (-38 and -42%, respectively) as compared to the 1 mg PN HCl/kg group (P<0.05, Fig. 1B).

Dietary supplemental vitamin B6 appeared to decrease colonic HSP70 protein in a dose-dependent manner (Fig. 2A). Compared to the 1 mg PN HCl/kg diet, the colonic level of HSP70 protein was significantly reduced in the 7 and 35 mg PN HCl/kg diet group (-49%) (P<0.05, Fig. 2A). Colonic levels of HO-1 protein were significantly lower in the 7 and 35 mg PN HCl/kg diet groups (-32 and -37%, respectively) compared to the 1 mg PN HCl/kg diet group (P<0.05, Fig. 2B). Colonic mRNA levels of HSP70 were unaffected by dietary levels of vitamin B6 (Fig. 3A). Colonic levels of HO-1 mRNA were significantly elevated in the 7 and 35 mg PN HCl/kg diet groups (+48 and +62%, respectively) compared to the 1 mg PN HCl/kg diet group (P<0.05, Fig. 3B).

Discussion

This study has shown that dietary supplemental vitamin B6 suppressed DMH-induced colon damage as evidenced by...
the data pertaining to fecal alkaline phosphatase activity (Fig. 1). Our previous study indicated that dietary vitamin B6 lowered colonic levels of oxidative stress markers, including 8-hydroxyguanosine and 4-hydroxynonenal, and inflammatory mediators, such as COX-2 and iNOS in AOM-treated mice (10). Therefore, the protective effect of vitamin B6 against colon damage appears to be, at least in part, explained by the lower oxidative stress and inflammation. Further study is in progress to fully elucidate the molecular basis of the underlying mechanisms of the protective role of vitamin B6.

The cytotoxicity of colonic components, including secondary bile acids and oxygen radicals, may cause epithelial cell loss in the large bowel, leading to a compensatory crypt cell proliferation and the development of colon cancer (13,14). In the DMH-treated rats, supplemental vitamin B6 caused a significant reduction in the PCNA labeling index, an index of cell proliferation as well as fecal intestinal phosphatase activity (Fig. 1). This finding suggests that the reduced intestinal damage by supplemental vitamin B6 decreases cell proliferation in the colonic epithelium. Previously, we reported that dietary supplemental vitamin B6 caused reduced colonic epithelial cell proliferation despite no alteration in apoptosis and the total cell number in the epithelium of AOM-treated mice (1). The reduced colonic cell proliferation by dietary vitamin B6 may be counteracted by the reduced loss of the epithelial cells.

Notably, this study has demonstrated that dietary supplemental vitamin B6 reduced the levels of colonic HSP70 and HO-1 proteins in DMH-treated rats (Fig. 2). Since tissue damage by stress exposure induces a higher expression of HSPs, reduced colon damage may result in a lower protein expression of HSP70 and HO-1. HSPs are considered to be novel targets for anti-tumor agents (16-18). HSP70 promotes cell survival under a wide range of otherwise lethal stimuli. HSP70 neutralization has been found to exert potent anti-tumor effects in animal models of colon cancer (23). It has been suggested that HO-1 protects tumor cells against oxidative stress and is regarded as an enzyme facilitating tumor progression (18,19). It has been demonstrated that pegylated zinc protoporphyrin, a potent HO inhibitor, administered in vitro induced apoptosis of human colon carcinoma SW480 cells and inhibited the growth of murine colon carcinoma in vivo (24). Thus, the results of the present study suggest that the anti-colon tumor effect of vitamin B6 is, at least in part, mediated by reducing the protein expression of HSP70 and HO-1. To the best of our knowledge, this is the first study to demonstrate the possible involvement of HSPs in the mechanism of anti-tumorigenesis by dietary nutrients. The expression of proinflammatory cytokines including TNF-α, IL-1α and IL-6 was up-regulated, and NF-kB was activated in monocytes by HSP70 (25). Our previous study indicated that dietary vitamin B6 decreased the colonic expression of COX-2 and iNOS in AOM-treated mice (10). Thus, it is of interest to test whether a reduced HSP70 protein expression by dietary vitamin B6 results in lower inflammatory mediators.

In the present study, colonic HSP70 and HO-1 mRNA levels were not reduced by dietary vitamin B6 (Fig. 3). Thus, the lower protein expression of HSP70 and HO-1 by dietary B6
appears not to be mediated by lowering the mRNA expression. These results are expected since a previous study has shown that HSP70 protein expression is regulated by the degradation of HSP70 proteins as well as transcriptional regulation (26). Further studies are required to examine the underlying mechanisms of a reduced protein expression of the HSPs by vitamin B6. In our preliminary study, no effect of dietary vitamin B6 was observed on the colonic protein expression of HSP90, HSP60 and HSP25 (Kayashima et al; unpublished data). At present, it is not known why the protein expression of HSP70 and HO-1 was specifically affected by dietary vitamin B6.

HSP70 has been suggested to be a cytoprotective factor by modulating protein unfolding and protein degradation (16). HO-1 has also been suggested to prevent cell damage and exhibit marked antioxidant activity (17). Thus, prior to this study, we considered the possibility that dietary vitamin B6 causes a higher expression of HSPs, leading to the protective function of vitamin B6 against colon damage by DMH. However, this possibility appears to be negated.

In conclusion, the present study provided the first evidence that dietary vitamin B6 lowers colonic damage and protein expression of HSP70 and HO-1 in rats exposed to DMH. These findings suggest novel mechanisms of an anti-colon tumor effect of dietary vitamin B6.

Acknowledgements

This study was supported in part by a Grant-in-Aid for Scientific Research from the Ministry of Education, Culture, Sports, Science and Technology of Japan.

References