Effect of level of feed intake and *Fusarium*-contaminated wheat on rumen fermentation in cows

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ABSTRACT

To study the effects of level of intake on ruminal fermentation, 14 dairy cows equipped with ruminal and duodenal cannulas were fed different amounts of the same diet. The diet consisted of 60% concentrate including 55% wheat (*Fusarium*-contaminated and control) and 40% maize- and grass silage (on DM-basis). Each cow was fed contaminated and control wheat. Rumen fermentation was not significantly influenced by the contamination, however the increased feed intake decreased pH, fermentation of organic matter (FOM), of protein and of neutral detergent fibre. Efficiency of microbial protein synthesis (g MP/kg FOM) was not linearly related to level of intake.

KEY WORDS: cow, rumen fermentation, level of feed intake, mycotoxins

INTRODUCTION

A high passage rate as a result of increased feed intake can decrease the ruminal digestibility of nutrients (Tyrrell and Moe, 1975) and the metabolization rate of mycotoxins, but increase the efficiency of the microbial protein synthesis (Sniffen and Robinson, 1987).

In most experiments increasing feed intake involved a higher concentrate portion in the ration. Then effects of roughage:concentrate can not be separated from effects of level of intake. Hence, the objective of the present study was to investigate the coherence between varying levels of feed intake on ruminal fermentation in cows feeding a constant ration with or without *Fusarium*-contamination of the wheat.

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MATERIAL AND METHODS

14 dairy cows of the German Friesian breed fitted with permanent fistulas in the rumen and the proximal duodenum were used. All diets contained 60% concentrate (on DM-basis), including 55% wheat (*Fusarium*-contaminated or not) and 40% maize- and grass-silage (50:50). Dry matter (DM)-intakes ranged from 5.7 to 20.9 kg/d. Each cow was fed the concentrate with the uncontaminated and with the contaminated wheat successively.

Each experimental period consisted of three weeks for adaptation to the diets and 5 days for duodenal-sample collection. Ruminal fluid for the determination of pH and volatile fatty acids was sampled in the 3rd week 3h after start of morning feeding. To estimate digesta flow at the duodenum, Cr$_2$O$_3$ was used as marker. The proportion of microbial N of non ammonia N (NAN) at the duodenum was determined by NIRS (Lebzien and Paul, 1997). Fermented organic matter (FOM) was calculated by:

$$\text{FOM (kg/d)} = \text{OM intake} - (\text{OM flow at the duodenum} - \text{microbial OM})$$

(Microbial OM = microbial N $\times$ 11.8)

The undegraded protein at the duodenum was estimated by subtracting endogenous protein (kg DM at the duodenum $\times$ 3.6) and the microbial protein from NAN $\times$ 6.25.

RESULTS AND DISCUSSION

The contents of crude nutrients and the *Fusarium*-toxins DON and ZON for the concentrates and silages are shown in Table 1.

Table 1. Chemical composition and mycotoxin concentrations of the feedstuffs

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Concentrate$^1$</th>
<th>Maize silage</th>
<th>Grass silage</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>mycotoxin period</td>
<td>control period</td>
<td></td>
</tr>
<tr>
<td>Dry matter, g/kg</td>
<td>877</td>
<td>885</td>
<td>395</td>
</tr>
<tr>
<td>Nutrients, g/kg DM</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>organic matter</td>
<td>945.6</td>
<td>944.0</td>
<td>960.1</td>
</tr>
<tr>
<td>crude protein</td>
<td>207.5</td>
<td>209.5</td>
<td>76.1</td>
</tr>
<tr>
<td>crude fibre</td>
<td>66.0</td>
<td>66.7</td>
<td>174.0</td>
</tr>
<tr>
<td>neutral detergent fibre</td>
<td>221.8</td>
<td>225.8</td>
<td>404.0</td>
</tr>
<tr>
<td>Mycotoxins</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>deoxynivalenol, mg/kg DM</td>
<td>5.2</td>
<td>0.1</td>
<td>0.6</td>
</tr>
<tr>
<td>zearalenone, µg/kg DM</td>
<td>59.4</td>
<td>21.7</td>
<td>62.4</td>
</tr>
</tbody>
</table>

$^1$55% (*Fusarium*-contaminated) wheat, 25% soya bean meal, 16% dried sugar beet pulp, 2% soya oil, 2% mineral feed  
$^2$DL - detection limit
Increased feed intake significantly decreased the ruminal pH (P<0.001; Figure 1).

In accordance to Tyrell and Moe (1975) the fermented portion of organic matter decreased moderately at increased organic matter intake but there was a quite large individual variation (P<0.001; Figure 2). The most pronounced decrease was observed with regard to NDF-fermentation.

Figure 1. pH-value in rumen fluid (collected 3 h after morning feeding) in dependence on the organic matter intake, —□— Control period, ▲— Mycotoxin period, numbers denote individual cows

Figure 2. Fermented organic matter (% of OM intake) in dependence on the organic matter intake, —□— Control period, ▲— Mycotoxin period, numbers denote individual animals
Although increasing organic matter intake significantly increased the quantities of microbial protein (P<0.001) and undegraded feed protein (P<0.001) at the duodenum, the relationship between efficiency of microbial protein synthesis and level of organic matter intake was not linear (Figure 3).

![Figure 3. Efficiency of microbial protein synthesis in dependence on the organic matter intake, □ Control period, ▲ Mycotoxin period, numbers denote individual animals](image)

In accordance to Sniffen and Robinson (1987), who recalculated from the studies of Tamminga (1981) the lowest efficiency of microbial protein synthesis at about 10 kg of DM-intake, the efficiency was lowest at 9 kg organic matter intake and higher at intakes which were higher or lower.

*Fusarium*-contamination of the wheat did not affect the daily flow (g/d) of microbial protein (P=0.904).

Further analyses of duodenal digesta will give information about the metabolization of the mycotoxins in the rumen.

REFERENCES


