Cocoa-rich diet prevents azoxymethane-induced colonic preneoplastic lesions in rats by restraining oxidative stress and cell proliferation and inducing apoptosis

Ildefonso Rodríguez-Ramiro¹, Sonia Ramos¹, Elvira López-Oliva², Angel Agis-Torres², Miren Gómez-Juaristi¹, Raquel Mateos¹, Laura Bravo¹, Luis Goya¹ and María Ángeles Martín¹,³

¹Instituto de Ciencia y Tecnología de Alimentos y Nutrición (ICTAN-CSIC), Madrid, Spain
²Facultad de Farmacia, Universidad Complutense de Madrid, Madrid, Spain
³CIBER de Diabetes y Enfermedades Metabólicas Asociadas (CIBERDEM), ISCIII, Madrid, Spain

Cocoa is a rich source of bioactive compounds with potential chemopreventive ability but up to date its effectiveness in animal models of colon carcinogenesis has not been addressed. Herein, we investigated the in vivo effect of a cocoa-rich diet in the prevention of azoxymethane (AOM)-induced colon cancer and the mechanisms involved. Our results showed that cocoa feeding significantly reduced AOM-induced colonic aberrant crypt foci formation and crypt multiplicity. Oxidative imbalance in colon tissues seems to be prevented by cocoa as indicated by reduced oxidation markers levels and increased enzymatic and non-enzymatic endogenous defences. Cocoa-rich diet also exhibited antiproliferative effects by decreasing the levels of extracellular regulated kinases, protein kinase B and cyclin D1 together with pro-apoptotic effects evidenced by reduced Bcl-xL levels and increased Bax levels and caspase-3 activity. Our findings provide the first in vivo evidence that a cocoa-rich diet may inhibit the early stage of colon carcinogenesis probably by preventing oxidative stress and cell proliferation and by inducing apoptosis.

Keywords:
Bioactive compounds / Cocoa flavanols / Colorectal cancer / Glutathione enzymes / Oxidative stress

Numerous epidemiological studies suggest that diets rich in natural antioxidants are associated with reduced risk of certain cancers, notably colorectal cancer (CRC) [1]. Cancer is a multistage process conventionally defined by three stages: initiation, promotion and progression. Along this process, oxidative stress has the potential to affect a large array of carcinogenic pathways involved in proliferation of initiated cells and enhanced malignant transformation [2]. Therefore, the suppression of oxidative stress by natural antioxidant compounds seems to be an effective approach in preventing the initiation and progression of CRC.

Cocoa and its phenolic compounds, the flavanols epicatechin, catechin and procyanidins, have recently attracted a great deal of interest because of their potential ability to act as highly effective chemopreventive agents. As antioxidants, cocoa flavanols have been shown to protect cell constituents, limiting the risk factor for cancer and other chronic diseases [3, 4]. In addition, they can display other chemopreventive properties which may be independent of conventional antioxidant activities [5]. In particular, the recent in vitro studies have demonstrated that cocoa phenolic compounds can regulate several signal transduction pathways [6, 7], activate redox-sensitive transcription factors [8] and modulate expression of specific genes involved in cell survival and cell...
death [9]. Moreover, as compared with other flavonoid-containing foods, cocoa products exhibit a high concentration of procyanidins that are poorly absorbed in the intestine and consequently its beneficial effects would be more focused on the gastrointestinal tract where they may have an important local function neutralising oxidants. Despite these evidences, the efficacy of cocoa against CRC initiation and development in vivo remains largely unexamined. Therefore, in the present study we have used the well-defined azoxymethane (AOM)-induced colon cancer model in rats to investigate the effectiveness of a cocoa-enriched diet in preventing the early phase of colon carcinogenesis and the mechanisms of action involved.

Natural Forastero cocoa powder (a kind gift from Nutrexpa, Barcelona, Spain) was used for this study. A detailed description of this cocoa is given elsewhere [10]. Diets were prepared from an AIN-93G formulation (Panlab S. L., Barcelona, Spain). The 12% cocoa diet was produced by adding 120 g/kg cocoa to AIN-93G. This powdered supplement was formulated to provide 1 g of polyphenols/kg of diet. A similar cocoa supplementation protected liver cells against oxidative stress by activating the antioxidant defence system [11]. The composition of the diets is given in Supporting Information Table 1. Rats were fed with control or cocoa-enriched diets during 8 wk and injected with saline or AOM (20 mg/kg body weight) during the second and third weeks (see experimental protocol in Supporting Information Fig. 1). At the end of the experiment, colon samples were evaluated for aberrant crypt foci (ACF) formation, and markers of oxidative stress, proliferation and apoptosis (Supporting Information, Material and Methods).

All animals had a steady body weight gain during the treatment and the administration of AOM did not affect the growth of the rats (Supporting Information Table 2). Interestingly, the body weight of animals fed with cocoa diet was slightly but significantly reduced as compared with control groups (about 10%). As described elsewhere [12, 13], this effect seems to be attributed to the cocoa polyphenolic fraction and its ability to reduce fat adipose tissue. Animals were treated according to the Institutional Care Instructions (Bioethical Commission from Consejo Superior de Investigaciones Científicas, CSIC).

Administration of the colon-specific carcinogen AOM to rodents induces the development of ACF, preneoplastic lesions that may progress into cancer later on [14]. Therefore, we first investigated the efficacy of cocoa-enriched diet on inhibiting AOM-induced ACF formation. We found that all the rats injected with AOM developed aberrant crypts; however, the cocoa-enriched diet significantly reduced the AOM-induced ACF formation (Supporting Information Fig. 2). More importantly, the number of ACF with a crypt multiplicity of four or more, that have been suggested to represent a higher risk for malignant tumour progression [14], was largely reduced by cocoa feeding (Fig. 1A).

Experimental studies with rodents have demonstrated that AOM can induce ACF formation in colon by causing oxidative stress leading to DNA damage and mutations in cancer related genes [15–17]. Owing to the reported antioxidant effects of cocoa and their flavonoids, oxidative stress biomarkers in the distal colon of rats were evaluated. As shown in Fig. 1B, the levels of carbonyl groups (marker of protein oxidation) and malondialdehyde (MDA), a lipid

| Figure 1. Number of crypt multiplicity of ACF (A), MDA and carbonyl levels (B), enzyme activity of GPx, GR and GST (C) and GSH levels (D) in colon tissues from rats injected with saline or AOM and fed with control or cocoa-enriched diet. The data represent mean ± SD value from 10 to 12 rats in each group. Results of crypt multiplicity were statistically analyzed with Student’s t test (*p<0.05; **p<0.01; compared with the AOM control group). In the rest of comparisons, one-way ANOVA was used. Means without a common letter differ significantly, p< 0.05. |
peroxidation end product, were significantly higher in the distal colon of control AOM-treated rats as compared with the untreated control group whereas both increases were prevented in the AOM-treated rats fed with cocoa. These results suggest that the antioxidant properties of cocoa could prevent the colonic oxidative damage induced by AOM. Supporting this, different natural antioxidant compounds have been found to prevent the generation of reactive oxygen species (ROS) and thereby inhibit AOM-induced colon carcinogenesis in animal models [18].

AOM-induced oxidative stress lowered the endogenous antioxidant/detoxificant defence capacity in colon tissues of control rats, as evidenced by the significant decrease in glutathione (GSH) levels and in the activity of its related enzymes glutathione peroxidase (GPx), glutathione reductase (GR) and glutathione S-transferase (GST) (Fig. 1C and D). Nevertheless, the levels of enzymatic and non-enzymatic defences were preserved in the colon of cocoa-fed animals. Since GSH and its related enzymes participate in the detoxification of xenobiotics, carcinogens, free radicals and peroxides [19], we can suggest that cocoa could prevent AOM-induced ACF formation by reinforcing the endogenous defence capacity in colon tissues to counteract carcinogen-induced toxicity. These results are consistent with recent reports indicating that promotion of antioxidant enzyme activity has a chemopreventive effect on CRC [15, 20].

Besides inducing oxidative damage and genomic instability, ROS can specifically activate certain redox-sensitive signalling pathways and contribute to CRC initiation/promotion through the regulation of cellular proliferation and survival [21]. The phosphatidylinositol 3-kinases/protein kinase B (PI3K/AKT) and the extracellular regulated kinase/mitogen-activated protein kinase (ERK/MAPK) are of the most important pathways activated in response to oxidative stress that promote carcinogenesis via target proteins involved in cell survival and cell cycle progression [2]. In this line, a recent in vitro study has showed that flavonoids such as luteolin and quercetin have anti-proliferative and pro-apoptotic effects in human CRC cells through the regulation of the ERK/MAPK and the PI3K pathways [22]. In the present work, the proliferating cell nuclear antigen (PCNA) assay was used to quantify the colonocyte proliferation. Microscopic examination of colonic tissue sections clearly showed an increased level of PCNA-positive cells in the colonic mucosa of AOM-treated rats fed with control diet and a higher percentage of proliferating cells per crypt column (Fig. 2A). Indeed, the AOM-treated group fed with control diet exhibited an increase in the phosphorylation of ERKs and AKT together with an increased expression of the proliferative marker cyclin D1 (Fig. 2B and C). Conversely, cocoa intake prevented all these processes induced by AOM, suggesting that cocoa, by its ability to restrain oxidative stress could also inhibit the consequent activation of signalling pathways involved in proliferation and thereby the progression of preneoplasic lesions in the colonic epithelial cells.

Modulation of apoptosis provides an additional protective mechanism against intestinal neoplasia. Therefore, the apoptotic process in colonic samples from rats by terminal

Figure 2. Representative photographs for immunohistochemical staining of PCNA-positive cells (400 × magnification) and proliferative index (A) in colon tissues from rats injected with saline or AOM and fed control or cocoa-enriched diet. Representative Western blot analyses (B) and percentage levels of total and phospho-ERKs and -AKT and cyclin D1 (C) in colon tissues from the indicated groups. Bars represent mean ± SD value from at least 10 rats in each group and for three independent experiments with two samples each in the Western blot analysis. Means without a common letter differ significantly, p < 0.05.
Deoxynucleotidyl transferase dUTP nick end labelling (TUNEL) staining was further investigated. The representative photographs for TUNEL-positive cells clearly showed a pro-apoptotic effect of dietary cocoa in the colonic tissue of AOM-treated rats (Fig. 3A). Besides, the majority of the apoptotic cells were localized on the luminal surface, implicating that cocoa feeding could ensure the reestablishment of homeostasis in the colonic mucosa by removal of damaged cells. To better characterize the mechanisms involved, the colonic expression of pro-apoptotic (Bax) and anti-apoptotic (Bcl-xL) members of the Bcl-2 protein family as well as the activity of caspase-3 were also examined. As shown in Fig. 3B and C, AOM-treatment in control rats decreased Bax expression along with a concomitant increase in Bcl-xL proteins. However, the cocoa-enriched diet was able to increase Bax expression and decrease Bcl-xL proteins levels. Similarly, the activity of caspase-3 was significantly increased in the AOM group that was fed with the cocoa diet (Fig. 3D). These findings indicate that cocoa-enriched diet also induces apoptosis as another complementary mechanism of chemoprevention during the progression of carcinogenesis. Taken together, our data provide evidence that consumption of cocoa would offer a natural therapeutic approach to improve individual health status including potential efficient cancer prevention with minimal toxicity.

This work was supported by the grant AGL2007-64042/ALI and project CSD 2007-00063 from Programa Consolidador Ingenio from the Spanish Ministry of Science and Innovation (CICYT). I. Rodríguez-Ramiro is a predoctoral fellow of the Consejo Superior de Investigaciones Científicas and M. Gómez-Juaristi is a predoctoral fellow of the Spanish Ministry of Science and Education.

The authors have declared no conflict of interest.

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