Phytase activity in the human and rat small intestine

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Abstract
Phytate is the major storage form of phosphorus in seeds and so is a common dietary constituent. Excessive ingestion of undegraded phytates can cause mineral deficiencies in humans. In addition, phytic acid is antimetastatic in animal models of both colon and breast carcinoma. There have been no previous studies quantifying phytase activity in the human small intestine although it is present in animals. Small intestinal phytase and alkaline phosphatase activity and distribution was measured in vitro in mucosal homogenates from two human small intestinal specimens obtained from transplant donors. Rat intestine was also studied for comparison. Phytase activity was found in human small intestine at low values (30 times less than that in rat tissue and 1000-fold lower than alkaline phosphatase in the same tissue). The activity was greatest in the duodenum and lowest in the ileum. In conclusion, the normal human small intestine has very limited ability to digest undegraded phytates. Although this may have adverse nutritional consequences with respect to metabolic cation imbalances, the presence of undigested phytates in the colon may protect against the development of colonic carcinoma.

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Phytate (myo-inositol hexaphosphate) is the major storage form of phosphorus in seeds and as such forms a variable large component of the diet of many animals and humans. Phytic acid is a strong chelator of cations and has been shown to cause deficiencies in minerals such as calcium, iron, zinc, and magnesium when present in excess in the human diet.2 Phytase (E.C.3.1.3.8.) is present in the small intestines of animals such as rats, calves, chickens, and pigs.3–5 Dietary phytate intake and the ability of the intestine to break down phytic acid have been extensively studied in animals because of the importance of phytic acid in mineral balance and the potentially deleterious effect of high concentrations of undegraded phosphates in livestock excreta.6 7 Despite the fact that humans consume large amounts of phytates, which can cause mineral deficiencies,8 published works on the ability of the human intestine to degrade phytic acid are surprisingly sparse. There has only been one previous study on phytase activity in the human small intestine, which found it to be present but made no mention of the activity of the enzyme or its distribution in the intestine.5 Metabolic balance studies in human volunteers have suggested that the human intestine is poor at metabolising phytic acid.9

Our aim was to measure human intestinal phytase activity in vitro on fresh small intestinal material and to compare this with the activity in the rat. As there has been controversy in the past as to whether phytase activity actually results from alkaline phosphatase,10 we planned to measure alkaline phosphatase activity in the same tissues and to use phenylalanine, which is known to inhibit alkaline phosphatase but not phytase.11

Methods
Phytase activity is determined by measuring the liberation of inorganic phosphate when inositol hexaphosphate is broken down. Most methods for measuring phosphate are based on the colour formed by the reduction of phosphomolybdate complexes.12 Various reducing agents have been used. The application of this principle to the determination of phytase activity has been complicated by the finding that phytase itself interferes in these colorimetric determinations of inorganic phosphate.13 A published method for the determination of phytase in duodenal biopsy specimens entails the reduction of phosphomolybdate complexes with ascorbic acid after the extraction of these complexes from remaining phytates using isobutyliheptane.14

In a preliminary trial of this method, however, we found that the variability at the extraction and reduction phases made the procedure too imprecise for the determination of the very small amounts of phosphate released by the human tissue. We therefore used a method for phosphate measurement using malachite green as the chromogen.15 This improved method does not entail phytate extraction before colour development. It also has the added advantage of not requiring protein precipitation as deproteinization has been shown to release phosphate groups from cellular constituents.15

REAGENTS
The following reagents were used: sodium phytate (30 mmol/l), hydrochloric acid (5 mol/l); malachite green made up in acidified molybdate (340 mg/l); aqueous TWEEN solution (15 ml/l); ammonium molybdate in hydrochloric acid (34 mmol/l); aqueous urea solution (360 g/l); aqueous zinc chloride (4 mmol/l); aqueous magnesium chloride (40 mmol/l).
SPECIMENS

Complete human small intestinal specimens were obtained from two previously fit subjects during organ harvesting for transplantation purposes. One was a 50 year old woman who died suddenly from a subarachnoid haemorrhage; the other a 40 year old man who died suddenly in a road traffic accident.

Rat small intestine was harvested from two Wistar male rats killed by cervical dislocation. All tissue samples were immediately frozen in liquid nitrogen. Subsequently the rat and human donor specimens were kept at −70°C.

PHYTASE DETERMINATION

For the determination of phytase activity the brush borders were scraped off underlying tissue with a glass microscope slide before homogenisation.

The tissue (approx 200 mg wet weight) was homogenised in 45 ml cold 50 mmol sucrose plus 2 mmol TRIS HCl buffer (pH 7·1) with 20 strokes of a Griffiths tube homogeniser. The homogenate was centrifuged at 6000 rpm for three minutes and the supernatant separated from the pellet, which was discarded. An aliquot was taken for alkaline phosphatase activity, which was estimated on the 'Cobas-Bio' automated enzyme analyser using paranirophenol phosphate as substrate,16 and for protein determination by the method of Bradford.17

Homogenate (2 ml) was combined with MgCl₂ solution (100 µl) and zinc chloride solution (100 µl). Sodium phytate substrate (100 µl) was then added to the tests and water added to the blanks. Reagent blanks consisting of buffer with or without phytate were included. The tubes were then incubated at 37°C for two hours after which an aliquot (100–500 µl) was taken for determination of inorganic phosphate content. Urea solution (1 ml) was added to this to prevent protein precipitation and then malachite green reagent (4 ml) containing acidified molybdate. After allowing 10 minutes for colour development, the samples were read at 640 nm on a Pye Unicam PU8610 U/V/Visible spectrophotometer.

For the measurements on rat tissue we found that 100 µl aliquots from the incubation mixture gave good colour formation. The much smaller phytase activity in the human intestine, however, required 500 µl of incubate and at this concentration there is a contribution from undegraded phytate to the final colour. This could be compensated for by reagent blanks consisting of buffer with and without phytate.

The concentrations of phosphate in the incubates and blanks were derived from the standard curve.

The phytase activity expressed as µmolP/min/mg tissue was calculated as:

\[ \text{Phyrase activity (µmolP/mg protein/min)} = \frac{(T_p - T) - (B_p - B)}{\text{protein conc} \times \text{time}} \]

where Tp: tissue + phytate, T: tissue blank, Bp: buffer + phytate, B: reagent blank.

Results

WORKING RANGE OF MALACHITE GREEN METHOD

By plotting the absorbance of varying concentrations of phosphate standards (Fig 1) we derived a linear working range of between 1 and 5 mmol/l.

FREEDOM FROM INTERFERENCE BY PHYTATE

Phosphate standards with added phytate up to 400 times that used in the final colour reaction product had negligible effect on the absorbance (Fig 2).

FREEDOM FROM SPONTANEOUS HYDROLYSIS OF PHYTATE

Reagent blanks comprising buffer with and without phytate showed negligible effects on the final absorbance from phytase showing that under the experimental conditions there was no spontaneous hydrolysis of phytic acid.

![Figure 1: Working range for malachite green method.](image1)

![Figure 2: Effect of phytate on malachite green method for phosphate.](image2)

![Figure 3: Rate of phosphate liberation during incubation of rat duodenal homogenate with phytate.](image3)
TABLE I Rat duodenal phytase and alkaline phosphatase activity and kinetics. Mean (SD) on six replicates shown

<table>
<thead>
<tr>
<th>Activity</th>
<th>Vmax</th>
<th>Km (mmol/l)</th>
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<tbody>
<tr>
<td>Phytase (umolP/mg/min)</td>
<td>270 (4.8)×10^{-1}</td>
<td>0.83</td>
</tr>
<tr>
<td>Alkaline phosphatase (umolP/mg/min)</td>
<td>273 (0.02)</td>
<td>4.21</td>
</tr>
<tr>
<td>Alkaline phosphatase phytase ratio</td>
<td>102</td>
<td></td>
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This is at variance with the much lower activity found in this study.

Nutritionists have estimated human phytase activity indirectly by studying the breakdown of various foodstuffs by the whole gut in normal controls and in patients with ileostomies and have suggested that intestinal phytase activity is very low.20,21 Our direct measurements of human intestinal phytase activity in vitro would support these findings.

The ability of the small intestine to degrade phytic acid is of considerable nutritional importance as evidenced by the mineral deficiencies seen in human populations consuming large quantities of undergraded phytates.20–23 Phytates in the diet are probably hydrolysed mainly by endogenous phytase in food (for example, yeast phytase) and this breakdown probably occurs largely in the acid environment of the stomach.9 This would explain why most people taking diets containing modest amounts of phytates do not become deficient in minerals.

An additional recent development in phytate metabolism has been the hypothesis that phytate may be protective against colonic carcinogenesis.24 Phytic acid has recently been found to be anti-neoplastic in both mammary and colon carcinogenesis models in vivo in the rat.25 26

The low level of phytase activity in the intestine dictates that, from the nutritional point of view, foodstuffs comprising high concentrations of phytic acid should undergo processing before human consumption.27 In view of the recent evidence, however, that phytic acid is anti-neoplastic28 it may be that consumption of undergraded phytates should be encouraged notwithstanding the nutritional implications.