



Review Article

Subacute ruminal acidosis in dairy herds: Microbiological and nutritional causes, consequences, and prevention strategies

Mawda E. Elmhadi ^a, Darien K. Ali ^b, Mawahib K. Khogali ^c, Hongrong Wang ^{a,*}^a Laboratory of Metabolic Manipulation of Herbivorous Animal Nutrition, College of Animal Science and Technology, Yangzhou University, Yangzhou 225009, China^b Department of Veterinary Preventive Medicine and Public Health, Faculty of Veterinary Medicine, University of Khartoum, Khartoum North, Sudan^c College of Animal Science and Technology, Yangzhou University, Yangzhou, 225009, China

ARTICLE INFO

Article history:

Received 23 August 2021

Received in revised form

25 November 2021

Accepted 13 December 2021

Available online 11 March 2022

Keywords:

Subacute ruminal acidosis

High-concentrate diet

Microbial diversity

Inflammatory and immune response

Prevention

ABSTRACT

Dairy cattle are frequently fed high-concentrate (HC) diets in modern intensive feeding systems, especially in the transition period. During this period, cows face many alterations that include hormonal changes and shifting to a lactating state. Switching to a HC diet that may disrupt the ruminal microbiota balance can lead to subacute ruminal acidosis (SARA). Moreover, the main factor shaping the rumen microbiota is dietary composition, especially the ratio of starch to fibrous carbohydrates. Feeding highly fermentable carbohydrate diets after adaptation to forage diets leads to a rumen fermentation rate that exceeds rumen absorption and buffering rates, resulting in a reduction in ruminal pH. As a result of Gram-negative bacterial cell lysis, an increase in harmful ruminal bacterial metabolites, including lipopolysaccharide, lactic acid, and histamine, is observed. The interactions between the host immune system and the ruminal microbiota play an essential role in many physiological processes and the development of the disorder. Progress in DNA sequencing and bioinformatics platforms provides new opportunities to investigate the composition of ruminal microbes and yields unique advances in understanding ecology of the rumen. Subacute ruminal acidosis is linked with a change in the ruminal microbiota structure and richness and with other metabolic disorders; such as rumenitis, milk fat depression, laminitis, and liver abscesses. Therefore, this review aims to explore a better understanding of the crosstalk between diet and microbiota in the prevalence of rumen acidosis and its consequences, which is crucial for control strategies such as feeding management, and supplementation with thiamine, prebiotics, and probiotics.

© 2022 Chinese Association of Animal Science and Veterinary Medicine. Publishing services by Elsevier B.V. on behalf of KeAi Communications Co. Ltd. This is an open access article under the CC BY-NC-ND license (<http://creativecommons.org/licenses/by-nc-nd/4.0/>).

1. Introduction

In the last 3 decades, global milk production has increased by more than 64%, from 538 million tonnes in 1989 to 883 million tonnes in 2019 (FAO, 2021). This improved milk production requires nutrients that cannot be met by forage alone (Hua et al., 2017). The

general practice in the modern intensive management system is feeding ruminants with massive amounts of high-concentrate (HC) diets. As a consequence, these animals suffer from metabolic diseases. In a dairy herd, one or several metabolic disorders may affect one out of every 2 cows (Ametaj, 2010). Subacute ruminal acidosis (SARA) is among the most crucial metabolic diseases in intensively fed cows, sheep, and goats (Howard, 1981; Huo et al., 2014).

The ruminant's gastrointestinal tract is an immune-dynamic organ continuously exposed to many microorganisms, toxins, and chemical stimuli. The complicated and varied microbial ecosystem is referred to as the microbiome or microbiota. Decades of studies have focused on the complex correlation between the host animal and its inhabitant microbiota, primarily to determine the variation in rumen bacterial communities with diet alterations that can lead to digestive disorders such as SARA. Consumption of a HC diet

* Corresponding author.

E-mail address: hrwang@yzu.edu.cn (H. Wang).

Peer review under responsibility of Chinese Association of Animal Science and Veterinary Medicine.



increases ruminal fermentation that produces short-chain fatty acids (SCFA), which causes a decrease in ruminal pH to a level below 5.6. The reduction in ruminal pH stimulates the Gram-negative bacteria to die and release lipopolysaccharide (LPS). When an animal fails to cope with the organic acid balance, SARA will occur (Herdt, 2000).

To date, the role of microbiota in SARA is not entirely understood. The functionality and capacity of gastrointestinal tract microorganisms to utilize the fractions of substrate resources are varied. Excessive microbiota richness, balance, and diversity are considered essential for promoting microbial stability through dietary challenge situations and allow the animals to utilize limited resources more effectively. This review outlines the current research performed on SARA to demonstrate the potential role of ruminal microbial alterations, consequences, and prevention strategies.

2. Subacute ruminal acidosis

The current definition of SARA is a ruminal pH reduction from 5.5 to 5.0 for 111 to 180 min per 24 h (Jaramillo-López et al., 2017). SARA is also known as a subclinical rumen acidosis or chronic acidosis. Lactic acidosis is a rare accidental pathology in which the ruminal ecosystem is completely disturbed. SARA is a predominant ruminant health and production problem that inflicts high economic losses on dairy herds by reducing dry matter intake, milk production, and profitability by increasing the loss of animals due to culling or death (Gozho et al., 2007). Field studies in Europe and the USA have estimated that SARA incidence is between 11% and 19% in cows in early lactation, and between 18% and 20% in cows in mid-lactation and at peak dietary dry matter intake (O'Grady et al., 2008; Enemark, 2008; Mao et al., 2013). Unlike acute ruminal acidosis, SARA is a more chronic status not associated with the increase in ruminal lactic acid. SARA is related to deleterious animal health problems, such as ruminal mucous epithelial damage, and decreased fibre degradation. Symptoms such as diarrhoea, poor body condition, weight loss, and reductions in dry matter intake and milk production appear (Lean, 2007; Plaizier et al., 2009).

SARA is associated with modifications in the ruminal microbial population and diet type (Calsamiglia et al., 2012). Previous evidence showed that feeding ruminants a 50% to 65% grain diet could induce SARA (Tao et al., 2014a, 2014b). Another important cause of SARA is related to the fibre present in the diet, which can occur in excessive amounts of long particles, or be present at sufficient levels, but the long or inedible particles cause the animal to eat a portion of the feed in which the physically adequate fibre is low (Krause and Oetzel, 2006; Oetzel, 2007). Feeding fibre and grain diets separately appears to be a high risk for inducing SARA compared with feeding as mixed diets (Krause and Oetzel, 2006). In addition, animals fail to accommodate abrupt changes in the diet. Ruminal bacterial populations and papillae need more time to adapt to the digestion of HC diets and the absorption of the resultant high levels of volatile fatty acids (VFA) (Ghaffari et al., 2017).

In healthy cows, ruminal bacteria ferment starch into VFA, glucose, and lactic acid. Then, VFA and lactic acid are immediately absorbed by the ruminal wall papillae and enter the blood vessels, where they can be used in milk production. However, under acidic conditions, VFA and lactic acid accumulate in the rumen and contribute to acidosis. Furthermore, the rapid release of glucose in the rumen has subsequent negative consequences: (1) the growth of starch-digesting bacteria such as *Streptococcus bovis* (*S. bovis*) increases lactate production, (2) rapid growth with further domination of Gram-positive bacteria and defaunation killing of Gram-negative bacteria (e.g., *Fibrobacter succinogenes*, *Ruminococcus flavefaciens*) lead to the release of endotoxins such as LPS, and (3)

the rumen osmolality and the accumulation of VFA and lactic acid increase.

3. Ruminal microbial changes associated with subacute ruminal acidosis

Until recently, traditional culture-based methods have been used for exploring the ruminal microbiome, but many microorganisms cannot be cultured. Moreover, quantitative polymerase chain reaction (PCR) techniques, which are based on a specific organism rather than all microbes, are limited. At the beginning of the 21st century, a series of DNA fragmentation procedures, such as terminal restriction fragment length polymorphism, became available for studying microbial diversity (Ghaffari et al., 2017).

The ruminal microbiota includes bacteria, archaea, ciliate protozoa, and fungi. A fibre-rich nutrient and a steady microbial population are essential for maintaining ruminant health. Various investigations have been carried out on ruminal microbiota changes related to ruminal acidosis, which have indicated that the most indispensable microbiotas are bacteria and ciliated protozoa that enhance the grain fermentation rate (Nagaraja and Titgemeyer, 2007). In SARA, the number of protozoa begins to decrease, followed by a further reduction in the number of Gram-negative bacteria (Howard, 1981), whereas Gram-positive bacteria tend to increase (Petri et al., 2013).

3.1. Ruminal bacteria

The ruminant gastrointestinal tract accommodates more than 5,000 bacterial species, which changes according to diets, feeding approaches, and geographical regions (Chaucheyras-Durand and Ossa, 2014; Khafipour et al., 2016). Feeding cows with a SARA induction diets is related to a decline in the ruminal bacterial richness and diversity (Ametaj, 2010; Khafipour et al., 2009; Mao et al., 2013; Plaizier et al., 2017; Hua et al., 2017). Many investigations have been conducted on ruminal microbial alterations related to SARA (Table 1). In the rumen, the most dominant bacterial phyla are Bacteroidetes and Firmicutes (Kong et al., 2010; Khafipour et al., 2016). An earlier study assessed the alterations in the ruminal bacterial communities of dairy cows following adaptation to SARA, which showed that a SARA induction diet decreases Bacteroidetes but increases Firmicutes (Mao et al., 2013). This result agrees with the results of Mao et al. (2016) and Hua et al. (2017) on the influence of HC feeding on the reduction in Bacteroidetes and the increase in Firmicutes. Bacteroidetes are the primary degraders of complex polysaccharides in the plant cell walls. However, a HC diet raises the ratio of Firmicutes to Bacteroidetes, which is undesirable (Khafipour et al., 2016). In contrast, Hook et al. (2011a), Hua et al. (2017), and Zhang et al. (2017) found that the most dominant bacteria at the phylum level were Firmicutes and Bacteroidetes when feeding HC diets to ruminants. This difference in Bacteroidetes and Firmicutes abundance may be due to the resilience of the bacteria to dietary changes, the susceptibility of the ruminants, and/or differences in the starch content in the diet.

The increased level of concentrate directly reduced the relative dominance of the cellulolytic bacteria of the genus *Fibrobacter* (Zhang et al., 2017; Mickdam et al., 2016) and *Ruminococcus* (Mickdam et al., 2016). In contrast, SARA induction is associated with a *Ruminococcus* abundance (Mao et al., 2013; McCann et al., 2016). However, the relative abundance of *Megasphaera elsdenii* (*M. elsdenii*), which is a lactic acid utilizer, was shown to be increased in mild grain-induced SARA as reported by Khafipour et al. (2009), in vitro severe SARA induction as reported by Mickdam et al. (2016), and in SARA induced using pellets of ground wheat and barley as reported by Plaizier et al. (2017). In contrast,

Table 1
Changes in the ruminal bacteria associated with the subacute ruminal acidosis (SARA).

Model	SARA induction diet	Rumen bacteria change	16S rRNA region and sequencing platform	Citation
Cow	<ul style="list-style-type: none"> Grains Alfalfa pellets 	Bacteroidetes were decreased with SARA, whereas <i>Megasphaera elsdenii</i> was found to dominate mild grain-induced SARA. There was a significant decrease in bacterial species variety in mild grain-induced SARA. Moreover, high-grain feeding increased the communities of ruminal pathogenic <i>Escherichia coli</i> and <i>Clostridium perfringens</i> .	V1–V2 (TRFLP and real-time PCR)	Khafipour et al. (2009)
Cow	<ul style="list-style-type: none"> Hay (week 0) High concentrate (HC) diet (week 1 to 3) Hay (week 4 to 6) 	Bacterial density was significantly reduced in the rumen solids with a HC diet. The most common bacteria were <i>Lactobacillus</i> spp. and <i>Streptococcus bovis</i> .	Real-time PCR	Hook et al. (2011a)
Cow	<ul style="list-style-type: none"> HC diet 	The percentages of Proteobacteria and Bacteroidetes were decreased, whereas those of Firmicutes and Actinobacteria were increased.	V1–V3 (454 pyrosequencing)	Mao et al. (2013)
Goat	<ul style="list-style-type: none"> HC diet 	SARA showed a reduction in Bacteroidetes and a rise in Firmicutes.	454 pyrosequencing	Huo et al. (2014); Mao et al. (2016)
Cow	<ul style="list-style-type: none"> Wheat/barley pellet 	Relative dominance of the genera <i>Prevotella</i> , <i>Ruminococcus</i> , <i>Streptococcus</i> and <i>Lactobacillus</i> within the ruminal sample liquid fraction was detected, while Firmicutes was the abundant phyla in the solid fraction.	V4 (MiSeq Illumina platform)	McCann et al. (2016)
In vitro	<ul style="list-style-type: none"> HC diet 	Under a severe SARA situation, the relative dominance of the fibrolytic <i>Ruminococcus albus</i> , <i>Fibrobacter succinogenes</i> , and <i>Ruminobacter amylophilus</i> was reduced. In contrast, the relative abundances of <i>Prevotella</i> , <i>Lactobacillus</i> group, <i>M. elsdenii</i> and <i>Entodinium</i> spp. were increased.	Real-time quantitative PCR	Mickdam et al. (2016)
Goat	<ul style="list-style-type: none"> HC diet 	The level of Firmicutes and Bacteroidetes were increased. In addition, a notable change in the Cyanobacteria and Verrucomicrobia were demonstrated.	V3–V4 (454 pyrosequencing and RT–PCR)	Hua et al. (2017)
Cow	<ul style="list-style-type: none"> Pellets of ground wheat and barley Pellets of ground alfalfa 	The dominance of the ruminal Bacteroidetes and Tenericutes was decreased by a diet of pellets of ground wheat and barley but not by pellets of ground alfalfa. Only pellets of ground wheat and barley could increase ruminal <i>M. elsdenii</i> . Additionally, both challenges reduced the dominance of <i>S. bovis</i> .	V1–V3 (pyrosequencing and qPCR)	Plaizier et al. (2017)
Heifer	<ul style="list-style-type: none"> HC diet 	The Bacteroidetes and Firmicutes were significantly increased, and the relative abundance of cellulolytic bacteria (<i>Fibrobacter</i> and <i>Succinimonas</i>) was reduced.	V3–V4 (Illumina MiSeq sequencing)	Zhang et al. (2017)

M. elsdenii was shown to decrease in the study of Wang et al. (2015). SARA induction is dominated by *Prevotella* (Khafipour et al., 2009; McCann et al., 2016; Mickdam et al., 2016), which is known for the utilization of starch and protein. The most common bacteria discovered in the cow's rumen related to SARA are starch-utilizing bacteria such as *Lactobacillus* spp. and *S. bovis* (Hook et al., 2011a; McCann et al., 2016; Wang et al., 2015). Moreover, feeding a HC diet was shown to increase the pathogenic *Clostridium perfringens* and *Escherichia coli* (Khafipour et al., 2016). These increases were limited when increased grain feeding did not lead to severe rumen and hindgut acidosis, indicating that *C. perfringens* and *E. coli* might be factors contributing to severe rumen and hindgut acidosis onset.

3.2. Ruminal archaea

A study of nonlactating dairy cattle fed a HC diet to induce SARA was conducted to assess alterations in methanogen (methane-producing archaea) diversity and community structure (Hook et al., 2011b). These authors found no significant influence of feed on the ruminal methanogen density. This is in line with the results investigating the effect of an increased dietary fibre to grain concentration ratio on the archaeal community (Zhang et al., 2017). These authors found that a different dietary fibre to grain concentration ratio did not affect the archaeal communities. This is in contrast to the study that examined the influence of increased dietary grain (0%, 25%, and 50% maize grain) on archaea in goats, in which HC feeding reduced the richness of the archaeal community and increased the methanogen community (Mao et al., 2016). Ruminal archaeal changes are illustrated in Table 2.

3.3. Ruminal protozoa

An alteration in rumen protozoa density was determined in a study on nonlactating dairy cows transitioned to a HC diet (Hook

et al., 2011b). SARA was induced during week 1 of the experiment. Up through week 3, there was a significant increase in the abundance of protozoa in the ruminal solids and fluid, followed by a significant decrease in ruminal fluid protozoa when the cattle transitioned back to a hay diet in week 6. Similarly, Mao et al. (2016) examined the influence of increased dietary grain (0%, 25%, and 50% maize grain) on goat ruminal protozoa and observed that HC feeding increased the ciliate community. In contrast, feeding lactating cows with diet-induced SARA reduced the ruminal protozoa diversity (Ishaq et al., 2017). Increasing the concentration level linearly reduced the relative dominance of ciliates, namely, *Polyplastron* and *Ostracodinium*, and linearly increased the relative dominance of *Entodinium* (a nonfibrous carbohydrate degrader) (Zhang et al., 2017). Ruminal protozoal changes are illustrated in Table 3.

3.4. Ruminal fungi

High-concentrate diet feeding decreased anaerobic fungal density in male goats (Mao et al., 2016) but increased fungal diversity in dairy cows (Ishaq et al., 2017). The predominant anaerobic fungi were *Neocallimastigomycota* and *Ascomycota*, whereas 4 diets with different dietary forage to concentrate ratios did not affect the density and diversity of anaerobic fungal communities in heifers (Zhang et al., 2017). The anaerobic fungal resistance in the 4 treatment diets might be due to the coexistence of HC and high forage diets among the treatments, which balance cellulose and carbohydrate degradation. Ruminal fungal changes are illustrated in Table 4.

4. Inflammatory and immune responses associated with subacute ruminal acidosis

The lower ruminal pH may trigger the cell lysis of Gram-negative bacteria that is associated with the release of free LPS.

Table 2
Changes in the ruminal archaea associated with the SARA.

Model	SARA induction diet	Rumen archaeal change	16S rRNA region and sequencing platform	Citation
Cow	<ul style="list-style-type: none"> • Hay (week 0) • High concentrate (HC) diet (week 1 to 3) • Hay (week 4 to 6) 	No significant influence on the ruminal methanogen density.	Quantitative real-time PCR	Hook et al. (2011b)
Goat	<ul style="list-style-type: none"> • Maize grain 	The richness of the archaeal community was decreased, and the richness of the methanogen community was increased.	454 pyrosequencing	Mao et al. (2016)
Heifer	<ul style="list-style-type: none"> • HC diet 	There were no effects on the communities of archaea.	V3–V4 (Illumina MiSeq sequencing)	Zhang et al. (2017)

Table 3
Changes in the ruminal protozoa associated with the SARA.

Model	SARA induction diet	Rumen protozoal change	18S rRNA region and sequencing platform	Citation
Cow	<ul style="list-style-type: none"> • Hay (week 0) • High concentrate (HC) diet (week 1 to 3) • Hay (week 4 to 6) 	There was a significant increase in the number of protozoa present in the ruminal fluid and solids in week 3 and a significant decrease in protozoa in the ruminal fluid in week 6.	Quantitative real-time PCR	Hook et al. (2011b)
Heifer	<ul style="list-style-type: none"> • HC diet 	The relative abundance of ciliates, namely, <i>Polyplastron</i> , and <i>Ostracodinium</i> , was reduced, and the relative abundance of <i>Entodinium</i> was increased.	Illumina MiSeq sequencing	Zhang et al. (2017)
Goat	<ul style="list-style-type: none"> • Maize grain 	The number of ciliates in the protozoal community was increased.	454 pyrosequencing	Mao et al. (2016)
Cow	<ul style="list-style-type: none"> • HC diet 	The abundance of ruminal protozoa was reduced.	V3–V4 (Illumina MiSeq sequencing)	Ishaq et al. (2017)

Table 4
Changes in the ruminal fungi associated with the SARA.

Model	SARA induction diet	Rumen fungal change	ITS1 region and sequencing platform	Citation
Goat	<ul style="list-style-type: none"> • Maize grain 	The density of the anaerobic fungi was decreased.	454 pyrosequencing	Mao et al. (2016)
Cow	<ul style="list-style-type: none"> • High concentrate (HC) diet 	The fungal diversity was increased.	Illumina MiSeq sequencing	Ishaq et al. (2017)
Heifer	<ul style="list-style-type: none"> • HC diet 	Four different dietary ratios of forage to concentrate did not affect the communities of anaerobic fungi.	Illumina MiSeq sequencing	Zhang et al. (2017)

Lipopolysaccharide is a portion of the Gram-negative bacterial outer membrane cell wall that acts as an immunogenic compound (Bilal et al., 2016; Ghaffari et al., 2017). Moreover, free ruminal LPS can also result from bacterial cell lysis due to excessive autolytic enzymes that enhance bacterial growth during the rapid growth phase (Gozho et al., 2007; Plaizier et al., 2009). Increased ruminal permeability may be an adaptive response to HC diets to increase SCFA uptake (Zebeli and Metzler-Zebeli, 2012) or acidification or hyperosmolality (Emmanuel et al., 2007; Schweigel et al., 2005). Depending on the failure of the ruminal epithelial barrier in decreased ruminal pH conditions, LPS may be translocated into the bloodstream and promote systemic inflammation by stimulating the release of proinflammatory cytokines, such as tumour necrosis factor- α , interleukin-1 β , and interleukin-6, which may be a key causative factor in SARA (Xia et al., 2020). In addition, there is an increase in acute-phase proteins such as serum amyloid A, haptoglobin, LPS-binding protein, C-reactive protein, and alpha-1-acid glycoprotein in peripheral blood (Eckel and Ametaj, 2016; Gozho et al., 2007; Zebeli and Metzler-Zebeli, 2012). Other harmful compounds, such as biogenic amines and ethanol, may be produced during SARA (Ametaj, 2010). Histamine is a biogenic amine produced during SARA that is either absorbed from the gut or generated endogenously through inflammation and has been suggested to play a crucial role in laminitis progression.

Ruminal and intestinal epithelia, in addition to their functions as a selective barrier, help immune functions by their interactions

with gut-associated lymphoid tissue (Goto and Kiyono, 2012). The microbial effects on mucosal functions and gut-associated lymphoid tissue occur through linear microbial component interactions with receptors and by-products of symbiotic microbiota, including SCFA (Brestoff and Artis, 2013). In an investigation, isolated rumen and colon tissue from steers showed that LPS and reduced pH worked synergistically to lower barrier function (Emmanuel et al., 2007). Once the epithelium is breached, gut-associated lymphoid tissue cells may respond by triggering local inflammation and altering the generation of cytokines; this may further enhance permeability, potentiate colonization by pathogenic organisms, increase the transfer of bacteria and toxins across the epithelium, and increase the inflammatory response (Kurashima et al., 2013; Mani et al., 2012).

5. Subacute ruminal acidosis consequences

Potential SARA consequences that may occur are rumenitis, milk fat depression (MFD), laminitis, and liver abscesses (Lean, 2007; Plaizier et al., 2009), which are illustrated in Fig. 1.

5.1. Rumenitis

Rumenitis is a common consequence of rumen acidosis (Enemark et al., 2002). The pathogenesis of rumenitis involves LPS resulting from rumen acidosis over-activating the nuclear factor-

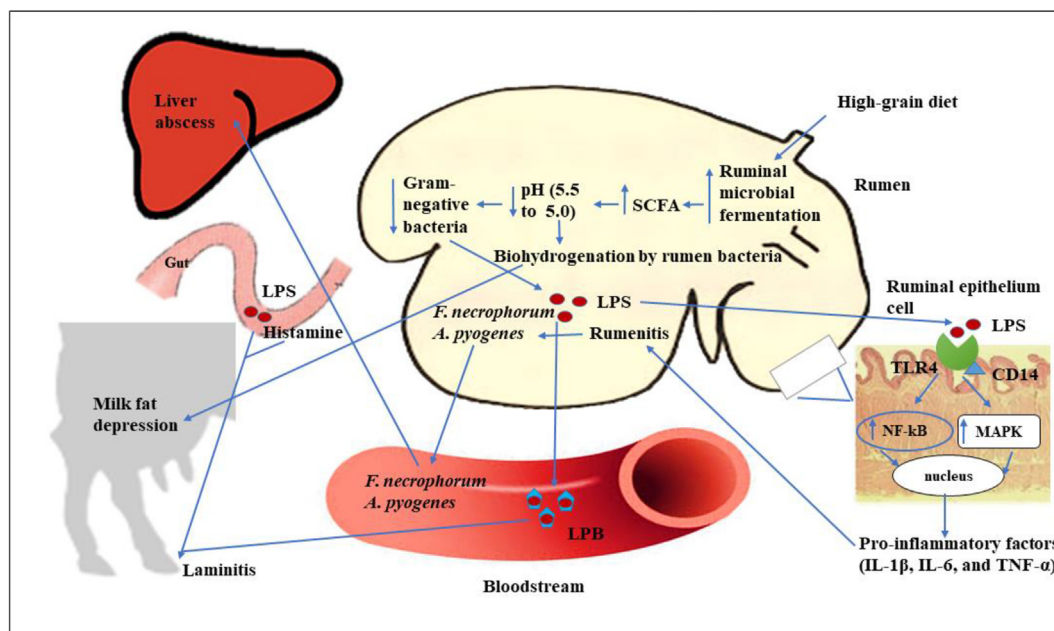


Fig. 1. The relationship between ruminal microbiota changes associated with subacute ruminal acidosis and other digestive disorders. The reduction in ruminal pH due to high-concentrate feeding alters Biohydrogenation by the ruminal bacteria, which leads to milk fat depression. In addition, the decreased ruminal pH stimulates rumenitis. Ruminal lesions resulting from acidosis influence *Fusobacterium necrophorum* and *Arcanobacterium pyogenes* entering the bloodstream, which are trapped and colonize in the liver, causing abscesses. Laminitis may also occur due to histamines and LPS. SCFA = short-chain fatty acids; LPS = lipopolysaccharide; LPB = lipopolysaccharide-binding protein; TLR-4 = Toll-like receptor 4; CD14 = coreceptor cluster of differentiation 14; NFκB = nuclear factor-kappa B; MAPK = mitogen-activated protein kinase; IL-1β = interleukin-1β; IL-6 = interleukin-6; TNF-α = tumour necrosis factor-α.

kappa B and mitogen-activated protein kinase inflammatory pathways in the ruminal epithelium. Consequently, this significantly increases the synthesis and expression of proinflammatory cytokines, thereby partly inducing rumenitis (Zhao et al., 2018).

5.2. Milk fat depression

Milk fat depression is a reduction in milk fat yield or percentage in response to highly fermentable diets and diets with high polyunsaturated fatty acids (Weimer et al., 2010). A depression in the milk fat concentration in cows induced by SARA has been demonstrated, due to alterations in ruminal fermentation patterns (Kleen et al., 2003). Milk fat depression illustrates the interactions among dietary feed, gastrointestinal microbiota, and tissue physiology (Pitta et al., 2018). Dietary components, particularly alterations in neutral detergent fibre and starch concentrations in the diet, can affect ruminal microbiota by modifying the ruminal environment (Lourenço et al., 2010).

During microbial fermentation in the rumen, polyunsaturated fatty acids undergo biohydrogenation to form saturated fatty acids with less toxic to microbes (Lourenço et al., 2010). Elevated concentrations of polyunsaturated fatty acids are considered toxic to rumen microbes because they form complexes with the cell walls of ruminal microbes and interfere with ruminal fermentation. Alterations in biohydrogenation pathways due to higher concentrations of polyunsaturated fatty acids and starch in the diet can lead to MFD.

Diet-induced MFD in dairy cows is commonly associated with alterations in the rumen microbe community due to the accumulation of VFA. Furthermore, it is associated with an increase in *S. bovis*, *M. elsdenii*, and *Selenomonas ruminantium* (Rico et al., 2015), an increase in specific genus levels of Actinobacteria and Firmicutes, such as unclassified Lachnospiraceae, *Butyrivibrio*, *Bulleidia*, and *Coriobacteriaceae* (Pitta et al., 2018), and a rise in the

relative dominance of *Prevotella* (Zeng et al., 2019). In contrast, a reduction in the dominance of *Prevotella bryantii*, *Fibrobacter succinogenes*, and the *Butyrivibrio fibrisolvens*/*Pseudobutyrvibrio* group (Rico et al., 2015), a reduction in the members of Bacteroidetes and Fibrobacteres (Pitta et al., 2018), and a decline in the relative abundance of unclassified Lachnospiraceae, *Oribacterium*, Veillonellaceae and *Pseudobutyrvibrio* in the ruminal fluid tend to occur (Zeng et al., 2019). In addition, the total number of ciliated protozoa and fungi was shown to rapidly decrease by more than 90% during the induction of MFD (Rico et al., 2015).

5.3. Laminitis

Laminitis is a clinical sign associated with SARA (Kleen et al., 2003; Krause and Oetzel, 2006). Laminitis is characterized by attachment loss between the distal phalanx and the inner hoof wall that causes the bone to be pressed down into the hoof, damaging the surrounding blood vessels and subsequently breaking the sole corium and the coronet (Li et al., 2017). Laminitis is considered the major cause of significant production loss due to compulsory culling and deaths (Oetzel, 2007; Thoenfer et al., 2005). Recently, it was found that the disturbance of ruminal microbiota plays a crucial role in laminitis onset (Li et al., 2017). Increasing levels of LPS, histamine, and lactic acid in the bloodstream during SARA play vital roles in causing laminitis. Feeding diets rich in HC has been identified as playing an important role in the pathogenesis of laminitis. Oligofructose administration was shown to induce a significant change in the ruminal microbiota and cause acute laminitis. Microbial data investigations have shown that oligofructose infusion increases the proportion of the phylum Firmicutes and the genera *Lactobacillus* and *Streptococcus*, and reduces the abundance of the phyla Bacteroidetes, Fibrobacteres, and the genera *Butyrivibrio* and *Ruminococcus* (Li et al., 2017).

5.4. Liver abscesses

Another indication of SARA at time of slaughter is liver abscess, which may reach a prevalence of 32% of the community, with more than half of the cases categorized as severe (Rezac et al., 2014). Liver abscess is a common disease of grain-fed cattle at all ages and within all types. The prevalence is most common in feedlots and is influenced by several dietary and management factors (Nagaraja and Chengappa, 1998). Liver abscess is often secondary to rumen acidosis, rumenitis, and rumen parakeratosis in cattle fed a grain diet. Liver abscesses are diet-induced, and the level of fibres negatively influence their predominance and severity.

Fusobacterium necrophorum, an anaerobic ruminal bacterium, is the primary cause of liver abscesses (Nagaraja and Chengappa, 1998). The amount of *F. necrophorum* is influenced by feed and some antimicrobial feed additives. Its role in ruminal fermentation is lactic acid metabolization of feed and epithelial protein degradation. It increases when cows are fed grain rather than forage. Ruminal *F. necrophorum* enters the portal circulation, is trapped by the liver, and then leads to the formation of abscesses. The term 'rumenitis-liver abscess complex' is usually used because of the close relationship between the ruminal pathology incidence and cattle liver abscesses (Tadepalli et al., 2009). Ruminal injuries occurring in acidosis are usually accepted as predisposing agents for liver abscesses (Nagaraja and Lechtenberg, 2007). The second most prevalent bacteria isolated from liver abscesses are *Arcanobacterium pyogenes*, which exist as a commensal microbe on the mucus membranes of the animal ruminal wall (Nagaraja and Lechtenberg, 2007).

6. Recent advanced studies in ruminal acidosis prevention

6.1. Feeding management strategies

The basic principles of prevention are a gradual adaptation to HC feeds, an adequate fibre source in the diet, and a reduction in concentrate amounts consumed by animals in a diet (Enemark, 2008; Krause and Oetzel, 2006). A well-balanced dietary fibre and starch content, and ratio of physically effective neutral detergent fibre to ruminal degradable starch, should be obtainable in diet formulations for high-producing ruminants (Xu et al., 2018). Approximately 15% of rumen-degradable starch from grains in total mixed rations may be considered a general optimum to assure normal rumen conditions and digestion, depending on the fibre content of the diet. The content of physically effective neutral detergent fibre >1.18 mm (expressed inclusive of particles-dry matter >1.18 mm) in the diet at approximately 30% to 32% is sufficient for maintaining a daily mean ruminal pH of 6.2, lowering the risk of SARA. In addition, a ratio of physically effective neutral detergent fibre >1.18 mm to ruminal degradable starch of grain that is lower than approximately 1.45 mm should be avoided (Zebeli et al., 2010). Feeding management requires a combination of proper diet formulation and good feed bunk management, including sufficient long-fibre particles in the feed (between 7% and 15% of the total mixed ration), which decreases SARA prevalence by enhancing salivation and increasing rumination (Oetzel, 2003; Ishaq et al., 2017).

6.2. Supplementation with buffer salts and organic acids

Supplementation with buffer salts has a direct effect on managing ruminal fluid pH. Supplementation of adequate buffer in the diet, such as sodium bicarbonate (from 110 to 225 g/day), neutralizes the ruminal pH changes and may prevent the overgrowth of acid-tolerant lactobacilli (Enemark, 2008). Additionally, adding

alkalinizing substances such as potassium carbonate or magnesium oxide from 270 to 410 and 50 to 90 g/day, respectively, increases ruminal pH (Enemark, 2008; Hernández et al., 2014). However, sodium bicarbonate supplementation may cause concern by upsetting the balance of ions in the body of ruminants. In addition, organic acids, such as malic acid, tannic acid and citric acid, which stimulate lactate utilization, are used directly as feed additives or through pre-treated (steeped) cereal grain (Krause and Oetzel, 2006; Zhao et al., 2021; Shen et al., 2019).

6.3. Supplementation with plant-derived extracts

Plant-derived extracts, e.g., alkaloids, terpenoids, and essential oils, have received extensive consideration from researchers due to their efficiency in maintaining ruminal pH and improving ruminal fermentation. Beta-sitosterol, a plant compound known as phytosterol, has the potential to relieve the inflammatory response and alter rumen fermentation in HC-intensive ruminant production (Xia et al., 2020). Supplementation with plant-derived alkaloids increases SCFA concentrations, which suggests advantages in improving gluconeogenesis and fermentation in ruminants (Mickdam et al., 2016).

6.4. Ionophores and probiotics

Supplementation with ionophores aims to inhibit lactate-producing bacteria, mainly *S. bovis* and *Lactobacillus* spp. Ionophores are a class of antibiotics that are not used in human medicine but may have more potential for animals. Administering ionophore rumen modifiers, such as monensin, may decrease the risk of rumen acidosis by controlling lactate production and reducing meal size. More than a decade ago, monensin has been approved for use in lactating dairy cattle in the US (Oetzel, 2007). Nevertheless, the European Union and China have banned antibiotic use in feed animals, and it has been limited in other countries due to antibiotic resistance and government concerns regarding food safety (Blanch et al., 2009; Xia et al., 2020). To enhance ruminal lactate utilization that may decrease the risk of SARA, diets can be supplemented with probiotics or direct microbial strains, such as *M. elsdenii*, or yeasts such as *Saccharomyces cerevisiae*, which stimulate the growth of ruminal protozoa, improve cellulolytic bacterial activity and suppress the proliferation of lactic acid-producing bacteria. Fungi such as *Aspergillus oryzae* have been proposed to improve rumen performance, normalize rumen fermentation, increase ruminal bacterial activity, and inhibit lactic acid production (Hernández et al., 2014).

6.5. Thiamine supplementation

Thiamine supplementation is a novel strategy to relieve SARA in animals fed HC diets, which can change rumen microbial communities and mitigate HC-induced local inflammation and ruminal epithelial disruption. It reduced the relative protein expression of interleukin-1 β , nuclear factor-kappa B unit p65, and phosphorylated nuclear factor-kappa B unit p65 in the ruminal epithelium (Ma et al., 2021; Wang et al., 2015; Zhang et al., 2020).

6.6. Immunization

Antibodies and vaccines against ruminal lactate-producing bacteria have successfully engendered a higher rumen pH and reduced the lactate concentration. In addition, the development of polyclonal antibodies against target bacteria may be effective in controlling rumen acidosis during a rapid transition to the HC diet (Blanch et al., 2009; Hernández et al., 2014).

7. Conclusion

SARA is a well-known metabolic disorder in early- and mid-lactation of high-producing dairy herds that harms animal production and health. Feeding animals HC diets leads to a reduction in the ruminal pH ≤ 5.5 and thus affects the ruminal microbial diversity and population. High-throughput sequencing, multivariate statistical techniques and bioinformatics advances, which are more accurate and sensitive, were used to estimate and monitor microbiota alterations in the rumen. Such approaches enable us to distinguish microbiota that may be numerically less abundant but functionally have an important role in the causes of the disorder. Additionally, we elucidated bacterial interactions with the host animal, which allowed us to use the power of the microbial diversity to improve animal productivity, health, and feed protection. The evidence shows that the ruminal microbiota plays a substantial role in SARA incidence, with cellulolytic bacteria decreasing and acid-tolerant bacteria such as *Streptococcus* and *Lactobacillus* spp. increasing. Free ruminal LPS resulting from Gram-negative bacterial cell lysis and histamine play a crucial role in the SARA prevalence, and consequences such as rumenitis, MFD, laminitis, and liver abscesses affect animal production and health. Understanding the dietary and microbial causes of SARA and coordinating efforts between feed and microbial management will help us develop prevention strategies. This understanding may provide further guidelines for strategies to maintain the fine balance between high growth performance and good host health. The research to date has focused on the bacterial microbiome role in SARA. However, more research is needed to fill the gaps in our knowledge on the role of the protozoal, fungal and archaeal microbiomes as potential cofactors in metabolic acidosis. In addition, we require further studies on the mechanisms by which they may contribute to disease.

Author contributions

Hongrong Wang reviewed, added intellectual content, and approved the final version of the manuscript to be published. **Mawda E. Elmhadi**: writing, reviewing, and editing. **Darien K. Ali** and **Mawahib K. Khogali** critically revised the manuscript.

Declaration of competing interest

We declare that we have no financial and personal relationships with other people or organizations that can inappropriately influence our work, and there is no professional or other personal interest of any nature or kind in any product, service and/or company that could be construed as influencing the content of this paper.

Acknowledgments

This research was partially supported by the National Natural Science Foundation of China (NSFC No. 31872988; No. 31572429) and the Priority Academic Program Development of Jiangsu Higher Education Institutions (PADA).

References

Ametaj BN. Veterinary science. In: Hudson R, Nielsen O, Bellamy J, Stephen C, editors. Metabolic disorders of dairy cattle. Paris: UNESCO- EOLSS Publishers; 2010. p. 56–73.

Bilal MS, Abaker JA, ul Aabdin Z, Xu T, Dai H, Zhang K, et al. Lipopolysaccharide derived from the digestive tract triggers an inflammatory response in the uterus of mid-lactating dairy cows during SARA. *BMC Vet Res* 2016;12(1):284.

Blanch M, Calsamiglia S, DiLorenzo N, DiCostanzo A, Muetzel S, Wallace RJ. Physiological changes in rumen fermentation during acidosis induction and its control using a multivalent polyclonal antibody preparation in heifers. *J Anim Sci* 2009;87(5):1722–30.

Brestoff JR, Artis D. Commensal bacteria at the interface of host metabolism and the immune system. *Nat Immunol* 2013;14(7):676–84.

Calsamiglia S, Blanch M, Ferret A, Moya D. Is subacute ruminal acidosis a pH related problem? Causes and tools for its control. *Anim Feed Sci Technol* 2012;172(1–2): 42–50.

Chaucheyras-Durand F, Ossa F. REVIEW: the rumen microbiome: composition, abundance, diversity, and new investigative tools. *Prof Anim Sci* 2014;30(1): 1–12.

Eckel EF, Ametaj BN. Invited review: role of bacterial endotoxins in the etiopathogenesis of periparturient diseases of transition dairy cows. *J Dairy Sci* 2016;99(8):5967–90.

Emmanuel DGV, Madsen KL, Churchill TA, Dunn SM, Ametaj BN. Acidosis and lipopolysaccharide from *Escherichia coli* B: 055 cause hyperpermeability of rumen and colon tissues. *J Dairy Sci* 2007;90(12):5552–7.

Enemark JM. The monitoring, prevention and treatment of sub-acute ruminal acidosis (SARA): a review. *Vet J* 2008;176(1):32–43.

Enemark JMD, Jorgensen R, Enemark PS. Rumen acidosis with special emphasis on diagnostic aspects of subclinical rumen acidosis: a review. *Vet Zootech* 2002;20(42):16–29.

FAO. Milk fact. Food and Agriculture Organization of United Nation; 2021.

Ghaffari MH, Khafipour E, Steele MA. Periparturient diseases of dairy cows. In: Ametaj BN, editor. Systems biology and ruminal acidosis. Switzerland: Springer; 2017. p. 51–69.

Goto Y, Kiyono H. Epithelial barrier: an interface for the cross-communication between gut flora and immune system. *Immunol Rev* 2012;245(1):147–63.

Gozho G, Krause D, Plaizier J. Ruminal lipopolysaccharide concentration and inflammatory response during grain-induced subacute ruminal acidosis in dairy cows. *J Dairy Sci* 2007;90(2):856–66.

Herdtr TH. Ruminant adaptation to negative energy balance: Influences on the etiology of ketosis and fatty liver. *Vet Clin N Am Food Anim Pract* 2000;16(2): 215–30.

Hernández J, Benedito JL, Abuelo A, Castillo C. Ruminal acidosis in feedlot: from aetiology to prevention. *Sci World J* 2014:702572.

Hook SE, Steele MA, Northwood KS, Dijkstra J, France J, Wright A-DG, et al. Impact of subacute ruminal acidosis (SARA) adaptation and recovery on the density and diversity of bacteria in the rumen of dairy cows. *FEMS Microbiol Ecol* 2011a;78(2):275–84.

Hook SE, Steele MA, Northwood KS, Wright A-DG, McBride BW. Impact of high-concentrate feeding and low ruminal pH on methanogens and protozoa in the rumen of dairy cows. *Microb Ecol* 2011b;62(1):94–105.

Howard JL. Ruminal metabolic acidosis. *Bov Pract* 1981:44–53.

Hua C, Tian J, Tian P, Cong R, Luo Y, Geng Y, et al. Feeding a high concentration diet induces unhealthy alterations in the composition and metabolism of ruminal microbiota and host response in a goat model. *Front Microbiol* 2017:8138.

Huo W, Zhu W, Mao S. Impact of subacute ruminal acidosis on the diversity of liquid and solid-associated bacteria in the rumen of goats. *World J Microbiol Biotechnol* 2014;30(2):669–80.

Ishaq SL, AlZahal O, Walker N, McBride B. An investigation into rumen fungal and protozoal diversity in three rumen fractions, during high-fiber or grain-induced sub-acute ruminal acidosis conditions, with or without active dry yeast supplementation. *Front Microbiol* 2017:81943.

Jaramillo-López E, Itza-Ortiz MF, Peraza-Mercado G, Carrera-Chávez JM. Ruminal acidosis: strategies for its control. *Austral J Vet Sci* 2017;49(3):139–48.

Khafipour E, Li S, Plaizier JC, Krause DO. Rumen microbiome composition determined using two nutritional models of subacute ruminal acidosis. *Appl Environ Microbiol* 2009;75(22):7115–24.

Khafipour E, Li S, Tun HM, Derakhshani H, Moossavi S, Plaizier JC. Effects of grain feeding on microbiota in the digestive tract of cattle. *Anim Front* 2016;6(2):13–9.

Kleen J, Hooijer G, Rehage J, Noordhuizen J. Subacute ruminal acidosis (SARA): a review. *J Vet Med* 2003;50(8):406–14.

Kong Y, Teather R, Forster R. Composition, spatial distribution, and diversity of the bacterial communities in the rumen of cows fed different forages. *FEMS Microbiol Ecol* 2010;74:612–22.

Krause KM, Oetzel GR. Understanding and preventing subacute ruminal acidosis in dairy herds: a review. *Anim Feed Sci Technol* 2006;126(3–4):215–36.

Kurashima Y, Goto Y, Kiyono H. Mucosal innate immune cells regulate both gut homeostasis and intestinal inflammation. *Eur J Immunol* 2013;43(12):3108–15.

Lean S. Ruminal acidosis—aetiopathogenesis, prevention and treatment. *Aust. Vet. Assoc* 2007:617–43.

Li H, Liu J, Zhu W, Mao S. Intraruminal infusion of oligofructose alters ruminal microbiota and induces acute laminitis in sheep. *J Anim Sci* 2017;95(12): 5407–19.

Ma Y, Wang C, Zhang H, Yu L, Dong L, Gong D, et al. Illumina sequencing and metabolomics analysis reveal thiamine modulation of ruminal microbiota and metabolome characteristics in goats fed a high-concentrate diet. *Front Microbiol* 2021;12:653283.

Mani V, Weber TE, Baumgard LH, Gabler NK. Growth and development symposium: endotoxin, inflammation, and intestinal function in livestock. *J Anim Sci* 2012;90(5):1452–65.

Mao S, Zhang R, Wang D, Zhu W. Impact of subacute ruminal acidosis (SARA) adaptation on rumen microbiota in dairy cattle using pyrosequencing. *Anaerobe* 2013:2412–9.

Mao SY, Huo WJ, Zhu WY. Microbiome–metabolome analysis reveals unhealthy alterations in the composition and metabolism of ruminal microbiota with increasing dietary grain in a goat model. *Environ Microbiol* 2016;18(2):525–41.

- McCann JC, Luan S, Cardoso FC, Derakhshani H, Khafipour E, Looor JJ. Induction of subacute ruminal acidosis affects the ruminal microbiome and epithelium. *Front Microbiol* 2016;7701.
- Mickdam E, Khaioosa-ard R, Metzler-Zebeli BU, Klevenhusen F, Chizzola R, Zebeli Q. Rumen microbial abundance and fermentation profile during severe subacute ruminal acidosis and its modulation by plant derived alkaloids in vitro. *Anaerobe* 2016;394–413.
- Nagaraja TG, Chengappa MM. Liver abscesses in feedlot cattle: a review. *J Anim Sci* 1998;76(1):287–98.
- Nagaraja TG, Lechtenberg KF. Liver abscesses in feedlot cattle. *Vet Clin North Am Food Anim Pract* 2007;23(2):351–69.
- Nagaraja TG, Titgemeyer EC. Ruminal acidosis in beef cattle: the current microbiological and nutritional outlook. *J Dairy Sci* 2007;90:E17–38.
- O'Grady L, Doherty ML, Mulligan FJ. Subacute ruminal acidosis (SARA) in grazing Irish dairy cows. *Vet J* 2008;176(1):44–9.
- Oetzel GR. Subacute ruminal acidosis in dairy cattle. *Adv Dairy Technol* 2003;15:307–17.
- Oetzel GR. Subacute ruminal acidosis in dairy herds: physiology, pathophysiology, milk fat responses, and nutritional management. 40th Annual Conference, American Association of Bovine Practitioners; 2007. p. 89–119.
- Petri R, Schwaiger T, Penner G, Beauchemin K, Forster R, McKinnon J, et al. Changes in the rumen epimural bacterial diversity of beef cattle as affected by diet and induced ruminal acidosis. *Appl Environ Microbiol* 2013;79(12):3744–55.
- Pitta DW, Indugu N, Vecchiarelli B, Rico DE, Harvatine KJ. Alterations in ruminal bacterial populations at induction and recovery from diet-induced milk fat depression in dairy cows. *J Dairy Sci* 2018;101(1):295–309.
- Plaizier JC, Krause DO, Gozho GN, McBride BW. Subacute ruminal acidosis in dairy cows: the physiological causes, incidence and consequences. *Vet J* 2009;176(1):21–31.
- Plaizier JC, Li S, Tun HM, Khafipour E. Nutritional models of experimentally-induced subacute ruminal acidosis (SARA) differ in their impact on rumen and hindgut bacterial communities in dairy cows. *Front Microbiol* 2017;7:72128.
- Rezac D, Thomson D, Siemens M, Prouty F, Reinhardt C, Bartle S. A survey of gross pathologic conditions in cull cows at slaughter in the Great Lakes region of the United States. *J Dairy Sci* 2014;97(7):4227–35.
- Ribeiro Alves Lourenço M, Ramos-Morales E, Wallace RJ. The role of microbes in rumen lipolysis and biohydrogenation and their manipulation. *Animal* 2010;4(7):1008–23.
- Rico DE, Preston SH, Risser JM, Harvatine KJ. Rapid changes in key ruminal microbial populations during the induction of and recovery from diet-induced milk fat depression in dairy cows. *Br J Nutr* 2015;114(3):358–67.
- Shen YZ, Ding LY, Chen LM, Xu JH, Zhao R, Yang WZ, et al. Feeding corn grain steeped in citric acid modulates rumen fermentation and inflammatory responses in dairy goats. *Animal* 2019;13(2):301–8.
- Schweigel M, Freyer M, Leclercq S, Etschmann B, Lodemann U, Böttcher A, et al. Luminal hyperosmolarity decreases Na transport and impairs barrier function of sheep rumen epithelium. *J Comp Physiol B* 2005;175(8):575–91.
- Tadepalli S, Narayanan SK, Stewart GC, Chengappa MM, Nagaraja TG. *Fusobacterium necrophorum*: a ruminal bacterium that invades liver to cause abscesses in cattle. *Anaerobe* 2009;15(1–2):36–43.
- Tao S, Duanmu Y, Dong H, Ni Y, Chen J, Shen X, et al. High concentrate diet induced mucosal injuries by enhancing epithelial apoptosis and inflammatory response in the hindgut of goats. *PLoS One* 2014a;9(10):e111596.
- Tao S, Duanmu Y, Dong H, Tian J, Ni Y, Zhao R. A high-concentrate diet induced colonic epithelial barrier disruption is associated with the activating of cell apoptosis in lactating goats. *BMC Vet Res* 2014b;10(1):235.
- Thoenfer MB, Wattle O, Pollitt CC, French KR, Nielsen SS. Histopathology of oligofructose-induced acute laminitis in heifers. *J Dairy Sci* 2005;88(8):2774–82.
- Wang H, Pan X, Wang C, Wang M, Yu L. Effects of different dietary concentrate to forage ratio and thiamine supplementation on the rumen fermentation and ruminal bacterial community in dairy cows. *Anim Prod Sci* 2015;55(2):189–93.
- Weimer PJ, Stevenson DM, Mertens DR. Shifts in bacterial community composition in the rumen of lactating dairy cows under milk fat-depressing conditions. *J Dairy Sci* 2010;93(1):265–78.
- Xia G, Sun J, Fan Y, Zhao F, Ahmed G, Jin Y, et al. β -sitosterol attenuates high grain diet-induced inflammatory stress and modifies rumen fermentation and microbiota in sheep. *Animals* 2020;10(1):171.
- Xu J, Wang H, Shen Y, Zhao R, Ge R, Yu L, et al. Impact of dietary carbohydrate balance on rumen fermentation, eating behaviour, growth and development of 8–10-month-old heifers. *Anim Prod Sci* 2018;58(11):2042–8.
- Zebeli Q, Mansmann D, Steingass H, Ametaj BN. Balancing diets for physically effective fibre and ruminally degradable starch: a key to lower the risk of subacute rumen acidosis and improve productivity of dairy cattle. *Livest Sci* 2010;127(1):1–10.
- Zebeli Q, Metzler-Zebeli BU. Interplay between rumen digestive disorders and diet-induced inflammation in dairy cattle. *Res Vet Sci* 2012;93(3):1099–108.
- Zeng H, Guo C, Sun D, Seddik H-e, Mao S. The ruminal microbiome and metabolome alterations associated with diet-induced milk fat depression in dairy cows. *Metabolites* 2019;9(7):154.
- Zhang H, Peng A, Zhao F, Yu L, Wang M, Osorio J, et al. Thiamine ameliorates inflammation of the ruminal epithelium of Saanen goats suffering from subacute ruminal acidosis. *J Dairy Sci* 2020;103(2):1931–43.
- Zhang J, Shi H, Wang Y, Li S, Cao Z, Ji S, et al. Effect of dietary forage to concentrate ratios on dynamic profile changes and interactions of ruminal microbiota and metabolites in Holstein heifers. *Front Microbiol* 2017;8:2206.
- Zhao C, Liu G, Li X, Guan Y, Wang Y, Yuan X, et al. Inflammatory mechanism of rumenitis in dairy cows with subacute ruminal acidosis. *BMC Vet Res* 2018;14(1):135.
- Zhao FF, Zhang XZ, Zhang Y, Elmhadi M, Qin YY, Sun H, et al. Tannic acid-steeped corn grain modulates in vitro ruminal fermentation pattern and microbial metabolic pathways. *Front Vet Sci* 2021;8:698108.