Utilization of raw or heat-treated starch fed in liquid diet to pre-ruminants. 1. Calves *

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Summary — Five-7 day-old Friesen bull calves, raised in Israel received twice a day a mixture of 40 g of soybean concentrate (65% protein) and 80 g of one of the following carbohydrates: glucose (G), expanded (heat-treated) (ES) or untreated (US) corn starch. In the afternoon the calves received in addition 400 g of milk replacer. Until weaning at experimental day 25, ES calves showed better growth and food utilization than their counterparts. Daily weight gain for the entire growing period up to slaughter was higher for the starch than for the glucose-fed group (P < 0.05 for ES vs G). Heat treatment of starch increased its in vitro availability to amylase and its in vivo digestibility. By-pass of the rumen was complete in all G calves. In the ES and US groups, partial diversion of the liquid feed into the rumen was evident. The G group showed hyperglycemia after meals, while almost no increase in blood glucose level was observed after soy-starch meals (either US or ES).

calf / milk replacer / soy-protein / starch / digestibility / blood glucose

Résumé — Utilisation de l’amidon cru ou traité à la chaleur chez les préruminants. 1. Veaux. Des veaux israéliens de race Frisonne âgés de 5–7 j ont reçu 2 fois par jour un mélange liquide contenant 40 g de concentré de soja (65% de protides) et 80 g d’un des glucides suivants : glucose (G), amidon de maïs traité à la chaleur (ES) ou non traité à la chaleur (US). Dans l’après-midi, les veaux ont reçu en supplément 400 g d’un aliment d’allaitement. Au sevrage, les veaux ES étaient plus lourds que les veaux G ou US et leur indice de consommation meilleur. Le gain de poids moyen quotidien durant toute la période de croissance, jusqu’à l’abattage, était supérieur dans le groupe ES. Le traitement thermique appliqué à l’amidon de maïs a amélioré sa digestibilité in vivo et in vitro. La fermeture de la gouttière œsophagienne était complète chez les veaux du groupe G contrairement aux veaux recevant l’amidon. Dans le groupe G, une hyperglycémie postprandiale a été constatée, alors que chez les veaux ES et US la concentration du glucose dans le sang n’a pas augmenté après le repas.

veau / aliment d’allaitement / protéine de soja / amidon / digestibilité / glycémie

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INTRODUCTION

The effect of expanding or pelleting the feed on digestibility in ruminants is well documented (Haenlein et al, 1962; Theurer, 1986). The effect of starch from different sources, as well as the effect of heat treatment of starch included in milk replacers for veal calves, is also known (Mathieu and Thivend, 1968; Toullc et al, 1973, 1979), but the above studies began with calves weighing 60 kg or more and fed about 2 kg of milk replacers daily. The effect of including different starches in milk replacers for perinatal pre-ruminants consuming about 0.5 kg milk replacer daily is not as well known. Improving the utilization of feeds of plant origin used in milk replacers may be of economical value. Dried milk products are used extensively in the production of commercial milk replacers for rearing calves, kids and lambs. Young calves fed milk or milk replacers ad libitum can consume and digest enough nutrients to achieve a daily average growth of 1 kg (Marshall and Smith, 1970). The amount of milk replacers fed in practice is limited due to economic reasons.

The cost of milk replacers can be reduced in substituting feeds of plant origin for some of the milk ingredients.

The digestive system of pre-ruminants is adapted to the efficient digestion and utilization of milk components. It was shown by direct and indirect methods that the capacity of young pre-ruminants to digest and utilize ingredients from sources other than milk is limited (Huber et al, 1961; Siddons, 1968; Siddons et al, 1969; Ben-Asher et al, 1981).

Toasted soybean concentrate (65% protein) may replace part of the milk proteins; this results in some reduction of calf performance. This reduction may be compensated after weaning, when dry feeds are given (Nitsan et al, 1972).

Contradictory reports exist regarding the utilization of starch by pre-ruminants when fed in liquid form with milk replacer. Poor growth of suckling calves fed milk replacers containing more than 10% of raw corn starch was reported by Flipse et al (1950) and by Noller et al (1956). Including 18.5% starch in milk replacer for calves slightly reduced the digestibility of organic matter compared with whole milk, but growth was similar in the 2 groups (Burt and Irvine, 1970). Thivend (1978) reported good performance of calves fed milk replacers containing about 20% starch, especially from 8 weeks of age onward. It was also shown that substituting part of the lipids in milk replacer by starch improved growth and reduced adiposity (Thivend et al, 1979). The utilization of starch from various sources may differ and be affected by cooking. Heat treatment of starch, which causes partial gelatinization, increases solubility and availability to amylolytic enzymes (Schoch and Maywald, 1967).

The purpose of this work was to compare the effects of the addition of raw vs heat-treated corn starch in milk replacer for young calves on growth, digestibility of nutrients, and rate of glucose absorption.

MATERIALS AND METHODS

Animals and feeds

Twelve Friesian bull calves, raised in Israel were purchased from various dairy farms after they had received colostrum for 3 d, followed by a commercial milk replacer. At 5–7 d of age, the calves were allotted to 3 experimental groups of 4 calves each according to body weight and kept in individual metabolic pens with wooden slatted floors.

For the morning meal (07 00 h), the calves received a mixture of 40 g of soybean concentrate (65% protein) and 80 g of one of the following carbohydrate sources: glucose (G), raw corn
starch (US) or expanded (heat-treated) corn starch (ES) (Galam Ltd, Maanit, Israel), mixed in 1.5 l of lukewarm (37 °C) water. At 17 00 h, 400 g of commercial milk replacer and 120 g of one of the soy-carbohydrate mixtures, described for the morning meal, were offered in 2.5 l of water. The commercial milk replacer was prepared from milk products (skim milk and whey powder). The composition of the milk replacer (% as fed) was: dry matter; 95.6, crude protein; 26.3, fat; 16.5, ash; 7.8, Mcal/kg; 4.7. Digestible energy was calculated using the following factors: 4, 9, 4 kcal/g for protein, fat and N-free extracts, respectively.

The calves were weighed at the beginning of the experiment and on days 10, 17 and 25. After weaning, they received a commercial concentrate (16% crude protein; 2 600 kcal/kg). Its consumption during the first 3 d was recorded for each calf. Later on, the calves were kept in 1 group; they received a concentrate (ad libitum) and hay (150 g/d), and their weights and age at slaughter were recorded.

On d 13–20 of the experiment, the faeces were collected in plastic bags, which were changed twice daily, and their contents frozen at -20 °C. On the same days, urine was collected in plastic bottles containing 200 ml of 5% sulfuric acid to prevent nitrogen loss. The pooled faecal samples of each calf were homogenized with water (1/1, bw/w) in a Waring Blender. Water intake was measured on the same days as the faeces and urine were collected.

In order to determine whether all liquid feed by-passed the rumen, iron oxide (2 g/kg dry food) was mixed with the diets and rumen liquor was obtained with a plastic tube connected to a suction bottle, before and 15 min after feeding on the 21st d of the experiment. The pH of the rumen liquor was determined immediately and the presence of red colour was recorded.

On d 22 of the experiment, blood samples were drawn from the jugular vein before the morning meal and at various intervals up to 120 min following the afternoon meal.

**Chemical analyses**

Dry matter, ash, total nitrogen, fat and starch in the feeds and the faeces, and total nitrogen in the urine were determined as described earlier (Ben-Asher et al, 1981). Blood glucose was determined by a glucose-oxidase method (Hestrin-Lerner and Ben-Yonah, 1963).

In vitro digestion of raw and heat-treated starch was determined by using pancreatic amylase (Type 1–A, Sigma Chemical Co, St Louis, Mo, USA) at a concentration of 0.5 mg/ml distilled water. The starch samples were prepared in phosphate buffer (pH 6.9, 0.1 mol/l) at a concentration of 0.5%. After incubation for 10 min at 37 °C, the amount of reducing sugars was determined with Summer reagent (Bernfeld, 1955).

**Statistical analysis**

Differences between means were assessed by Duncan's multiple range test (Duncan, 1955).

**RESULTS**

The body weight of ES calves increased linearly, from the beginning of the experiment, while that of G and US calves increased more slowly during the first 10 d. Later on, the growth of all groups was parallel (fig 1).

**Fig 1.** Body weights of calves fed milk replacer and a mixture of soy concentrate and glucose, (G), heat-treated starch, (ES) or untreated starch (US). Vertical lines = SEM.
There were no statistically significant differences in the performance of calves in the different groups up to weaning. However, the ES calves grew and utilized dietary energy and protein better than the other 2 groups (table I). For 3 d after weaning the ES calves consumed the highest amounts of dry concentrate feed. The calves were slaughtered when they reached a mean body weight of about 420 kg. The ES calves reached slaughter weight 23 d earlier than the G group ($P < 0.05$) and 11 d before the US group (NS).

Apparent digestibility of the dry matter was highest in the G group; no significant difference was observed between the ES and US groups. Expanded starch was digested better than raw starch ($P < 0.05$). Apparent digestibility of protein was similar in the G and ES groups and about 7% lower in the US group (NS). There were no statistically significant differences among the groups in fat or in ash apparent digestibility. Nitrogen retention was highest in the G group and lowest in the US group ($P < 0.05$). The ES group was intermediate and did not differ significantly from either group (table II).

Water intake of the calves was about 5.3 l/d or about 9 l/kg of dry matter intake and did not differ between the groups.

Diarrhoea was not observed and water concentration in the faeces was 16–20% in the different groups.

The pH of the rumen liquor was similar in all groups before the morning meal. Fifteen min following the meal, the pH was unchanged in the G group but dropped markedly in the ES one. In the US calves this drop was moderate (table III). The drop in rumen liquor pH occurred when the liquid feed did not completely by-pass the rumen. No marker was detected in the rumen liquor in any of the G calves. In the ES group the marker was detected in all the calves and in the US group in 2 out of 4 calves.

The faeces pH was lower in the calves fed diets containing US or ES compared with the G one, and the difference was statistically significant in the evening. There

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**Table I. Growth, food intake and utilization of calves fed milk replacer and a mixture of soy-concentrate and glucose (G), heat-treated starch (ES) or untreated starch (US) (means of 4 calves/group).**

<table>
<thead>
<tr>
<th></th>
<th>G</th>
<th>ES</th>
<th>US</th>
<th>SEM</th>
</tr>
</thead>
<tbody>
<tr>
<td>Initial body weight (kg)</td>
<td>40.9</td>
<td>40.1</td>
<td>40.7</td>
<td>2.35</td>
</tr>
<tr>
<td>Final body weight (kg)</td>
<td>46.8</td>
<td>48.4</td>
<td>47.0</td>
<td>2.31</td>
</tr>
<tr>
<td>Daily gain (g)</td>
<td>236</td>
<td>332</td>
<td>250</td>
<td>52</td>
</tr>
<tr>
<td>Intake:</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Digestible energy (Mcal)</td>
<td>66.9</td>
<td>67.5</td>
<td>67.2</td>
<td></td>
</tr>
<tr>
<td>Crude protein (kg)</td>
<td>3.58</td>
<td>3.58</td>
<td>3.58</td>
<td></td>
</tr>
<tr>
<td>Utilization:</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mcal/kg wt gain</td>
<td>11.3</td>
<td>8.1</td>
<td>10.7</td>
<td>1.65</td>
</tr>
<tr>
<td>Protein kg/kg wt gain</td>
<td>0.61</td>
<td>0.43</td>
<td>0.57</td>
<td>0.11</td>
</tr>
<tr>
<td>Dry concentrate intake</td>
<td>3 d after weaning (g)</td>
<td>456$^b$</td>
<td>887$^a$</td>
<td>541$^{ab}$</td>
</tr>
</tbody>
</table>

Means in the same raw with different superscript letters differ significantly ($P < 0.05$)
were no differences between the ES and US groups in this respect (table I).

Blood glucose levels before the morning meal were 80–90 mg/100 ml in all calves (fig 2). After the morning meal, blood glucose level in the G group increased markedly and reached 196 mg/100 ml 2.5 h post feeding. It resumed the basal level 6 h post feeding. In the ES and US groups, which were fed soy-starch mixtures in the morning meal, the changes in blood glucose levels were non-significant. In the second meal, 6 h after the morning meal, all groups received 400 g of milk replacer in addition to 120 g of soy-carbohydrate mixture. Blood glucose levels increased markedly in all groups; 60 min after the meal these levels were highest in the ES group and lowest in the US one (NS), they increased further 120 min following the meal in the G group and reached 200 mg/100 ml, similar to the peak reached in the morning. In the 2 groups that received soy-starch mixtures, the peaks were below 150 mg/100 ml.

Hyperglycemia in the G group was accompanied by higher secretion of glucose in the urine compared with the ES and US

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**Table II.** Apparent digestibility (%) of feed ingredients by calves fed milk replacer and a mixture of soy-concentrate and glucose (G), heat-treated (ES) or raw (US) starch (means of 4 calves/group).

<table>
<thead>
<tr>
<th></th>
<th>G</th>
<th>ES</th>
<th>US</th>
<th>SEM</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dry matter</td>
<td>92.4&lt;sup&gt;a&lt;/sup&gt;</td>
<td>88.1&lt;sup&gt;b&lt;/sup&gt;</td>
<td>86.0&lt;sup&gt;b&lt;/sup&gt;</td>
<td>1.48</td>
</tr>
<tr>
<td>Crude protein</td>
<td>82.2</td>
<td>82.2</td>
<td>76.9</td>
<td>3.07</td>
</tr>
<tr>
<td>Fat</td>
<td>91.1</td>
<td>90.7</td>
<td>87.4</td>
<td>2.96</td>
</tr>
<tr>
<td>Starch</td>
<td>—</td>
<td>95.4&lt;sup&gt;a&lt;/sup&gt;</td>
<td>87.0&lt;sup&gt;b&lt;/sup&gt;</td>
<td>2.50</td>
</tr>
<tr>
<td>Ash</td>
<td>81.7</td>
<td>80.8</td>
<td>81.8</td>
<td>2.39</td>
</tr>
<tr>
<td>Nitrogen retention</td>
<td>60.6&lt;sup&gt;a&lt;/sup&gt;</td>
<td>55.8&lt;sup&gt;ab&lt;/sup&gt;</td>
<td>48.0&lt;sup&gt;b&lt;/sup&gt;</td>
<td>3.14</td>
</tr>
</tbody>
</table>

Means in the same raw with different superscript letters differ significantly \((P < 0.05)\)

**Table III.** pH of rumen liquor and faeces of calves fed milk replacer and a mixture of soy-concentrate and glucose, heat treated starch or raw starch (means of 4 calves/group ± SE).

<table>
<thead>
<tr>
<th></th>
<th>Rumor liquor pH</th>
<th>Faeces pH</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>before meal</td>
<td>after meal</td>
</tr>
<tr>
<td>Glucose</td>
<td>7.3 ± 0.16</td>
<td>7.2&lt;sup&gt;a&lt;/sup&gt; ± 0.06</td>
</tr>
<tr>
<td>Treated starch</td>
<td>7.2 ± 0.23</td>
<td>4.9&lt;sup&gt;b&lt;/sup&gt; ± 0.18</td>
</tr>
<tr>
<td>Raw starch</td>
<td>7.3 ± 0.06</td>
<td>6.3&lt;sup&gt;ab&lt;/sup&gt; ± 0.43</td>
</tr>
</tbody>
</table>

Means in the same column with different superscripts differ significantly \((P < 0.05)\)
In vitro digestion of the treated starch by pancreatic amylase was about 3-fold higher than that of raw starch. The amount of reducing sugars released from 1 g of starch during 3 min of incubation was 250 and 82 mg, respectively.

DISCUSSION

The daily gains of the calves were 230–330 g. These gains were low since the calves received limited amounts of liquid feeds, according to the amounts fed to calves designed for beef production. However, while in practice calves are fed dry concentrate and hay in addition to milk replacers, the calves in the present experiment did not receive any dry feeds in order to evaluate the digestibility, utilization and glycemia obtained by the liquid diets.

Heat treatment of starch increased its in vitro availability to amylase. This was concurrent to about 10% improved digestibility of starch in vivo, confirming the previous results of Mathieu et al (1970). Improved digestibility of starch was accompanied by a 7% increase in protein digestibility and retention which was not statistically significant. A relationship between starch and protein digestion was also reported by Huber et al (1968) and by Thivend et al (1979). The lower pH of the faeces of calves fed starch could indicate that part of the starch escaped digestion and absorption in the small intestine and was fermented in the caecum or colon. Intense fermentation and thus high bacterial proliferation could cause an underestimate of protein apparent digestibility. A similar phenomenon was observed in kids fed milk replacers containing starch (Nitsan et al, 1985).

In young calves, milk by-passes the rumen and is diverted to the abomasum due to the closure of the oesophageal groove. Liquids other than milk may cause incomplete closure of this groove (Guilhermet et al, 1973), and part of the liquid feed may pass into the rumen. In the present experiment using iron oxide as a marker, it was found that inclusion of the starch-soy mixtures was accompanied by incomplete closure of the oesophageal groove. This was also evident from the lower pH in the rumen liquor of the ES calves compared to the other group as was reported for kids (Nitsan et al, 1990). The passage of small amounts of liquid feed into the rumen could enhance the development of this site. Indeed, during the first 3 d after weaning, when food intake was still recorded for each calf individually, the ES calves consumed far more dry concentrate than the
other groups (table I). This early effect on food intake probably persisted since the ES calves reached a slaughter weight of about 420 kg 23 d earlier than the G group, while the US calves were intermediate in this respect.

The post-feeding increase in blood glucose levels in the G group was quite similar to that reported by Thivend and Martin-Rosset (1971) for calves fed milk replacer containing a mixture of glucides. Blood glucose levels did not differ between the ES and the US groups and were considerably lower than in the G group, in accordance with Natrajan et al (1972).

Although the in vitro study showed a 3-fold increase in the digestibility of the treated, compared with the untreated raw starch, the post-feeding changes in blood glucose levels did not differ between the 2 groups. This could be related to the higher amounts of ES trapped in the curd, causing slower release to the small intestine and delayed evacuation from the gastrointestinal tract compared with the US (Coombe and Smith, 1974; Golan et al, 1990). When calves were fed a glucose-soy mixture, the high solubility of the glucose was accompanied by a rapid flow of glucose to the intestine, inducing a substantial rise in glycemia. It seems that post-feeding blood glucose levels may reflect the rate of absorption of soluble carbohydrates but are not a reliable measurement of total absorption or utilization of less soluble carbohydrates, such as various starches. The absorption of glucose from starches may be delayed due to their slower release from the abomasum or slower breakdown by amylase. Moreover, rapid absorption of glucose did not relate to a better growth rate.

In conclusion, suckling calves receiving 160 g of corn starch daily, or 250 g of starch per 1 kg of dry feed grew and utilized feed as well as their counterparts receiving a similar amount of glucose, without showing any ill effects or diarrhoea. Heat treatment improved the apparent digestibility of starch \( (P < 0.05) \). In the present study suckling calves absorbed about 152 g of ES and 139 g of US per d.

**REFERENCES**


Duncan DB (1955) Multiple range and multiple \( F \) tests. *Biometrics* 11, 1-44


