Bioavailability of the flavonol quercetin in neonatal calves after oral administration of quercetin aglycone or rutin

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ABSTRACT

Polyphenols, such as flavonoids, are secondary plant metabolites with potentially health-promoting properties. In newborn calves flavonoids may improve health status, but little is known about the systemically availability of flavonoids in calves to exert biological effects. The aim of this study was to investigate the oral bioavailability of the flavonol quercetin, applied either as quercetin aglycone (QA) or as its glucorhamnoside rutin (RU), in newborn dairy calves. Twenty-one male newborn German Holstein calves were fed equal amounts of colostrum and milk replacer according to body weight. On d 2 and 29 of life, 9 mg of quercetin equivalents/kg of body weight, either fed as QA or as RU, or no quercetin (control group) were fed together with the morning meal. Blood samples were taken before and 0.5, 1, 1.5, 2, 2.5, 3, 4, 5, 6, 12, 24, and 48 h after feed intake. Quercetin and quercetin metabolites with an intact flavonol structure (isorhamnetin, tamarixetin, and kaempferol) were analyzed in blood plasma after treatment with glucuronidase or sulfatase by HPLC with fluorescence detection. Maximum individual plasma concentration was depicted from the concentration-time-curve on d 2 and 29, respectively. Additional blood samples were taken to measure basal plasma concentrations of total protein, albumin, urea, and lactate as well as pre- and postprandial plasma concentrations of glucose, nonesterified fatty acids, insulin, and cortisol. Plasma concentrations of quercetin and its metabolites were significantly higher on d 2 than on d 29 of life, and administration of QA resulted in higher plasma concentrations of quercetin and its metabolites than RU. The relative bioavailability of total flavonols (sum of quercetin and its metabolites isorhamnetin, tamarixetin, and kaempferol) from RU was 72.5% on d 2 and 49.6% on d 29 when compared

with QA (100%). Calves fed QA reached maximum plasma concentrations of total flavonols much earlier than did RU-fed calves. Plasma metabolites and hormones were barely affected by QA and RU feeding in this experiment. Taken together, orally administrated QA resulted in a greater bioavailability of quercetin than RU on d 2 and 29, respectively, and quercetin bioavailability of quercetin and its metabolites differed markedly between calves aged 2 and 29 d.

Key words: bioavailability, calf, flavonoid, quercetin, rutin

INTRODUCTION

Flavonoids are secondary plant metabolites occurring ubiquitously in all higher plants (Manach et al., 2004; Besle et al., 2010). They are known for their health-promoting properties (e.g., antioxidative and anti-inflammatory; Middleton et al., 2000; Nijveldt et al., 2001; Williams et al., 2004). Quercetin is one of the most abundant flavonoids and is present in high concentrations in onions, apples, and kale (Hertog et al., 1992; Nijveldt et al., 2001), and in low concentrations in milk (Besle et al., 2010; Bhagwat et al., 2013). In addition to their health-promoting properties, quercetin and its metabolites modulate the expression and activity of several metabolic key enzymes, and therefore might be involved in regulation of lipid and carbohydrate metabolism (Middleton et al., 2000; Gasparin et al., 2003; Kobayashi et al., 2010).

Newborn calves undergo tremendous immunological and metabolic changes after birth to adapt for extra-uterine life (Blum, 2006; Chase et al., 2008; Hammon et al., 2012) and colostrum management is one of the most important factors to support neonatal health and development (Godden, 2008; Hammon et al., 2012). Nevertheless, morbidity and mortality rates are still high during first weeks of life, and calves often suffer from diarrhea and respiratory disease (McGuirk, 2008; Mee, 2008; Uetake, 2013) as well as high levels of oxidative stress (Inanami et al., 1999; Gaál et al.,

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2006). Frequent problems on farms are the lack of high-quality colostrum availability and the insufficient colostrum supply (Quigley and Drewry, 1998; Godden, 2008). Colostrum and mature milk contain antioxidant agents to protect neonatal calves from oxidative stress (Lindmark-Månsson and Åkesson, 2000; Besle et al., 2010). Poor quality of colostrum is reflected by low concentrations of antioxidative substances. Thus, supplementing colostrum and milk at the beginning of lactation with the natural antioxidant agent quercetin may improve neonatal oxidative status. However, there is no information about the bioavailability (BV) of orally applied quercetin with colostrum and milk in newborn calves, although feeding industry already offers flavonoid-supplemented feed for dairy calves.

Studies on BV of quercetin from quercetin aglycone (QA) or its glucorhamnoside rutin (RU) in different monogastric species, such as rats (Manach et al., 1997), pigs (Ader et al., 2000; Cermak et al., 2003; Lesser et al., 2004), dogs (Reinboth et al., 2010), and humans (Erlund et al., 2000; Egert et al., 2008), showed marked differences to ruminant species (e.g., cows with intraruminal QA and RU application; Berger et al., 2012). These differences may be due to differences in gastrointestinal anatomy and physiology (Arts et al., 2004; Berger et al., 2012; Gohlke et al., 2013). Conversely, in newborn calves the forestomach system is just developing, thus newborn calves are functionally monogastrics (Drackley, 2008). The aim of the present study was to investigate the relative BV of quercetin after oral administration of QA or RU in calves during first month of life. We hypothesized that BV in calves depends on the form of application (QA and RU) and changes with age due to ontogenetic development and maturation of the gastrointestinal tract during first month of life. We further tested the hypothesis that quercetin application might affect metabolic and endocrine traits, especially concerning glucose metabolism in neonatal calves, because findings in literature pointed to impaired carbohydrate digestion and glucose absorption after flavonoid intake (Cermak et al., 2004; Tadera et al., 2006).

MATERIALS AND METHODS

Animals and Feeding

The experimental procedures were carried out according to the animal care guidelines and were approved by the relevant authorities of the State Mecklenburg-West Pomerania, Germany (LVL M-V/TSD/7221.3-2.1-019/10). Twenty-one male German Holstein calves were examined on d 2 and 29 of life. All calves were spontaneously born from multiparous

cows on neighboring farms and transported directly after birth to the experimental barn. Calves were kept in single boxes with straw bedding and had free access to water. Calves were fed twice daily (0700 and 1500 h) with a nipple bottle or nipple bucket. On the first 3 d of life calves received pooled colostrum obtained from milkings 1, 3, and 5 (d 1, 2, and 3 after parturition, respectively; Table 1) at amounts of 8% of BW on d 1 and 10% of BW on d 2 and 3 (Steinhoff-Wagner et al., 2011). From d 4 until 29, calves received milk replacer (150 g/L; SalvaLac MiraPro 45, Salvana Tiernahrung GmbH, Klein-Offenseth Sparrieshoop, Germany) at 12% of BW/d (Table 1). To ensure uptake of equal amounts of feed, refused amounts of colostrum or milk were tube-fed to calves. Milk intake was adapted to BW data once a week.

Colostrum or milk replacer was supplemented with chicken egg-derived immunoglobulins (Globigen Life Start 25%, EW Nutrition GmbH, Visbek, Germany) composed of 75% dextrose and 25% whole egg powder (10.75% CP, 10.50% crude fat, 0.10% crude fiber, and 2.50% ash), with high antibody titer against *Escherichia coli* type K 99, *Salmonella* Typhimurium and *Salmonella* Dublin, bovine rotavirus type G6 and G10, bovine coronavirus, *Cryptosporidium parvum*, and *Clostridium perfringens* serotype C. Immunoglobulins were added from d 2 to 6. Respective amounts of immunoglobulins fed twice daily were 40, 32, 24, 16, and 8 g/d.

From d 4 on calves had free access to pelleted concentrate (Kälber Start 18/3 pell., Vollkraft Mischfutterwerke GmbH, Karstädt, Germany; Table 1) and hay. Concentrate intake was measured daily after morning milk feeding. To avoid iron deficiency, calves received 600 mg of iron dextran subcutaneously (Ursoferran, Serumwerk Bernburg, Germany) on their first day of life. Navel disinfection was performed with 10% iodine solution (vet sept Lösung, Albrecht GmbH, Aulendorf, Germany) immediately after birth. Health status of calves was determined daily by measuring rectal temperature, heart rate, and respiratory rate, by evaluation of behavioral abnormalities, nasal discharge, respiratory sounds, fecal consistence, and by navel inspection.

Treatment and Blood Sampling

Calves were randomly assigned to 1 of 3 feeding groups (n = 7 per group) receiving either no flavonoids (control group; **CTRL**), 9 mg of QA/kg of BW (quercetin aglycone dihydrate, Carl Roth GmbH, Karlsruhe, Germany), or 18 mg of RU/kg of BW (rutin trihydrate, Carl Roth GmbH), each resulting in a dose of 9 mg of quercetin equivalents (**QE**)/kg of BW (30 μ mol of QE/kg of BW) on d 2 and 29 of life. Calves received the

Table 1. Composition and amounts of colostrum, milk replacer, and concentrate fed to calves

Item	d 1 colostrum ¹ milking 1	d 2 colostrum ¹ milking 3	d 3 colostrum ¹ milking 5	d 4 to 30 ²	
				Milk replacer ³	Concentrate ⁴
DM, g/kg	206.9	154.7	139.3	110.0	880.0
CP, g/kg of DM	597.4	390.4	333.1	201.8	180.0
Crude fat, g/kg of DM	169.2	240.5	274.9	190.9	43.0
Ash, g/kg of DM	48.3	55.6	56.0	73.6	74.0
Lactose, g/kg of DM	136.3	260.5	308.7	514.6	—

¹Colostrum was derived from multiparous cows and separately pooled from milkings 1, 3, and 5 after parturition, respectively.

²Hay was offered ad libitum except for days of bioavailability studies.

³Milk replacer (SalvaLac MiraPro 45, Salvana Tiernahrung GmbH, Klein-Offenseth Sparrieshoop, Germany), was composed of 45% spray skim milk powder, 35% sweet whey powder, 18% plant oil (palm, coconut, and soy oil), 1.7% Lys, 0.8% Ca, 0.75% P, and added with 50,000 IU of vitamin A, 4,000 IU of vitamin D₃, 200 mg of vitamin E (α -tocopherol acetate), 9 mg of Cu (glycerine-copper chelate), citric acid, sorbic acid, butylhydroxytoluene (BHT), and 1.2×10^9 cfu of *Enterococcus faecium* E1706.

⁴Concentrate (Kälber Start 18/3, Vollkraft, Mischfutterwerke GmbH, Karstädt, Germany) was composed of wheat gluten, wheat bran, grain mash, sugar beet pulp, rapeseed meal, oat, rye, molasses, linseed, oat bran, soybean meal extract, calcium carbonate, sodium chloride, 0.6% phosphorous, 0.2% sodium, 10,800 IU of vitamin A, 1,215 IU of vitamin D₃, 30 mg of vitamin E, 0.7 mg of I, 0.4 mg of Co, 54 mg of Mg, 81 mg of Zn, and 0.4 mg of Se per kg of DM.

whole dose of QA or RU during morning feeding, applying the QE suspension with a 10-mL syringe directly into the mouth.

The day before the study started, a catheter (CertoFix Mono 340, Braun Melsungen AG, Melsungen, Germany) was inserted into the calves' right jugular vein and blood samples were taken before (time point 0), and 0.5, 1, 1.5, 2, 2.5, 3, 4, 6, 8, 12, 24, and 48 h after feeding using S-Monovette tubes (Sarstedt AG & Co., Nümbrecht, Germany) containing lithium heparin (16 IU/mL of blood) for analysis of plasma flavonols. Additional blood samples, except at 1.5, 2.5, and 48 h, were taken for measurement of plasma concentrations of total protein, albumin, glucose, NEFA, urea, and lactate using tubes containing sodium fluoride and K₃EDTA (1.0 mg/mL fluoride and 1.2 mg/mL EDTA). Blood sampled into tubes containing dipotassium EDTA (1.8 mg/mL) was used for determination of insulin and cortisol plasma concentrations. Catheters were flushed with 10 mL of sodium chloride solution (0.9% sodium chloride, Braun Melsungen AG) after each blood sampling. Blood was immediately put on ice, centrifuged for 20 min at $1,500 \times g$ at 4°C, and plasma was stored at -80°C for flavonol analyses, and at -20°C for analyses of metabolites and hormones.

Analytical Methods

Flavonols were extracted from plasma as described by Egert et al. (2008) and concentrations of quercetin aglycone and its methylated (isorhamnetin, tamarixetin) and dehydroxylated (kaempferol) derivatives were analyzed by HPLC with fluorescence detection as previously described (Ader et al., 2000; Berger et al., 2012; Gohlke et al., 2013). Briefly, plasma samples

were thawed in a 40°C water bath. Then 980 μ L of plasma was acidified with 130 μ L of acetic acid (0.583 mol/L), spiked with 20 μ L of methanolic rhamnetin solution (internal standard, 1 mg/20 mL in methanol; Rotichrom HPLC, Carl Roth GmbH), and treated with a mixture of β -glucuronidase and sulfatase (from *Helix pomatia* Type H-1, with final activities of 7,300 and 130 U/mL for glucuronidase and sulfatase, respectively; Sigma-Aldrich Chemie GmbH, Dreieich, Germany) to cleave the ester bonds of glucuronides and sulfates. After incubation at 37°C for 1 h, 3 mL of acetone was added and the samples were centrifuged at $3,700 \times g$ for 45 min at 4°C. The supernatant was evaporated until dry. Residues were resolved in 200 μ L of methanol, and after 15 min in an ultrasonic bath 77.5 μ L of nanopure water and 22.5 μ L of hydrochloric acid (10 mol/L) were added. For flavonol analyses by HPLC, 30 μ L of the final solution was injected by a cooled (4°C) autosampler (AS-2057 Plus, Jasco Deutschland GmbH, Groß-Umstadt, Germany) onto a C-18 Kromasil 100 column (250 $\times$ 4 mm, particle size = 5 μ m, Jasco Deutschland GmbH) guarded by a precolumn (C-18 Inertsil ODS-2, 10 $\times$ 4 mm, particle size = 5 μ m, Jasco Deutschland GmbH). The eluent (flow rate: 1 mL/min) was composed of 0.025 mol/L of sodium dihydrogen phosphate (pH 2.4), acetonitrile, and methanol (68:27:5 vol/vol/vol). In a postcolumn reactor, the effluent was mixed with aluminum nitrate [flow rate = 0.4 mL/min, 1 mmol/L in methanol containing 7.5% (vol/vol) acetic acid] for derivatization, and fluorescence of the flavonol-aluminum complex was measured using a fluorescence detector (excitation wavelength = 422 nm, emission wavelength = 485 nm; FP920, Jasco Deutschland GmbH). Identification of peaks obtained was performed using the retention times of the standards, which were

prepared with pure flavonols and treated in the same way as samples. The detection limit of flavonols was ≤ 10 nmol/L and the recovery rate was $92 \pm 2\%$ (mean $\pm$ SE). Inter- and intraassay variances for quercetin were 7.2 and 0.5%. The sum of plasma concentrations of quercetin and its metabolites in plasma is referred to as total flavonols.

Plasma metabolites were analyzed spectrophotometrically (ABX Penta 400; Horiba ABX SAS, Montpellier Cedex, France) by the Clinic for Cattle (University of Veterinary Medicine, Foundation, Hannover, Germany) using the respective kits: albumin (no. A11A01664) and lactate (no. A11A01721; Horiba Europe GmbH, Hannover, Germany), NEFA (no. 434-91795; Wako Chemicals GmbH, Neuss, Germany), glucose (no. 553-230) and total protein (no. 553-412; MTI Diagnostics GmbH, Idstein, Germany), and urea (no. LT-UR 0050; Labor + Technik E. Lehmann GmbH, Berlin, Germany).

Plasma insulin concentrations were determined using an RIA as previously described (Vicari et al., 2008). Plasma cortisol concentrations were analyzed using a commercial ELISA kit validated for bovine plasma (EIA-1887; DRG Instruments GmbH, Marburg, Germany; Weber et al., 2013). Cross reactivities of the monoclonal antibody against corticosterone and progesterone were 45 and 9%, respectively, and to any further plasma steroids lower than 2%. Test sensitivity was 3.4 ng/mL; inter- and intraassay variations were 5.3 and 12.1%, respectively.

Colostrum was analyzed for DM, CP, crude fat, and ash content (Qualitätsprüfungs- und Dienstleistungsgesellschaft Mecklenburg-Vorpommern mbH, Güstrow, Germany) according to the Weender standard procedure (Naumann and Bassler, 2004), as shown in Table 1. Chemical composition of milk replacer and concentrates were according to the manufacturers' declarations (Table 1).

Calculations and Statistical Analyses

Quercetin measurements were corrected by baseline concentration (first blood sample, before QA or RU administration), then area under the curve (AUC) was calculated for quercetin and its metabolites (isorhamnetin, tamarixetin, and kaempferol) according to the trapezoidale rule between 0 and 24 h using GraphPad Prism 3 (GraphPad Software Inc., San Diego, CA, version 3.03). For single quercetin metabolites, AUC proportion relative to AUC of total flavonols (sum of quercetin, isorhamnetin, tamarixetin, and kaempferol) was calculated in addition. Maximum plasma flavonol concentrations ($C_{\max}$) and time after intake until $C_{\max}$ was reached ($T_{\max}$) were determined from individual

plasma concentration-time curves. Relative BV of quercetin from RU was calculated as

$$\text{Relative BV} = \text{AUC}_{\text{RU}} / \text{AUC}_{\text{QA}} \times 100\%.$$

Statistical analyses were performed using SAS/STAT software (SAS Institute, 2010). All data are presented as LSM $\pm$ SE. Body weight, concentrate intake, and pharmacokinetic parameters (AUC, $C_{\max}$, $T_{\max}$), as well as proportions of single metabolites on d 2 and 29, were analyzed by repeated measurement ANOVA using the Mixed procedure with group (levels: CTRL, QA, RU; for proportional calculations only QA and RU), repeated variable day (levels: d 2, 29), and group $\times$ day interaction as fixed effects in the model. Repeated measures on each calf were taken into account by using the repeated statement of the Mixed procedure with an autoregressive residual covariance structure. Plasma concentration of total flavonols as well as metabolite and hormone data were analyzed by repeated measurement ANOVA using the Mixed procedure, as described previously, with group (levels: CTRL, QA, RU), day (levels: d 2, 29), time changes within d 2 and 29 (several time points before and after feed intake on d 2 and 29, respectively), and respective interactions as fixed effects. Pairwise differences among treatments and time changes were tested by the Tukey-Kramer-test. Effects were considered significant if $P < 0.05$.

RESULTS

BW and Feed Intake

All calves were born spontaneously and healthy. Birth weight was 47.4 ± 1.0 kg. Body weight increased with time in all groups ($P < 0.01$) and was 65.5 ± 1.3 kg on d 29 without differences between groups. Milk intake was 3.8 ± 0.1 kg/d on d 1 (8% of BW), 4.8 ± 0.1 kg/d on d 2 (10% of BW), and 6.9 ± 0.1 kg on d 29. Concentrate intake was 1.59 ± 0.28 kg/calf and increased equally with time ($P < 0.01$) in all groups. Average daily gain from birth to d 29 was 612 ± 28 g/d without group differences.

BV Studies

Plasma concentrations of total flavonols of CTRL were close to or below the detection limit on d 2 and 29, respectively (Figure 1A, B). In QA- and RU-fed calves, plasma concentrations of total flavonols increased ($P < 0.05$) after flavonoid administration on d 2 and returned to baseline concentrations 48 h after administration (Figure 1A). On d 2, plasma concentrations of total

Figure 1. Plasma concentration-time-curves of total flavonols after oral administration of 9 mg of quercetin/kg of BW $\times$ day as quercetin aglycone ($\square$), rutin trihydrate (Δ), or no flavonoid ($\bullet$) on d 2 (A) and 29 (B) of life. Values are LSM $\pm$ SE; $n = 7$.

flavonols were greater at 8 and 12 h ($P < 0.01$), but were smaller at 24 h after flavonoid administration ($P < 0.01$) in QA- than in RU-fed calves.

On d 29, postprandial plasma concentrations of total flavonols increased far less distinctly ($P < 0.01$) when compared with d 2 (Figure 1A, B). Further, plasma concentrations of total flavonols increased until 8 h after administration ($P < 0.01$) in QA-, but not in RU-fed calves (Figure 1B). Plasma concentrations of total flavonols on d 29 tended to be greater ($P < 0.1$) in QA- than in RU-fed calves.

Compared with QA- (100%), RU-fed calves showed a relative BV of quercetin of 72.5% on d 2 and of 49.6% on d 29. On d 2, AUC of total flavonols, quercetin, and quercetin metabolites in QA- and RU-fed calves differed ($P < 0.05$) from CTRL, and were greater ($P < 0.001$) on d 2 than on 29 for both treatments (Table 2). On d 2, AUC of quercetin and isorhamnetin was greater ($P < 0.05$) and tended to be greater for total flavonols ($P < 0.1$), but was lower for tamarixetin ($P < 0.05$) in QA- than in RU-fed calves. On d 29, AUC of quercetin was greater ($P < 0.05$) in QA-fed calves than in CTRL.

On d 2, $C_{\max}$ of total flavonols, quercetin, and quercetin metabolites except kaempferol of QA- and RU-fed calves differed from CTRL ($P < 0.05$; Table 2), whereas on d 29 only $C_{\max}$ of total flavonols and quercetin of QA-fed calves differed ($P < 0.05$) from CTRL. Maximal plasma concentrations of total flavonols, quercetin, and isorhamnetin decreased ($P < 0.05$) or tended to decrease (kaempferol, tamarixetin; $P < 0.1$) in QA- and RU-fed calves from d 2 to 29, respectively. On d 2, $C_{\max}$

was greater ($P < 0.05$) for quercetin and isorhamnetin, but was lower for tamarixetin in QA- than in RU-fed calves.

Time to reach $C_{\max}$ in QA- and RU-fed calves was not affected by day, but indicated treatment effects for total flavonols, isorhamnetin, and tamarixetin (Table 2). For isorhamnetin, $T_{\max}$ was greater ($P < 0.05$) in RU- than in QA-fed calves on both days.

Relative distribution of single flavonols (quercetin, isorhamnetin, tamarixetin, and kaempferol) proportionately to total flavonols in plasma showed highest values for quercetin on both days and in both groups (QA and RU). Proportion of single flavonols changed with time, namely increased for quercetin ($P < 0.05$) but decreased for isorhamnetin and tamarixetin ($P < 0.05$) from d 2 to 29 (Figure 2A, B). On d 2, proportion of isorhamnetin was greater ($P < 0.05$) and proportion of tamarixetin was smaller ($P < 0.05$) in QA- than in RU-fed calves (Figure 2A).

Plasma Concentrations of Metabolites and Hormones

Plasma concentrations of total protein decreased ($P < 0.05$) from d 2 to 29 in CTRL and were greater on d 2 in CTRL than in QA-fed calves ($P < 0.05$; Table 3). Plasma concentrations of albumin increased ($P < 0.01$) and plasma concentrations of lactate and urea decreased ($P < 0.01$) from d 2 to 29 without any group effect (Table 3). Mean glucose and NEFA plasma concentrations decreased ($P < 0.05$) from d 2 to 29 in all groups, but concentrations did not differ among groups

Table 2. Pharmacokinetic parameters after oral administration of quercetin as quercetin aglycone (QA) or rutin trihydrate (RU) on d 2 and 29 of life; calves of the control group (CTRL) received no flavonoids

Item ¹	Day	Group ²			SE	ANOVA (<i>P</i> -value)		
		CTRL	QA	RU		Group	Day	Group × day
AUC, nmol/L × h								
AUC _{Total}	2	154 ^c	5947 ^a	4312 ^{ab}	395	0.001	0.001	0.001
	29	7.70	1336	662				
AUC _Q	2	102 ^c	3413 ^a	1967 ^b	228	0.001	0.001	0.001
	29	26.7 ^b	1090 ^a	535 ^{ab}				
AUC _K	2	31.9 ^b	152 ^a	159 ^a	24.4	0.02	0.001	0.05
	29	0.00	11.1	6.39				
AUC _I	2	0.00 ^c	1390 ^a	447 ^b	70.1	0.001	0.001	0.001
	29	0.00	67.5	23.9				
AUC _T	2	0.00 ^c	1070 ^b	1768 ^a	154	0.001	0.001	0.001
	29	0.00	58.3	19.6				
C _{max} ⁴ , nmol/L								
C _{max, Total}	2	13.9 ^b	407 ^a	330 ^a	35.9	0.001	0.001	0.001
	29	2.24 ^b	168 ^a	74.7 ^{ab}				
C _{max, Q}	2	10.7 ^c	238 ^a	129 ^b	19.3	0.001	0.001	0.03
	29	6.28 ^b	137 ^a	60.3 ^{ab}				
C _{max, K}	2	1.96 ^b	12.4 ^a	10.9 ^{ab}	2.07	0.01	0.001	0.16
	29	0.00	3.86	2.12				
C _{max, I}	2	0.00 ^c	94.5 ^a	39.6 ^b	5.85	0.001	0.001	0.001
	29	0.00	15.5	3.99				
C _{max, T}	2	0.00 ^c	77.1 ^b	163 ^a	15.6	0.001	0.001	0.001
	29	0.00	9.45	3.27				
T _{max} ⁵ , h								
T _{max, Total}	2	9.50	10.6	16.4	2.84	0.05	0.12	0.90
	29	6.49	6.21	10.4				
T _{max, Q}	2	6.65	8.86	11.7	2.77	0.31	0.47	0.76
	29	6.44	5.93	11.1				
T _{max, K}	2	10.0	9.67	12.3	4.01	0.93	0.49	0.76
	29	ND ⁶	8.00	8.00				
T _{max, I}	2	ND	11.4 ^b	19.3 ^a	2.56	0.001	0.88	0.09
	29	ND	5.90 ^b	24.0 ^a				
T _{max, T}	2	ND	11.1	16.6	3.57	0.01	0.63	0.18
	29	ND	7.63	24.0				

^{a-c}LSM in a row with different superscripts differ ($P < 0.05$).

¹Values are LSM with SE, $n = 7$ per group.

²Group: QA = 30 $\mu\text{mol}/(\text{kg of BW} \times \text{d})$ quercetin aglycone; RU = 30 $\mu\text{mol}/(\text{kg of BW} \times \text{d})$ rutin trihydrate; CTRL = no flavonol fed.

³AUC = area under the plasma concentration-time curve of flavonols from 0 to 24 h, calculated by subtracting baseline values (time point 0) from all other time points according to the trapezoidal rule; AUC_{Total}, AUC_Q, AUC_K, AUC_I, AUC_T = AUC of total flavonols (Total) or of the respective quercetin metabolites quercetin aglycone (Q), kaempferol (K), isorhamnetin (I), and tamarixetin (T), respectively.

⁴C_{max} = maximum plasma concentration of flavonols subtracted by basal plasma concentration before feed intake; C_{max, Total}, C_{max, Q}, C_{max, K}, C_{max, I}, C_{max, T} = maximum plasma concentration of total flavonols (Total), quercetin aglycone (Q), kaempferol (K), isorhamnetin (I), and tamarixetin (T), respectively.

⁵T_{max} = time until C_{max} was reached; T_{max, Total}, T_{max, Q}, T_{max, K}, T_{max, I}, T_{max, T} = time until C_{max, Total}, C_{max, Q}, C_{max, K}, C_{max, I}, C_{max, T} was reached, respectively.

⁶ND = Not definable.

(Table 3). Considering the day course of the plasma metabolites, plasma concentrations of glucose increased ($P < 0.01$), whereas NEFA decreased ($P < 0.01$) after feed intake in all groups on d 2 and 29, respectively (Figure 3A-D). The increase of plasma insulin concentrations after feed intake was more pronounced ($P < 0.05$) on d 29 than on d 2, but without group effects (Figure 4A, B; Table 3). Plasma concentrations of cortisol decreased ($P < 0.05$) after feed intake on both days with mean concentrations being greater ($P < 0.05$) on d 2 than on d 29 (Figure 4C, D; Table 3).

DISCUSSION

Without dietary supplementation of quercetin (CTRL group) plasma concentrations of total flavonols were close to or below the detection limit on d 2 and 29. However, small amounts of quercetin and its dehydroxylated metabolite kaempferol were found in blood plasma on d 2 and 29 in the control group. According to Besle et al. (2010), flavonoids are natural components of bovine milk and their concentration is affected by cow's diet. Although colostrum and milk

Figure 2. Proportion of single flavonols (quercetin, isorhamnetin, tamarixetin, and kaempferol) in plasma after oral administration of 9 mg of quercetin/kg of BW × day as quercetin aglycone (QA) and rutin trihydrate (RU) on d 2 and 29 of life. Values are LSM ± SE; n = 7.

replacer were not analyzed for flavonoid content in the present study, it can be assumed that colostrum is most likely the source for flavonoids found in CTRL calves at d 2. Apart from colostrum, feed components such as concentrates and hay may also contain flavonoids (Fraisie et al., 2007; Reynaud et al., 2010), which could explain measurable, albeit very low, plasma concentrations of quercetin and its metabolites at least in older calves (d 29).

In calves, quercetin was systemically available from both supplements, QA as well as RU. Interestingly, plasma values obtained for quercetin and its metabolites with an intact flavonol structure were greater on d 2 than on d 29, indicating better absorption or less excretion in younger animals. This may be explained by a different degree of gastrointestinal maturation and intestinal permeability on d 2 and 29, respectively (Blum, 2006). Thus, the epithelium in newborn animals

Table 3. Blood plasma concentrations of metabolites and hormones of calves after oral administration of quercetin as quercetin aglycone (QA), rutin trihydrate (RU), or no flavonoid (CTRL) on d 2 and d 29

Item ¹	Day	Group ²			SE	ANOVA (<i>P</i> -value)		
		CTRL	QA	RU		Group	Day	Group × day
Total protein, g/L	2	64.0 ^a	59.9 ^b	61.7 ^{ab}	0.98	0.03	0.001	0.07
	29	58.0	56.5	59.6				
Albumin, g/L	2	22.9	22.6	22.5	0.37	0.95	0.001	0.42
	29	26.1	26.5	26.3				
Lactate, mmol/L	2	1.68	2.37	2.65	0.27	0.31	0.001	0.04
	29	0.63	0.68	0.69				
Glucose, mmol/L	2	7.37	7.14	7.61	0.28	0.35	0.001	0.99
	29	6.58	6.40	6.79				
NEFA, μmol/L	2	298	281	298	13	0.87	0.001	0.65
	29	172	177	173				
Urea, mmol/L	2	3.80	3.75	3.55	0.26	0.91	0.001	0.82
	29	1.55	1.56	1.55				
Insulin, μg/L	2	2.38	1.79	2.41	0.65	0.12	0.08	0.29
	29	4.38	2.09	2.77				
Cortisol, nmol/L	2	67.2	62.9	77.9	7.0	0.43	0.001	0.67
	29	11.1	8.6	13.0				

^{a,b}LSM in a row with different superscripts differ (*P* < 0.05).
¹Values are LSM based on plasma concentrations in blood samples taken before and until 24 h after morning feeding on d 2 and 29, respectively; n = 7 per group.
²Group: QA = 30 μmol/(kg of BW × d) quercetin aglycone; RU = 30 μmol/(kg of BW × d) rutin trihydrate; CTRL = no flavonol fed.

Figure 3. Plasma concentrations of glucose (A, B) and NEFA (C, D) after oral administration of 9 mg of quercetin/kg of BW $\times$ day as quercetin aglycone (□), rutin trihydrate (Δ), or no flavonoid (●) on d 2 and 29 of life. Values are LSM $\pm$ SE; n = 7.

is characterized by vacuolated epithelial cells, which are not present in older animals (Bainter, 2002; Blum, 2006). This has possibly contributed to a greater absorption of quercetin from both sources (QA and RU) on d 2 compared with d 29 in calves. Furthermore, the reticulorumen and microbial activities might be already more developed in 29- than in 2-d-old calves, resulting in partial ruminal degradation of the flavonoids applied (Heinrichs and Jones, 2003; Guilloteau et al., 2009; Berger et al., 2012; Li et al., 2012). The BV of quercetin from both, QA and RU, especially on d 29 could probably be improved when encapsulated quercetin preparations will be used (Ding et al., 2014).

On both days investigated, feeding of QA resulted in greater plasma concentrations of total flavonols, quercetin, and its metabolites than feeding of RU, indicating QA as a better source of quercetin than RU with respect to the BV of quercetin. Markedly lower plasma concentrations of total flavonols were also seen in monogastric species after feeding of RU instead of QA at dosages comparable to the current study (Manach et al., 1997; Cermak et al., 2003; Reinboth et al., 2010). Thus, the present results obtained in calves are in principal accordance with studies on the BV of quercetin in monogastric species, such as rats (Manach et al., 1997), dogs (Reinboth et al., 2010), pigs (Cermak et al., 2003;

Figure 4. Plasma concentrations of insulin (A, B) and cortisol (C, D) after oral administration of 9 mg of quercetin/kg of BW $\times$ day as quercetin aglycone (□), rutin trihydrate (Δ), or no flavonoid (●) on d 2 and 29 of life. Values are LSM $\pm$ SE; n = 7.

Lesser et al., 2004), and humans (Erlund et al., 2000; Egert et al., 2008). As the newborn calf is considered as a functionally monogastric animal (Drackley, 2008), these findings are not surprising.

Concerning RU application, studies in monogastric species often indicate a delayed increase of quercetin and its metabolites in blood plasma (Erlund et al., 2000; Cermak et al., 2003; Reinboth et al., 2010). In the present study, maximal plasma concentrations of quercetin and its metabolites were measured much later after RU than after QA feeding, indicating different intestinal sites for absorption (Erlund et al., 2000). In RU-fed calves, quercetin and its metabolites,

at least on d 2, most likely derive from absorption in both the small and the large intestine. In addition, we found the same slow increase of quercetin and its metabolites in blood plasma after QA and RU feeding, but the increase stopped much earlier in RU- than in QA-fed calves. In general, the sugar moiety in flavonol molecules determines the absorption of flavonoids in organisms (Day et al., 1998; Hollman et al., 1999) and QA is supposed to already be absorbed in the small intestine. More complex glycosides, such as the glucorhamnoside RU, are absorbed in distal parts of the small intestine or in the colon of monogastric animals, where flavonol molecules are hydrolyzed by microorganisms.

Thus, the lack of rhamnosidase activity in the small intestine can be considered as one reason for delayed and lower absorption of RU than of QA (Erlund et al., 2000; Cermak et al., 2003). Our pattern of relative BV of total flavonols after RU feeding fits to relative BV of total flavonols in blood plasma after duodenal, but not after ruminal application in dairy cows (Berger et al., 2012; Gohlke et al., 2013), indicating a low ruminal function in our calves during first month of life.

Interestingly, pharmacokinetics of quercetin absorption after QA feeding markedly differed between calves and adult cattle (Berger et al., 2012; Gohlke et al., 2013) or monogastric animals (Erlund et al., 2000; Cermak et al., 2003; Reinboth et al., 2010). The typical peak of total flavonol plasma concentrations some minutes after QA administration was not seen in newborn calves. As a consequence, time for reaching $C_{\max}$ was much greater in calves than in adult ruminants and monogastrics. Comparable slow increases of quercetin and its metabolites as in newborn calves' blood plasma were only seen in rats (Manach et al., 1997). These differences in pharmacokinetics after QA feeding are difficult to explain, but a probable reason is the low release of milk protein into the duodenum due to abomasal casein clotting (Heinrichs and Jones, 2003; Guilloteau et al., 2009) and the potential binding of flavonoids to milk proteins (Gugler et al. 1975; Boulton et al., 1998; Janisch et al., 2004).

All measured metabolites and hormones were in physiological ranges and most of these findings were in accordance with previous results indicating postnatal growth, especially protein accretion, and development in breeding as well as suckling calves (Egli and Blum, 1998; Nussbaum et al., 2002; Schiessler et al., 2002). Conversely, except for total protein concentrations on d 2, we found no treatment effects by QA or RU feeding in these calves. Plasma protein concentrations were lower in QA-fed than in CTRL calves, a finding that was hard to explain because all calves received the same pooled colostrum with identical IgG content at first days of life. Postprandial changes of metabolites and hormones have been reported previously (Hadorn et al., 1997; Hammon and Blum, 1998), but systemic glucose metabolism was not affected by QA or RU feeding, as we have assumed from findings in literature (Cermak et al., 2004; Tadera et al., 2006).

In conclusion, BV of quercetin and its metabolites in newborn calves is more pronounced when quercetin is fed as QA than as RU. These findings go along with data on BV of quercetin in lactating cows with duodenal application (Gohlke et al., 2013) and in monogastric species such as rats, dogs, and pigs (Manach et al., 1997; Cermak et al., 2003; Reinboth et al., 2010). Greater BV of quercetin on d 2 than on d 29 of life

was probably due to reduced maturation status of the gastrointestinal tract in 2-d-old calves. However, effects of quercetin feeding on the antioxidative status should be investigated to validate health-protecting effects of quercetin feeding in neonatal calves.

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