

Influence of some food additives on IgG plasma concentrations in newborn calves fed an immunoglobulin solution extracted from colostrum

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Summary — A lot of newborn ruminants have no free access to their mother's colostrum, due to various reasons. For them, colostro-replacers are now on the market. These products contain immunoglobulins but their capacity to pass through the intestinal wall is often doubtful. So, an experiment has been designed to test the influence of three supplements on the absorption of immunoglobulins extracted from colostrum by ultrafiltration. Fifty newborn calves were divided into five groups of ten. The first group received three meals of colostrum, exactly 2, 10, and 18 h after birth. According to the same kinetics, the four other groups received the same amount of immunoglobulins but previously extracted from colostrum and diluted in a saline solution. The diets given to the last three groups were supplemented with isobutyric acid, caseino-macropéptide and colostrum extract, respectively. Immunoglobulins extracted from colostrum were badly absorbed compared with control colostrum diet. In spite of liberal supply of immunoglobulins given soon after birth, the IgG plasmatic levels of the calves fed immunoglobulin solution remained below what is usually admitted as a good passive immunity. All the three additives were unable to improve the immunoglobulins' absorption.

calf / birth / colostrum / immunoglobulin

Résumé — Influence de quelques additifs sur les taux plasmatiques d'immunoglobulines G mesurés chez le veau nouveau-né après ingestion d'immunoglobulines extraites de colostrum.

De nombreux ruminants nouveau-nés n'ont pas accès au colostrum de leur mère, pour des raisons diverses. À leur intention, des substituts de colostrum sont maintenant commercialisés. Ils sont riches en immunoglobulines, dont l'aptitude à être absorbées par l'intestin du jeune animal, n'est pas solidement établie. Nous avons testé l'influence de trois additifs, sur l'absorption des immunoglobulines extraites du colostrum par ultrafiltration. Cinquante veaux nouveau-nés de race Holstein, ont été divisés en cinq groupes de dix. Le premier a reçu trois repas de colostrum 2, 10 et 18 h après la naissance, très exactement. Suivant une cinétique identique, les quatre autres ont reçu la même quantité d'immunoglobulines extraites du colostrum et remises en solution saline. Les régimes des trois derniers lots ont été supplémentés en acide isobutyrique, caséino-macropéptide ou extrait protéique de colostrum. Les immunoglobulines extraites du colostrum ont été beaucoup moins bien absorbées que les immunoglobulines restées dans leur milieu d'origine. Malgré une administration

abondante effectuée très précocement après la naissance, les concentrations plasmatiques en IgG obtenues sont restées inférieures au taux de 10 g/L qui est habituellement considéré comme nécessaire pour assurer la protection immunitaire du nouveau-né. Les trois additifs testés se sont montrés sans effet positif sur l'absorption des immunoglobulines.

veau / naissance / colostrum / immunoglobuline

INTRODUCTION

Due to the absence of immunoglobulin placental transfer, young ungulates are born agammaglobulinemic (Levieux, 1984) and, consequently, are highly sensitive to infections. To protect them, a sufficient amount of colostrum well provided with immunoglobulins should be given as soon as possible (Dardillat et al, 1978). However, for many newborn calves, lambs, kids or foals, several reasons may impair colostrum intake: insufficient production in case of multiple births, acute mastitis, lack of adequate maternal behaviour mostly at first parturition, etc. These difficulties seem to be smoothed out by giving frozen colostrum and more recently, colostrum replacers, which are essentially issued from the newly established colostrum industry. Nevertheless, it is important to be very cautious regarding these nutritional specialities because of low IgG plasma concentrations frequently obtained (Zaremba et al, 1993). Drying standard colostrum collected from farm to farm, leads to products which are often poorly provided with immunoglobulins because farmers are tempted to deliver not only first milking colostrum but also those from second or third milking. In order to preserve an adequate concentration of immunoglobulins, liquid reconstitution before supplying the newborn, with a reduced amount of water, is attractive. This is not suitable, resulting in a beverage too viscous to be delivered via a bucket or an artificial teat. Therefore, before drying, removal of the major part of fat and other proteins is needed. Extraction of immunoglobulins from colostrum, followed by concentration, may

be considered but according to Grongnet et al (1986), in such conditions, immunoglobulins appear to be poorly absorbed by the intestine of the newborn calf. That's why an experiment was made to test the ability of some food additives to improve the absorption efficiency of immunoglobulins extracted from colostrum. Three products were used. Two of them, isobutyric acid (IA) (Hardy, 1969) and freeze dried colostrum extract (FDCE) (Balfour and Comline, 1962) have already been favourably tested in related conditions. The third one, caseinomacropetide (CMP) could reduce gastric secretion (Chernikov et al, 1974), thus limiting abomasal denaturation of immunoglobulins.

ANIMALS, MATERIALS AND METHODS

Animals

Fifty newborn Holstein calves (41 male, 9 female) were separated at birth from their dams, housed in individual pens and affected, birth after birth, by systematic circular permutation (table I) to one of five treatments: colostrum, immunoglobulin solution either alone or supplemented with IA, CMP or FDCE. All the animals were born spontaneously, at term and without dystocia. Calving was often slightly accelerated by a mild traction exerted on the forelimbs of the animals. No chemical compound was used to facilitate parturition.

Diets

First milking colostrum were obtained from about 40 cows of the ENSAR dairy herd. Colostrum were frozen just after milking, all thawed, at the end of the calving season, pooled and frozen again, for 20% of the total amount, by aliquot fractions

Table 1. Body weight and sex of 50 newborn calves fed colostrum or immunoglobulin solution. *Poids vif et sexe de 50 veaux nouveau-nés alimentés avec du colostrum ou une solution d'immunoglobulines.*

<i>Diets</i>	<i>Body weight ($x \pm \text{sem/kg}$)</i>	<i>Male</i>	<i>Female</i>
Colostrum	42.7 \pm 2.2	8	2
Immunoglobulin solution (IS)	39.7 \pm 2.4	7	3
Immunoglobulin solution + isobutyric acid (0.5%) (IS + IA)	38.3 \pm 1.0	7	3
Immunoglobulin solution + caseinomacropéptide (0.4%) (IS + CMP)	43.4 \pm 1.4	9	1
Immunoglobulin solution + freeze dried colostrum extract (IS + FDCE)	39.7 \pm 2.2	10	0

of 1L. The immunoglobulin solution (IS) was prepared with the remaining 80%, according to figure 1. Diafiltration was stopped when immunoglobulin G (IgG) concentration reached the initial colostrum concentration, ie, 44 g/L. FDCE issued from the first permeate, was prepared during the same technological operations (fig 1), in order to collect all the low molecular mass soluble proteins, present in colostrum at the beginning. Before use, FDCE was added to immunoglobulin solution in a proportion suited to restore the initial colostrum concentrations.

Aliquot fractions of colostrum and immunoglobulin solution were thawed in a warm water bath and supplemented if necessary with IA, CMP or FDCE, just before feeding the animals. Exactly 2, 10 and 18 h after birth, the calves were fed either colostrum or one of the immunoglobulin solutions. The meals were three times delivered at the rate of 25 g/kg body weight and given in a bucket fitted with a teat at the bottom. In order to equalize the quantities of immunoglobulin ingested, in case of carefully recorded obvious inappetence, repeated pressure on the teat to force deglutition or oesophageal intubation were practised.

Measurements, analysis and statistics

Blood was collected in heparinised vials by puncture of a jugular vein just before each meal and

26 h after birth. It was immediately centrifuged and resulting plasma was frozen at -20°C till analysis. IgG concentrations were measured on a immunonephelometer Behring for plasma, colostrum and immunoglobulin solution according to Lebreton et al (1981) with an anti IgG (H + L) Behring rabbit immunoserum. During the extraction process, HPLC on gel filtration column (GF 250, DuPont) was used in order to control real time the evolution of immunoglobulin concentration in the solution. Rectal temperature was noted before each blood sampling. Anova and test were used for statistical analysis.

RESULTS

Weight, vitality, appetite and health of calves

Vitality was excellent during the whole experimental period except for two animals suffering from transient general atony after the second meal. They recovered about 4 h later.

Colostrum was very well accepted for the three meals but not the immunoglobulin solutions, whose acceptability depended on the nature of the additive. Highest refusal occurred with isobutyric acid: 30% for the first meal, 80% for the second and 70% for

First milking colostrum

- | 1) thawing
- ↓ 2) dilution 1/2 with 35°C water

SKIMMING =====>cream

↓
skim colostrum

- | -dilution 1/3 with 35°C water

PRECIPITATION OF CASEIN

- | -by lowering pH at 4.1 with HCl 6N
- | -decanting 10 mn
- | -restoration of 4.6 pH with NaOH 5N

SIPHONING and DE-SLUDGING by CENTRIFUGATION

====>Caseins

Whey from colostrum

CLARIFICATION BY FILTRATION

- | -on Seitz Supra EK frontal filter system

ULTRAFILTRATION =====> First permeate

- | -on Romicon PM 100 ultrafiltration system

First retentate

Second permeate ←— **ULTRAFILTRATION**

(to the waste)

- | on Romicon PM10 ultrafiltration system

DIAFILTRATION

- | -Constant volume batch process
- | by introduction of salted water solution
- | (KCl : 2.9 g/l ; NaCl : 1.27 g/l)
- | -on Romicon PM 100 ultrafiltration system
- | -elimination of most lactose, mineral salts,
- | α-lactalbumin and β-lactoglobulin.

Second retentate

FREEZE DRYING

Freeze-dried

Immunoglobulin solution (IS)

Colostrum extract (FDCE)

Fig 1. Extraction process of immunoglobulin from colostrum.
Procédé d'extraction des immunoglobulines à partir du colostrum.

Table II. Postnatal evolution of rectal temperature ($^{\circ}\text{C}$) of 50 newborn calves fed colostrum or immunoglobulin solution ($x \pm \text{SD}$).*Évolution postnatale de la température rectale ($^{\circ}\text{C}$) de 50 veaux nouveau-nés alimentés avec du colostrum ou une solution d'immunoglobulines (moyenne \pm écart type).*

Groups	Time after birth (h)			
	2	10	18	26
Colostrum	37.9 \pm 0.4	38.2 \pm 0.1	38.7 \pm 0.1	38.7 \pm 0.1
IS	38.0 \pm 0.4	38.0 \pm 0.3	38.2 \pm 0.2	38.6 \pm 0.2
IS + IA	37.6 \pm 0.2	38.0 \pm 0.1	38.2 \pm 0.2	38.3 \pm 0.1
IS + CMP	37.8 \pm 0.2	38.0 \pm 0.1	38.3 \pm 0.1	38.7 \pm 0.2
IS + FDCE	38.2 \pm 0.2	38.0 \pm 0.2	38.4 \pm 0.2	38.7 \pm 0.1

the last. For immunoglobulin solution alone, it was 30, 70, 40% and only 0, 30, 20% for CMP and FDCE, respectively. The strict rule used to affect the animals to the various treatments, led to slight and non-significant differences in average weight of the five groups and to slightly unbalanced sex-ratios.

Rectal temperature and IgG plasma concentrations

Rectal temperature (table II) rose slightly and significantly from the beginning and

stabilized thereafter. No significant differences were observed among treatments.

Before the first meal, IgG plasma concentrations (table III) were below the detection level of the nephelometer and considered equal to zero. For all treatments, feeding triggered an obvious increase which lasted during the whole experimental period except for IA group which exhibited the same values at 18 h and 26 h of life. The colostrum group differs strongly from all the other groups characterized by dramatically weak

Table III. Postnatal evolution of plasma IgG concentrations (g/L) of 50 calves fed colostrum or immunoglobulin solution ($x \pm \text{SD}$).*Évolution post-natale de la concentration plasmatique en immunoglobulines G de 50 veaux nouveau-nés alimentés avec du colostrum ou une solution d'immunoglobulines (moyenne \pm écart type).*

Groups	Time after birth (h)			
	2	10	18	26
Colostrum	0	5.8 \pm 2.6 ^A	11.0 \pm 3.7 ^A	12.8 \pm 4.4 ^A
IS	0	2.3 \pm 0.7 ^{Ba}	4.2 \pm 2.8 ^{Bab}	5.6 \pm 3.7 ^{Ba}
IS + IA	0	1.4 \pm 0.9 ^{Bb}	2.8 \pm 1.9 ^{Ba}	3.3 \pm 2.6 ^{Bb}
IS + CMP	0	1.9 \pm 1.4 ^{Ba}	4.2 \pm 2.6 ^{Bab}	4.9 \pm 2.8 ^{Bab}
IS + FDCE	0	2.1 \pm 1.2 ^{Ba}	4.9 \pm 1.6 ^{Bb}	5.6 \pm 2.8 ^{Ba}

For every time, values with different capital letters are significantly different at $P < 0.001$; values with different small letters are significantly different at $P < 0.05$.

Pour chaque temps, les valeurs avec des lettres majuscules différentes sont significativement différentes à $p < 0,001$; les valeurs avec lettres minuscules différentes sont significativement différentes à $p < 0,05$.

values: always lower than 6 g/L instead of 12.8 g/L in colostrum group 26 h after birth.

DISCUSSION

A concentration of 44 g/L IgG in the first milking colostrum pool used here is weak when compared with 100 g/L recorded by Leveux (1984). But his result was obtained in Charolais cows producing a scarce but highly concentrated colostrum. Today, in high-producing Holstein cows, 40 to 50 g/L may be considered as an average (Grongnet, unpublished data; Leveux, personal communication).

Good general vitality emphasizes that the immunoglobulin solution, except for its disappointing bad absorption efficiency and supplemented or not, was harmless, as opposed to the results obtained by Grongnet et al (1986). Similar evolution of rectal temperature in all groups shows that the low amount of energy provided by immunoglobulin solution did not impair thermoregulation.

Colostrum-fed animals exhibit maximal IgG plasma concentrations lower than those recorded in quite similar conditions by Grongnet et al (1986) but in agreement with Stott et al (1979), Gay et al (1983) and enough for ensuring a satisfactory protection. Taking into account such weak levels registered with the immunoglobulin solution, it can be said, first of all, that oesophageal intubation needed by frequent inappetence cannot impair so strongly the absorption mechanism. This particular way of colostrum administration has been systematically and successfully used by some authors (Al-Jawad and Lees, 1985). Birth weight differences can no more be involved: they are very small among groups and colostrum-fed animals are neither the heaviest nor the lightest. Besides, influence of birthweight on Ig absorption has always been recorded as slight or negligible (Bekele et al, 1992). It is the same for influence of sex (Villette et Leveux, 1981; Donovan et al, 1986; Bekele et al, 1992). Finally, the

composition of the solution seems to be the only factor involved in the insufficient absorption.

Additives were all ineffective. Isobutyric acid was even injurious, disagreeing with the results of Hardy (1969) who observed an improvement with this compound under quite similar experimental conditions. CMP is a 64-aminoacid peptide, released in the abomasum, following action of chymosin on κ -casein. In the preruminant calf, CMP enters rapidly the duodenum (Guilloteau et al, 1987). Vasilevskaya et al (1977) established, in the dog, that CMP injection restricted gastric secretions, previously stimulated by histamine or gastrin C-terminal tetrapeptide. As reduction of hydrochloric secretion, pepsin and other proteases would have a favourable effect on acquisition of passive immunity in mammals (Chernikov et al, 1974), addition of CMP to immunoglobulin solution was attractive but this hypothesis was not confirmed by the results presented here. FDCE was tested with reference to Balfour and Comline (1962). A positive effect was obtained by these authors with a low molecular mass protein, from colostrum origin but not identified. Our negative result seems to indicate that this component has a molecular mass below 10 000 kDa, level of cut-off of the half-permeable membrane here used to retain our FDCE (fig 1).

Colostrum has an antitryptic activity, exerting probably a protective effect on IgG in the gut. Grongnet et al (1986) have established that this beneficial property was preserved in an immunoglobulin solution prepared in the same way as described here. Therefore, large differences in absorption efficiency do not appear to be related to variations of antitryptic activity.

Many hypotheses can be advanced to explain these low IgG plasmatic levels. Lack of energy must be considered. Immunoglobulin solution was free of fat and lactose. Grongnet et al (1986) found that the addition of milk powder to such a solution im-

proved IgG plasma concentrations in newborn calves. Secondly, it must be taken into account that before joining blood, a great part of IgG is found in the lymphatic system (Kiryama, 1992), a way also used by lipids. This may suggest that IgG leaves the enterocytes via a mechanism involving lipids.

Colostrum is a complex biological medium still insufficiently known. Precise physico-biochemical conditions seem to be needed for an efficient IgG transport from intestinal lumen to lymph and blood. Additional research is still required to ensure valuable utilization of colostrum substitutes in field conditions.

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