Calcium carbonate suppresses haem toxicity markers without calcium phosphate side effects on colon carcinogenesis

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Red meat intake is associated with an increased risk of colorectal cancer. We have previously shown that haemin, Hb and red meat promote carcinogen-induced preneoplastic lesions, aberrant crypt foci (ACF), in the colon of rats. We have also shown that dietary calcium phosphate inhibits haemin-induced promotion and normalises faecal lipoperoxides and cytotoxicity. Unexpectedly, high-calcium phosphate control diet-fed rats had more preneoplastic lesions in the colon than low-Ca control diet-fed rats. The present study was designed to find a Ca supplementation with no adverse effect, by testing several doses and types of Ca salts. One in vitro study and two short-term studies in rats identified calcium carbonate as the most effective Ca salt to bind haem in vitro and to decrease faecal biomarkers previously associated with increased carcinogenesis: faecal water cytotoxicity and thiobarbituric acid-reactive substances. A long-term carcinogenesis study in dimethylhydrazine-injected rats demonstrated that a diet containing 100 µmol/g calcium carbonate did not promote ACF, in contrast with a previously tested calcium phosphate diet. The results suggest that calcium carbonate, and not calcium phosphate, should be used to reduce haem-associated colorectal cancer risk in meat eaters. They support the concept that the nature of the associated anion to a protective metal ion is important for chemoprevention.

Red meat: Haem: Calcium: Colorectal cancer: Prevention

Red and processed meat consumption increases the risk of colorectal cancer, according to meta-analyses of epidemiological studies\(^\text{(1,2)}\). In its 2007 report, the World Cancer Research Fund panel judges that ‘the evidence that red meat and processed meat are a cause of colorectal cancer is convincing’\(^\text{(3)}\). The panel thus recommends: ‘Limit intake of red and processed meat are a cause of colorectal cancer is convincing’\(^\text{(3)}\). The panel thus recommends: ‘Limit intake of red meat to healthy levels to promote colon cancer among women of the Swedish Mammography Cohort\(^\text{(13)}\). Experimental animal studies support the hypothesis that haem Fe promotes colorectal carcinogenesis\(^\text{(14)}\). We have shown that dietary haem, in the form of haemin (haem stabilised by a Cl atom, also called ferrisaproporphyrin IX chloride), Hb or red meat, promotes putative precancerous lesions, aberrant crypt foci (ACF) and mucin-depleted foci (MDF), in the colon of rats fed a low-Ca diet\(^\text{(15–17)}\). This haem-induced promotion is associated with increased lipoperoxidation and cytotoxicity of faecal water, and linked to urinary 1,4-dihydroxynonane mercapturic acid excretion, a fat peroxidation biomarker\(^\text{(18)}\). The addition of calcium phosphate to the diet inhibits haemin-induced lipoperoxidation, cytotoxicity and promotion of carcinogenesis in rats\(^\text{(17)}\), and we can explain the lack of effect of meat diets on colon carcinogenesis in previous study\(^\text{(19)}\) by a high concentration of Ca in the diet. The addition of calcium phosphate to beef meat diet fully prevents red meat promotion of colon carcinogenesis, but, unexpectedly, the rats fed the no-meat high-calcium phosphate high-fat diet had more ACF and more MDF than the rats fed the no-meat low-calcium phosphate diet\(^\text{(16)}\). In the absence of meat, dietary calcium phosphate appears to promote carcinogenesis in this rodent model, which precludes its use in human volunteers.

Abbreviations: ACF, aberrant crypt foci; MDF, mucin-depleted foci; TBARS, thiobarbituric acid-reactive substances.

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The present study was designed to determine a Ca supplementation with no adverse effect by testing different types of Ca salts and finding the minimum effective Ca dose.

Materials and methods

Animals and diets

Eighty-five Fischer 344 female rats were purchased at 4 weeks of age from Iffa Credo (St Germain sur l’Arbresle, France). Animal care was in accordance with the guidelines of the European Council on animals used in experimental studies. The animal colony and staff got official agreement no. 31-121 for animal studies by the French government.

Short-term studies. Two short-term studies were designed to find the most effective Ca salt and the minimum effective Ca dose to suppress early biomarkers of haem toxicity. Rats were housed individually in metabolism cages. The room was kept at a temperature of 22°C on a 12 h light–dark cycle. After 5 d of acclimatisation to the AIN76-powdered diet, rats were provided with the experimental diets. Body weights and food intake were monitored at the start and at the end of each study. In the first study, forty rats were randomly allocated to eight dietary groups (five rats per group) for 1 week. Faeces were collected at day 7 and were frozen at −20°C. Animals were killed 7 d after the start of the experimental diets. In the second study, twenty-five rats were randomly allocated to five dietary groups (five rats per group) for 2 weeks. Faeces were collected at day 14 and were frozen at −20°C. Animals were killed 14 d after the start of the experimental diets.

Long-term study. A 3-month study was designed to test the hypothesis that the minimum effective dose of the most effective Ca salt, selected from the results of the short-term studies, does not promote carcinogenesis in rats. Twenty rats were housed in pairs in stainless steel wire-bottomed cages. The room was kept at a temperature of 22°C on a 12 h light–dark cycle. Rats were allowed 7 d of acclimatisation to the room and to the control diet, before being injected intraperitoneally with the carcinogen 1,2-dimethylhydrazine (190 mg/kg body weight; Sigma Chemical, St Quentin, France) in NaCl (9 g/l). In most published studies, several carcinogen injections are administered to rats. We reasoned here that the first shot initiates carcinogenesis, but that the following shots would promote carcinogenesis, blurring diet-induced promotion. We thus chose a single-shot protocol, following Karkare et al. After 7 d, rats were randomly allocated to two groups of ten, and allowed free access to calcium carbonate-based diets for 94 d. We chose to initiate with the carcinogen in all rats, since the study was designed to show dietary promotion of carcinogenesis.

Diets. Experimental diets shown in Table 1 were based on a modified standard AIN76 diet prepared and formulated in a powdered form by UPAE (INRA, Jouy-en-Josas, France). For the first short-term study, high-beef meat diets were formulated to contain 60 % meat (w/w, freeze dried, 0.6 mmol haem/g of meat), and dibasic calcium phosphate was included in all the diets at a low concentration of 20 μmol/g. To determine the minimum effective dose of calcium phosphate, the effect of six concentrations of Ca was tested in six groups of rats fed the low-Ca beef meat diet supplemented with calcium phosphate (doses range: 33–250 μmol/g, diet names: Phos33…Phos250). To compare the efficacy of calcium phosphate with other salts, the effect of two Ca salts was tested in two groups of rats fed a calcium carbonate (Carb250) or calcium gluconate (Gluc250) diet at the highest tested dose (250 μmol/g). For the second short-term study, the effect of calcium carbonate was tested in five groups of rats fed a Hb diet, supplying 0.6 mmol/g haem to mimic the 60 % beef meat diet, supplemented with calcium carbonate (doses range: 33–155 μmol/g, diet names: Carb33…Carb155). For the long-term carcinogenesis study, two experimental diets were formulated to contain two levels of calcium carbonate (33 and 100 μmol/g, diet names: control diet and Carb100) added into the control low-calcium phosphate diet.

Preparation of faecal water

Faecal pellets were collected for 24 h under each cage (one rat/cage, short-term studies; two rats/cage, long-term study), thus leading to five samples per diet. Freeze-dried faeces were used to calculate dry faecal mass and to prepare faecal water by adding 1 ml of sterilised water to 0.3 g of dry faeces. Samples were then incubated at 37°C for 1 h, stirring thoroughly every 20 min, followed by centrifugation at 20 000 g for 10 min. The aqueous phase was re-centrifuged at the same speed and duration, and the subsequent supernatant (faecal water) was collected and stored at −20°C until use.

Thiobarbituric acid-reactive substances and haem assay

Thiobarbituric acid-reactive substances (TBARS) were measured in faecal water according to Ohkawa et al., exactly as described previously, and results are expressed as malondialdehyde equivalent. Haem contents of faecal water were measured by fluorescence according to Van den Berg et al. and Sesink et al., respectively, as described previously. Both TBARS and haem in faeces were measured on faecal samples from both the short-term studies.

In vitro experiment

The ability of five different Ca salts (1 – calcium phosphate, 2 – calcium chloride, 3 – calcium gluconate, 4 – calcium carbonate, 5 – calcium lactate) to bind and to precipitate haem from bovine Hb solution was studied in vitro. Hb (0.36 mmol; Sigma Chemical) was acidified at pH 1.4 with 2 mol HCl and then heated 1 h at 105°C to dissociate haem from Hb. This solution was centrifuged at 20000 g for 10 min. Supernatant with free haem was collected.

Six concentrations of each of five Ca salts were prepared (25, 50, 75, 100, 125 and 150 mmol/l). According to Sesink et al., 1 ml Hb added to 1 ml Ca salt was incubated for 90 min at 37.5°C in the dark and mixed every 10 min. The mixture was centrifuged for 2 min at 10000 g. Supernatant (100 μl) was assayed for the remaining haem after twenty-fold dilution in distilled water, by absorbance being measured at 400 nm with a Lambda 2 spectrophotometer (Pye Unicam-PU8600 UV/VIS-spectrophotometer Philips). Different Hb concentrations were prepared as standard
Table 1. Formulation of diets (g/100 g)*

<table>
<thead>
<tr>
<th></th>
<th>First short-term study: beef meat-based diets</th>
<th>Second short-term study: casein-based diets</th>
<th>Long-term study</th>
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<tr>
<td></td>
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<td>Phos55</td>
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<td>Calcium gluconate</td>
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</table>

* Phos20–Phos250 are low-Ca beef diets supplemented with calcium phosphate from 20 to 250 μmol/g. Carb250 and Gluc250 are low-Ca beef diets supplemented, respectively, with calcium carbonate and calcium gluconate at 250 μmol/g. Carb33–Carb155 are low-Ca Hb diets supplemented with calcium carbonate from 33 to 155 μmol/g. Control diet and Carb100 low-Ca Hb diet supplemented with calcium phosphate from 33 to 100 μmol/g.
† Low-Ca casein.
‡ AIN76 mix, but 500 g/kg of dibasic calcium phosphate were replaced by sucrose in mineral mix.
solutions: 1/1 (100%), 1/5, 1/10, 1/20 and 1/50 in water. The assay was repeated three times, and the mean values were recorded.

Aberrant crypt foci assay

Rats were killed by CO2 asphyxiation in a random order at day 94 of the long-term study. Colons were excised from rats immediately post-mortem, flushed with cold Krebs solution (Sigma Chemical), opened longitudinally and fixed flat between two sheets of filter paper in 10% formalin (Sigma Chemical), marked with a two-digit blinding code. Colons picked up in random order were stained for 6 min in a 0·05% filtered solution of methylene blue25. The number of ACF per colon and number of crypts in each ACF were recorded. The assay was repeated three times, and the mean values were reported.

Statistical analysis

Results were analysed using SYSTAT 10 software for Windows and reported as means and standard deviations. Long-term study data were analysed by Student’s t test, except ACF data that were analysed by two-way ANOVA (dietary group and reader). Short-term study data were considered first using one-way ANOVA. If a significant difference was found between the groups (P<0·05), then pairwise comparisons were made with Fisher’s least significant difference multiple comparison test.

The meta-analysis of the relation between calcium carbonate and colon cancer in carcinogen-injected rats was done as a sub-analysis of a study that we published in 200526. From this previous meta-analysis, we selected studies where calcium carbonate had been fed to experimental groups of rats. The studies were homogeneous (Q Cochran P>0·05); we thus used the ‘Fixed Effects’ model, using EasyMA DOS software (2001 version) to compute data.

Results

Weight and food intake during in vivo studies

Final body weight of rats was 107 (sd 2) g after 7 d on the experimental diets in the first short-term study, without significant differences between the groups. Food intake was not significantly different (data not shown), but Gluc250-fed rats tended to eat less than the other groups (Gluc250: 7·2 (sd 2·0) g/d, all other groups: 9·1 (sd 1·0) g/d, P = 0·09). No difference in body weight and diet intake between the groups was observed in the second short-term study or in the long-term study. Mean body weights were, respectively, 166 (sd 2·0) g after 14 d and 201 (sd 8·0) g after 94 d on the experimental diets.

First short-term study: thiobarbituric acid-reactive substances of faecal water

Haem can induce the formation of peroxyl radicals in rats, which may be cytotoxic and cleave DNA in vivo. This might explain the promoting effect of haem on colon carcinogenesis. Lipid peroxidation was thus measured in faecal water by TBARS assay. Ca can prevent haem-induced promotion of colon carcinogenesis by precipitating haem in the colonic content; thus haem concentration in faecal water was measured.

Haem concentration in faecal water from beef meat-fed rats was significantly reduced in rats fed 150 and 250 mmol/g calcium phosphate (P<0·01; Fig. 1(a)), but not in rats fed lower levels of calcium phosphate. Dietary calcium carbonate and 250 mmol/g gluconate produced the same effect on haem than calcium phosphate (Fig. 1(a)). Surprisingly, haem-induced lipid peroxidation measured by TBARS level was not significantly reduced by the calcium phosphate diets, but only the high-calcium carbonate diet decreases TBARS level in faecal water (P<0·01; Fig. 1(b)).

In vitro study

Any tested concentration of calcium carbonate higher than 25 mmol was much more effective to precipitate haem than any other tested Ca salt. Proportion of trapped haem was more than three times higher with calcium carbonate than with calcium phosphate (100 mmol) (P<0·001; Fig. 2).

Second short-term study: thiobarbituric acid-reactive substances of faecal water

Beef meat was replaced by Hb in the second study diets, since we have demonstrated previously that they produce similar effects on colon carcinogenesis and on faecal water peroxidation15. The high-calcium phosphate diet (250 mmol/g) did not prevent haem toxicity in the 7 d study, in contrast with our previous 50 d study17. We suggest that this duration was too short to reduce the level of TBARS; thus we chose the second short-term study to last for 14 d. As calcium carbonate was the only effective salt against haem-induced lipid peroxidation in vivo and the only effective salt to precipitate haem in vitro, this salt was used to determine the smallest effective Ca dose.

Haem concentration was lower in faecal water from rats fed a diet containing at least 80 mmol/g calcium carbonate (P<0·001; Fig. 3(a)). Accordingly, diets containing at least 80 mmol/g calcium carbonate reduced haem-induced lipid peroxidation of faecal water (P<0·001; Fig. 3(b)).

Aberrant crypt foci assay

We chose to test if a diet containing 100 mmol/g calcium carbonate would promote colon carcinogenesis because the second short-term study showed that a 80 mmol/g calcium carbonate diet reduces faecal TBARS concentration, and that 110 mmol/g seemed twice more potent than 80 mmol/g to bind faecal haem. In contrast with a calcium phosphate diet16, the consumption of a diet containing 100 mmol/g calcium carbonate for 94 d did not increase significantly the number of ACF per colon, 101 d after a dimethylhydrazine injection (Fig. 4).

Discussion

These data confirm that dietary Ca reduces faecal water lipoperoxides level, a biomarker linked with haem-induced
carcinogenesis promotion. They also establish that the type of Ca salt is paramount in this protection, and that calcium carbonate is effective without promoting the side effect of calcium phosphate.

Red meat promotes ACF and MDF in the colon of carcinogen-induced rats, and haem-induced promotion by meat is associated with faecal water cytotoxicity and TBARS\(^{15,16}\). The mechanism of promotion by haem is not known and may be linked to lipid peroxidation. We have explored the effect of faecal water from beef meat-fed rats on normal (Apc\(^{+}/^{+}\)) and premalignant colonic cells (Apc\(^{-}/^{-}\)). Results show that Apc-mutated cells survive in haem-induced faecal lipoperoxides, notably 4-hydroxynonenal that is toxic to normal cells. Selection of mutated cells by cytotoxic lipoperoxides may thus explain haem-induced promotion of colon carcinogenesis\(^{27}\). Ca added on top of beef meat or haemin diets normalises the number of ACF and MDF at the same low level as in the control group\(^{17}\). Calcium phosphate precipitates haem in the gut, and little haem remains available to induce lipid peroxidation\(^{17,24}\). The trapping of haem by Ca thus abolishes carcinogenesis promotion. In the present study, we established that different Ca salts are able to precipitate haem \textit{in vitro} and in faecal water of haem-fed rats and by consequence are able to reduce faecal water peroxidation (Figs. 1 and 2). We observed that only high concentrations of calcium phosphate were able to precipitate haem (Fig. 1(a)). Furthermore, the ability of the three tested salts (phosphate, carbonate and gluconate) to trap haem was very similar in high-Ca diets (Fig. 1(a)). However, only calcium carbonate decreased significantly the level of lipid peroxidation in faecal water (Fig. 1(b)). So, to screen more precisely
the efficiency of different Ca salts, we measured the in vitro trapping of haem: only calcium carbonate precipitated haem significantly. In conclusion, the present study shows for the first time a difference of efficiency between Ca salts in vitro (Fig. 2) and in vivo, in the trapping of haem and the limitation of peroxidation (Fig. 1(b)). We have no explanation for the relatively high TBARS level in controls and in rats administered with high doses of calcium phosphate or gluconate: we speculate that the beef meat in the first study may have contained already oxidised fat or oxidising agents different from haem Fe (Fig. 1(b)).

Our previous study showed that rats consuming a high-Ca control diet had more ACF and more MDF than rats consuming a low-Ca control diet. This promotion was observed with calcium phosphate in a high-fat diet context, but was not observed in a low-fat diet context without a clear explanation for the high-fat/low-fat discrepancy. Bull et al. also showed that the promotion of tumours by calcium phosphate was more important in 30%-fat diet-fed rats than in 3%-fat diet-fed rats. Phosphate, not Ca, might be the promoting nutrient, an issue already discussed by Bruce et al. Since our in vitro study indicated that calcium carbonate was more effective than calcium phosphate to trap haem (Fig. 2) and since the first in vivo study indicated that a high concentration of calcium carbonate was able to precipitate haem and to limit peroxidation in vivo, we determined the lower effective dose of calcium carbonate in a second short-term study (Fig. 3). We showed that 80 μmol calcium carbonate/g diet is enough to trap haem and to reduce peroxidation. The first aim of the present study was to identify the lower dose and type of Ca sufficient to limit the effect of haem in the colon without side effects. We thus tested the effect of an addition of 100 μmol calcium carbonate/g diet on the promotion of colon carcinogenesis at the level of ACF (Fig. 4). In contrast with the promoting effect of calcium phosphate at 250 μmol/g, we did not observe a promoting effect.

**Fig. 3.** Effect of calcium carbonate-based diets (30–155 μmol/g) on faecal water values after the second short-term study. (a) Haem concentration in faecal water (μmol/l). (b) Thiobarbituric acid-reactive substances concentration in faecal water, expressed as μmol/ml malondialdehyde (MDA) equivalents. * Mean values were significantly different from Carb30 by Fisher’s least significant difference test (P<0.05). Data are means, and bars are standard deviations (n 5 in each group). Note: Carb33–Carb155 are low-Ca Hb diets supplemented with calcium carbonate from 33 to 155 μmol/g.

**Fig. 4.** Effect of calcium salts in diet on putative precancerous lesions (aberrant crypt foci (ACF)) per rat colon 101 d after the injection of dimethylhydrazine. Results of the present study with 100 μmol/g calcium carbonate are represented by □. Results of the previous study with 250 μmol/g calcium phosphate (Pierre et al.) are represented by □. Results are expressed after normalisation of the control diet groups to 100. * Mean values were significantly different from Carb33 by Fisher’s least significant difference test (P<0.05). Data are means, and bars are standard deviations (n 5 in each group).
CaCO₃ blocks haem without side effects

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References


