Latest published research in phytate-protein interactions highlights differences between phytase enzymes in practice

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A breakthrough in our understanding of phytase enzyme mode of action and the associated matrix values in feed formulation has been provided in a recent paper published by Tran et al. in the January 2011 edition of Analytical Biochemistry entitled ‘A simple and fast kinetic assay for phytases using phytic acid-protein complex as substrate’.

While at first glance the relevance of the title to animal nutritionists may not be immediately obvious, this collaborative research by scientists from Lund University (Sweden) and Genencor studied interactions of phytate with dietary proteins as measured by the formation of insoluble complexes and the suitability of using the phytate-protein complex as a substrate for phytase assays. The authors demonstrated that, as phytate-protein complexes occurred in feed and the digestive tract of animals, this may be a more relevant substrate for the evaluation of phytases than sodium phytate, a laboratory chemical routinely used in traditional phytase assays. Of commercial interest was that the study used in the evaluation of phytases than sodium phytate, which is used as the substrate to determine phytase activity in-vitro, in nature also in the acid part of the digestive tract of poultry (proventriculus and gizzard) and swine (stomach), phytic acid does not occur in its free acid or sodium salt form, but rather in protein interactions highlights differences between phytase enzymes in practice.

To fully understand the research results shown in Table 1 it is important to know that while sodium phytate is used as the substrate to determine phytase activity in-vitro, in nature also in the acid part of the digestive tract of poultry (proventriculus and gizzard) and swine (stomach), phytic acid does not occur in its free acid or sodium salt form, but rather in the gut is that most proteins of plant origin, such as those derived from maize, soybean, sunflower and rapeseed (canola) meal, have their iso-electric points (pI values) in the acidic range (pH 4.5). Hence, when the pH in the digestive tract drops below the iso-electric point of the protein, the protein is now positively charged, which allows the negatively charged phytic acid molecule to bind to it, thereby altering the proteins iso-electric point and rendering the protein insoluble (Reddy et al., 1989; Konietzny and Greiner, 2003). This formation of insoluble protein-phytate complexes in the acid stomach this data provides the first strong evidence that the nutritional benefit from phytase enzymes can no longer be viewed as being constant within classes of phytase such as E. coli. Consequently, in practice, different matrix values should be applied in feed formulation to account for these differences.

To converge to the negative effects of phytate on energy and amino acid digestibility, nutritional advantages of phytase enzymes will depend primarily on the ability of the source of phytase to rapidly hydrolyze phytate-protein complexes at a low pH (gizzard and proventriculus region), thereby reducing their anti-nutritional effects and resulting in net improvements in energy and protein utilization.

Table 1: Activity of different commercial phytases at pH 3.0 and 37˚C on soy protein or lysozyme protein complexes with phytate (IP6) as compared with sodium-phytate (IP6-Na) as substrate. All phytases were added at a dose of 0.1 FTU/ml (1FU from Tran et al., 2011).

<table>
<thead>
<tr>
<th>Phytase source</th>
<th>IP6-soy protein</th>
<th>IP6-lysozyme</th>
<th>IP6-Na</th>
<th>Relative activity (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>E. coli phytase 1*</td>
<td>164</td>
<td>229</td>
<td>100</td>
<td></td>
</tr>
<tr>
<td>E. coli phytase 2*</td>
<td>138</td>
<td>152</td>
<td>103</td>
<td></td>
</tr>
<tr>
<td>A. niger phytase</td>
<td>32</td>
<td>23</td>
<td>37</td>
<td></td>
</tr>
<tr>
<td>P. lycii phytase</td>
<td>25</td>
<td>13</td>
<td>10</td>
<td></td>
</tr>
</tbody>
</table>

1 Activities of the phytases (amount of phosphate released) on the other substrates are reported relative to the activity of E. coli phytase variant 1 on sodium phytate (IP6-Na).
* E. coli phytase 1 was from Schizosaccharomyces pombe and E. coli phytase 2 from Pichia pastoris.

Of further importance are new data that have shown the formation of insoluble protein-phytate complexes to be proportional to the amount of phytate present. This is shown in Figure 1 where the presence of insoluble protein-phytate complexes from either soy protein or casein, measured by absorbance at 600 nm, increased almost linearly as the phytate concentration increased. Stated another way, the anti-nutritional effect of phytate in the acid stomach of the animal will be proportional to the concentration of undigested phytate (IP6) present in the acid stomach and is further supported by previous work by Cowieson et al. (2009), who demonstrated a step-wise increase in endogenous amino acid losses at the terminal ileum as dietary phytate levels increased. Converse to the negative effects of phytate on energy and amino acid digestibility, nutritional advantages of phytase enzymes will depend primarily on the ability of the source of phytase to rapidly hydrolyze phytate-protein complexes at a low pH (gizzard and proventriculus region), thereby reducing their anti-nutritional effects and resulting in net improvements in energy and protein utilization.
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In conclusion, the past decade of research has shown anti-nutritional effects of phytate to extend beyond a simple reduction in phosphorus and calcium availability and include negative effects of phytate on protein-binding, endogenous nutrient losses, and energy and amino acid utilization. An understanding by nutritionists that the formation of insoluble protein-phytate complexes in the acid stomach is a primary mode of action whereby phytate exerts its anti-nutritional effects and conversely, whereby phytase enzymes contribute to improved energy and amino acid utilization is essential in order to make informed decisions on selecting appropriate phytase sources for in-feed application.

The clear differences between the sources of phytase evaluated in the research by Tran and co-workers would suggest that the nutritional benefit of the different phytases cannot be the same, which makes a critical reassessment of the associated matrix values provided by the phytase suppliers imperative. Furthermore, as the effects of phytate on binding protein in the acid environment are also proportional to the phytate concentration and dose of phytase in feed, the matrix value applied to phytases in feed formulation can be summarised as being a function of these three factors and should be adjusted based on:

1. The source of phytase enzyme and its ability to hydrolyze phytate-protein complexes at low pH.
2. The phytate level of the diet, with reduced benefits in energy and amino acid utilization being derived from phytases at lower levels of dietary phytate.
3. Higher phytase doses provide a greater ratio of enzyme:substrate in the digesta, allowing for a more rapid degradation of the protein-phytate complex, further reducing its anti-nutritional effects and improving net dietary energy and amino acid utilization.

Key references

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