

Expression of the *Tas1r3* and *Pept1* genes in the digestive tract of wagyu cattle

Weihong Fan,^{*} Shoichiro Saito,[†] and Shuichi Matsumura^{†,1}

^{*}Graduate School of Natural Science and Technology, Gifu University, 1-1 Yanagido, Gifu 501-1193, Japan; and

[†]Faculty of Applied Biological Sciences, Gifu University, 1-1 Yanagido, Gifu 501-1193, Japan

ABSTRACT: Animals have precise recognition systems for amino acids and peptides that regulate their feeding behavior as well as metabolic responses. Because of their particular gastrointestinal structure, ruminants are expected to have unique mechanisms of amino acid regulation in the digestive tract. To better understand these mechanisms in the ruminant digestive tract, the expression of *Tas1r3* and *Pept1* was studied along the gastrointestinal tract of Japanese Black cattle through quantitative RT-PCR and immunohistochemistry. *Tas1r3* mRNA was detected ubiquitously along the gastrointestinal tract, and the most predominant expression was observed in the reticulum. In addition, the presence of

Tas1r3 receptor was confirmed in the rumen through immunohistochemistry. The expression level of *Pept1* mRNA was higher in the forestomach (rumen, reticulum, and omasum) and small intestine (duodenum) than that in the tongue, and predominant expression was observed in the rumen. By contrast, a negligible amount of *Pept1* mRNA was detected in the abomasum and large intestine. Further studies on the roles of *Tas1r3* and *Pept1* in the digestive tract, in particular, in the four components of the stomach, will help us to understand the mechanisms of amino acids regulation in ruminants and provide the basis for formulating cattle diets to improve the health and productivity of cattle.

Key words: amino acids, digestive tract, peptides, *Pept1*, *Tas1r3*, wagyu cattle

© The Author(s) 2020. Published by Oxford University Press on behalf of the American Society of Animal Science.

This is an Open Access article distributed under the terms of the Creative Commons Attribution Non-Commercial License (<http://creativecommons.org/licenses/by-nc/4.0/>), which permits non-commercial re-use, distribution, and reproduction in any medium, provided the original work is properly cited. For commercial re-use, please contact journals.permissions@oup.com

Transl. Anim. Sci. 2020.4:980-985
doi: 10.1093/tas/txaa019

INTRODUCTION

Animals need to ensure adequate absorption of amino acids from the environment. They must respond appropriately to the available sources of nutrients in their environment. Therefore, they have developed precise amino acid recognition systems that regulate their feeding behavior as well as metabolic responses.

Taste receptors of the *Tas1rs* family play a central role in their amino acid recognition (Zhao et al., 2003). The *Tas1rs* family belong to the G

protein-coupled receptor (GPCR) superfamily C subtype and consist of three principal domains: an amino-terminal domain (ATD), a cysteine-rich domain (CRD), and a transmembrane domain (TMD) (Kunishima et al., 2000). *Tas1rs* receptors function as heterodimers; *Tas1r2* and *Tas1r3* bind to form sweet receptors, whereas *Tas1r1* and *Tas1r3* constitute a receptor for umami substances including L-amino acids (Nelson et al., 2002; Zhao et al., 2003). Recently, it has been demonstrated that *Tas1rs* receptors exist not only in oral tissues but also in various nonoral tissues (Behrens and Meyerhof, 2011).

Pept1, also known as solute carrier family 15 member 1 (SLC15A1), utilizes a proton gradient

¹Corresponding author: matsumur@gifu-u.ac.jp

Received September 26, 2019.

Accepted February 7, 2020.

(Adibi, 1997) to transport di/tri-peptides from the intestinal lumen into epithelial cells (Zhang et al., 2016). In the intestine, the *Pept1* pathway constitutes a major mechanism for absorption of the products of protein digestion (Fei et al., 1994). In recent years, many researchers have reported that *Pept1* is expressed in different organs and has multiple biological functions (Zhang et al., 2016; Spanier and Rohm, 2018; Viennois et al., 2018).

Although the expression of taste receptors and transporters for amino acid regulation in the digestive tract has been studied intensively in many mammalian species, little information is available for ruminants. Ruminants are characterized by their four compartments of the stomach, that is, the rumen, reticulum, omasum, and abomasum. Because of the particular gastrointestinal structure, some of the amino acids consumed by ruminants are degraded by rumen microorganisms (Diao et al., 2019). However, the mechanisms of amino acid regulation in the digestive tract of ruminants are still unclear. Because the gastrointestinal structure of ruminants is unique compared with nonruminants, the roles of taste receptors and peptide transporters must also be unique. The pattern of the expression of taste receptor and transporter genes, that is, whether they are expressed in a particular organ or not is quite different between species (e.g., Gonda et al., 2013). To date, however, there have been few reports about the expression of taste receptor and peptide transporter genes in the digestive tract of ruminants. Therefore, one receptor gene (*Tas1r3*) and one transporter gene (*Pept1*) were chosen as targets of the present study. To better understand their roles in the ruminant digestive tract, the expression of *Tas1r3* and *Pept1* was studied along the gastrointestinal tract of Japanese Black cattle through quantitative RT-PCR and immunohistochemistry.

MATERIALS AND METHODS

Wagyu cattle include four types of Japanese cattle: the Black, Brown, Shorthorn, and Polled breeds (Gotoh et al., 2018). Of this, Japanese Black cattle are the most predominant breed of them because of their high-performance marbled beef and extremely high economic value (Gotoh et al., 2018). Samples of the gastrointestinal tract from Japanese Black cattle were acquired from the meat inspection center of Gifu City, Japan, following humane slaughter procedures. Tissue samples for the RT-qPCR analysis were obtained from three individuals (one female and two castrated males) reared in the same farm. Small blocks of tissues were incubated overnight at 4 °C in

RNA later (Invitrogen), and stored at -20 °C until use. Total RNA was extracted from the tissues using the RNeasy Plus mini kit (QIAGEN, Germany). Elimination of genomic DNA from the RNA samples and reverse transcription of RNA into cDNA were carried out using PrimeScript RT reagent kit with gDNA Eraser (TaKaRa, Japan). The following primers for the target genes (*Tas1r3* and *Pept1*) and an internal marker (*Gapdh*) were designed using Primer 3 plus software (<http://www.bioinformatics.nl/cgi-bin/primer3plus/primer3plus.cgi>) (Thornton and Basu, 2011): 5'-TGTGCACCAGGTTGTCGG-3' and 5'-AAAGGTCATAGCCCAGACGC-3' (*Tas1r3*), 5'-ATGGCCTTAACCAGAAGCCAGA-3' and 5'-TCACTGGCATTGTGACTGGAGAC-3' (*Pept1*), and 5'-AAGGTCGGAGTGAACGGATTTC-3' and R: 5'-ATGGCGACGATGTCCACTTT-3' (*Gapdh*). The PCR products were checked using the standard 2% agarose gel electrophoresis. Real-time PCR was carried out with StepOne Plus (Applied Biosystems) using THUNDERBIRD SYBR qPCR Mix (Toyobo, Japan). Standard real-time PCR condition (Pre-denaturation: 95 °C, 3 min; denaturation: 95 °C, 15 s; annealing: 60 °C, 30 s; extension: 60 °C, 1 min; 40 cycles) were applied to all analyses, which were repeated three times. The relative expression of *Tas1r3* and *Pept1* were determined by comparative calculation with the internal marker *Gapdh* (Livak and Schmittgen, 2001). The ΔC_t was determined by subtracting the *Gapdh* C_t value from the C_t value of the target gene from each organ, and the $\Delta\Delta C_t$ was determined by subtracting the ΔC_t from the baseline organ (tongue in the present study) from the ΔC_t at each organ. The relative quantity of target gene mRNA was expressed as $2^{-\Delta\Delta C_t}$.

Tissues for immunohistochemistry were embedded overnight in 4% paraformaldehyde. An antigen retrieval treatment with normal horse serum was applied to unmask immunogenic epitopes for 30 min. Sections were incubated overnight at 4 °C with a primary antibody (G-2, SC-398996, CosMo Bio Co, Japan) diluted in solution (1:200 concentration). Sections were incubated at room temperature with a secondary antibody (biotinylated anti-mouse IgG horse serum; Funakoshi, Japan) at a dilution of 1:500 for 30 min. Antibody complexes were visualized with avidin-biotin-peroxidase conjugate (ABC Elite; Funakoshi, Japan) and 3,3'-diaminobenzidine (DAB; Dojindo, Japan).

RESULTS AND DISCUSSION

When the PCR products were checked using 2% agarose gel electrophoresis, a clear single target band was observed for each gene, indicating the

high specificity of the PCR primers. Therefore, the relative expression of the *Tas1r3* and *Pept1* genes in the digestive tract was quantified using real-time PCR (Fig. 1).

Expression of *Tas1r3* in the Gastrointestinal Tract

The mRNA of *Tas1r3* was detected ubiquitously in both the digestive tract and tongue (Fig. 1a). The expression level in most parts of the digestive tract was higher than that in the tongue. The most predominant expression was observed in the reticulum. Following real-time PCR, the existence of the *Tas1r3* protein was validated by immunohistochemistry. The *Tas1r3* protein was clearly expressed in the rumen (Fig. 2).

The *Tas1r* family members (*Tas1r1*, *Tas1r2*, and *Tas1r3*) mediate sweet and umami tastes in mammals: a heterodimer of *Tas1r2* and *Tas1r3* functions as a sweet taste receptor while a heterodimer of *Tas1r1* and *Tas1r3* functions as an umami taste receptor (Li, 2009). Disruption of the *Tas1r3* gene in knockout mice diminishes the taste responses to both sweet and umami taste stimuli (Damak et al.,

2003; Zhao et al., 2003). It has been well demonstrated that the *Tas1r3* receptor is expressed in small numbers of cells in the anterior and posterior taste buds (Chaudhari et al., 2009). Recently, it has been reported that *Tas1r3* is also expressed in extra-oral tissues, including the gastrointestinal tract, brain, bladder, pancreas, male reproductive organs, immune system, adipose tissue, and bone, in many animal species including humans and mice (Laffitte et al., 2014). However, for ruminants, few reports are available except for Moran et al. (2014), which reported the existence of *Tas1r3* protein in the small intestine of sheep and cattle. The present study demonstrated that *Tas1r3* mRNA was expressed ubiquitously in the cattle digestive tract, and confirmed the existence of *Tas1r3* protein in the cattle rumen.

Although the *Tas1r3* gene was ubiquitously expressed in the cattle digestive tract, the expression level was relatively high in the reticulum, omasum, and cecum. The mRNA of *Tas1r1* and *Tas1r3* are known to be expressed in the stomach of mice and humans as well as in the small intestine and colon (Bezençon et al., 2007; Janssen and Depoortere, 2013). The existence of *Tas1r3* protein was confirmed in the mouse stomach by immunohistochemistry (Hass et al., 2010). However, studies on the expression of the *Tas1rs* family in the stomach

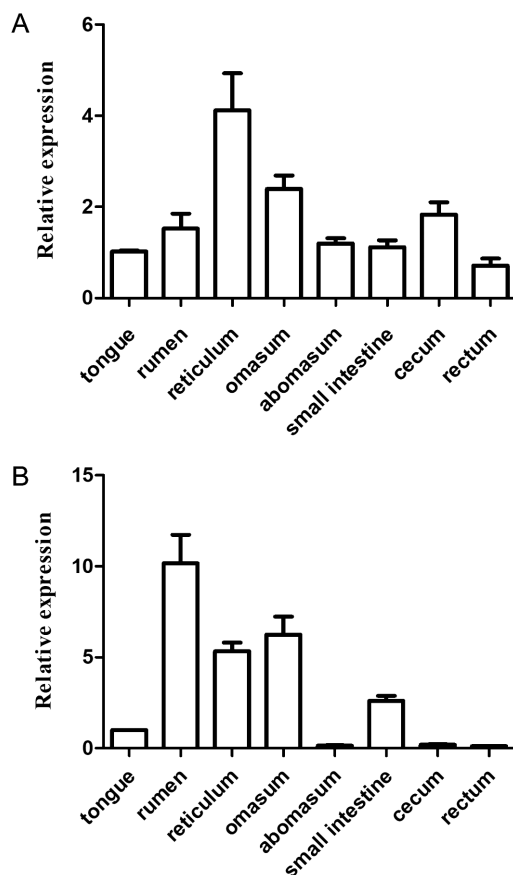


Figure 1. Relative expression of the (a) *Tas1r3* and (b) *Pept1* genes in the digestive tract of waxy cattle. The error bars indicate the standard error of the mean (SEM) from three individuals. The expression level of tongue was set at 1.

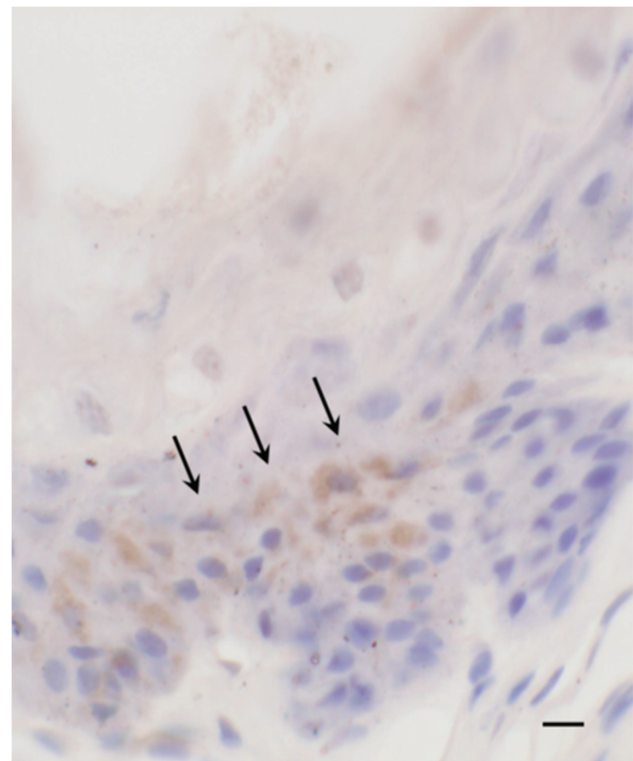


Figure 2. Immunohistochemistry of the *Tas1r3* receptors in the rumen of Japanese Black cattle. The scale bar represents 10 μ m.

are few compared with their expression in the intestine, and little is known about their function. Intestinal *Tas1r2* has been suggested to be a possible sensor of glucose or other nutritional ingredients (Cumming and Overduin, 2007; Depoortere, 2014; Calvo and Egan, 2015). The functions of intestinal *Tas1rs* might be related to hormonal regulation and/or defense mechanisms, as suggested for other types of taste receptors (Sternini, 2007; Breer et al., 2012; Prandi et al., 2013). Further studies, in particular, on the type of cells in the cattle stomach are important to clarify the role of *Tas1rs* in the stomach of ruminants.

Expression of Pept1 in the Gastrointestinal Tract

Although *Pept1* mRNA was also detected in the gut and tongue, its expression pattern was different from that of *Tas1r3* (Fig. 1b). The expression level in the forestomach (rumen, reticulum, and omasum) was more than five times higher than that in the tongue. Predominant expression was observed in the rumen. Considerable expression of *Pept1* mRNA was also observed in the small intestine (duodenum). By contrast, a negligible amount of *Pept1* mRNA was detected in the abomasum and as large intestine.

To the best of our knowledge, no other study except Wang et al. (2016) compared the expression of the *Pept1* gene quantitatively in the gastrointestinal tract of ruminants. As a peptide transporter, it is well known that *Pept1* plays a vital role in the absorption of peptides in the small intestine (Spanier and Rohm, 2018). Correspondingly, it has been reported that *Pept1* mRNA was highly expressed in the small intestine compared with other organs (Spanier and Rohm, 2018). The very low level of expression in the large intestine of cattle demonstrated by the present study agrees with previous studies in humans and mice (e.g., Groneberg et al., 2001; Terada et al., 2005). By contrast, the predominant expression of *Pept1* mRNA in the forestomach of cattle in the present study is a little puzzling because very weak expression in the stomach has been reported in humans and mice (Terada et al., 2005; Lu and Klaassen, 2006). Wang et al. (2016) also reported considerable amounts of mRNA expression in the forestomach of yak (*Bos grunniens*) and indigenous cattle in the Qinghai-Tibetan plateau.

In ruminants, protein from the diet is generally degraded quickly and converted into microbial protein in the rumen, and then microbial protein flows into the small intestine (Gruninger et al., 2019).

Therefore, it was believed that peptide absorption occurred mostly in the small intestine. However, several authors have studied amino acid net flux in the portal vein in sheep and cattle and suggested that peptide absorption occurs in the stomach (DiRienzo, 1992; Rémond et al., 2000; Tagari et al., 2004, 2008). Based on these studies, Gilbert et al. (2008) pointed out a possible important role of the rumen and omasum in the absorption of amino acids in the form of small peptides. The present study is consistent with these results suggesting significant uptake of peptides from the forestomach in ruminants. For ruminants, peptide absorption appears to be more important than was previously believed. The idea that absorption from the small intestine is the only way to ingest amino acids into the body should be reconsidered.

Implications for the Future

The results described above suggested that the function of the digestive tract of ruminants is not as simple as previously believed. It was believed that the absorption of peptides by animals mainly depended on *Pept1*, which is located on the brush border membrane of the intestine (Fei et al., 1994; Spanier and Rohm, 2018). The high level of *Pept1* mRNA expression in the present study suggested that the forestomach of ruminants might play a role in the absorption of nutrients and/or regulation of homeostasis. *Tas1r3* in the gastrointestinal tract is believed to detect amino acids and induce a series of downstream signals such as the release of gastrointestinal hormones (Vancleef et al., 2015). The secretion of hormones might affect the ingestion of nutrients via feeding behavior as well as the absorption of nutrients in the gastrointestinal tract. However, our knowledge about the roles of *Tas1r3* and *Pept1* in the digestive tract of ruminants is quite limited. Because the gastrointestinal structure of ruminants is unique compared with nonruminants, the roles of *Tas1r3* and *Pept1* must also be unique. Comprehensive studies on *Tas1rs* and *Pept1* should be promising because a recent study in rats has suggested an interaction between them (Arakawa et al., 2016).

Research on *Tas1r3* and *Pept1* in the digestive tract will be applied to improve the efficiency of cattle production, dietary protein use, and to solve environmental issues such as reduction of the emission of greenhouse gasses in the future. Ruminants are, and continue to be, the main source of protein in areas of environmentally harsh conditions and weak infrastructures because they can digest

fibrous feeds that cannot be directly consumed by humans (Gerber et al., 2015). The global consumption of cattle meat is predicted to increase as the global population increases (Salter, 2017). However, cattle meat production is sometimes considered inefficient in terms of the conversion of natural resources into edible products (Gerber et al., 2015; Salter, 2017). Therefore, it is imminent to improve the efficiency of converting energy and protein in the cattle diet into human edible energy and protein. Further knowledge about the roles of *Tas1r3* and *Pept1* in the digestive tract will be indispensable to tackle this challenging task because they play central roles in the regulation of amino acids and peptides in the gut.

Japanese wagyu beef is famous for its marbling and high quality, specific sweet and fatty aroma, sweet rather than greasy taste, and a high quality umami flavor (Matsuishi et al., 2001). Appropriate nutritional management is indispensable to produce beef that satisfies the requirements of the Japanese beef market (Gotoh et al., 2018). However, there are many problems with the management of wagyu cattle, including dependency on imported grains for feed and high production costs accounting for more than 90% of the sale price (Gotoh et al., 2018). Further studies on *Tas1r3* and *Pept1* can contribute to the wagyu industry by providing the knowledge necessary for stable and efficient feeding systems and production of high quality meat.

CONCLUSION

The present study determined the mRNA expression of *Tas1r3* and *Pept1* quantitatively in the digestive tract of Japanese wagyu cattle. Further studies are required on the roles of *Tas1r3* and *Pept1* in the digestive tract, in particular, in the four components of the stomach. Such studies will help us to understand the mechanism of amino acids regulation in ruminants and provide a basis for formulating cattle diets to improve the health and productivity of cattle.

ACKNOWLEDGMENTS

The authors thank to Prof. Hiroo Imai and Misa Hayashi, Molecular Biology Section, Primate Research Institute, Kyoto University, Prof. Yasuro Atoji, Faculty of Applied Biological Sciences, Gifu University, and the staff of Division of Genomics Research, Life Science Research Center, Gifu University for their support during the experiments. The authors also thank to the staff of the

meat inspection center of Gifu City for their support during sample collection.

Conflict of interest statement. None declared

LITERATURE CITED

- Adibi, S. A. 1997. The oligopeptide transporter (Pept-1) in human intestine: biology and function. *Gastroenterology*. 113:332–340. doi:10.1016/s0016-5085(97)70112-4.
- Arakawa, H., T. Ohmachi, K. Ichiba, H. Kamioka, T. Tomono, M. Kanagawa, Y. Idota, Y. Hatano, K. Yano, K. Morimoto, et al. 2016. Interaction of peptide transporter 1 with D-glucose and L-glutamic acid; possible involvement of taste receptors. *J. Pharm. Sci.* 105:339–342. doi:10.1016/j.xphs.2015.11.024.
- Behrens, M., and W. Meyerhof. 2011. Gustatory and extragustatory functions of mammalian taste receptors. *Physiol. Behav.* 105:4–13. doi:10.1016/j.physbeh.2011.02.010.
- Bezençon, C., J. le Coutre, and S. Damak. 2007. Taste-signaling proteins are coexpressed in solitary intestinal epithelial cells. *Chem. Senses*. 32:41–49. doi:10.1093/chemse/bj034.
- Breer, H., J. Eberle, C. Frick, D. Haid, and P. Widmayer. 2012. Gastrointestinal chemosensation: chemosensory cells in the alimentary tract. *Histochem. Cell Biol.* 138:13–24. doi:10.1007/s00418-012-0954-z.
- Calvo, S. S., and J. M. Egan. 2015. The endocrinology of taste receptors. *Nat. Rev. Endocrinol.* 11:213–227. doi:10.1038/nrendo.2015.7.
- Chaudhari, N., E. Pereira, and S. D. Roper. 2009. Taste receptors for umami: the case for multiple receptors. *Am. J. Clin. Nutr.* 90:738S–742S. doi:10.3945/ajcn.2009.27462H.
- Cumming, D. E., and J. Overduin. 2007. Gastrointestinal regulation of food intake. *J. Clin. Invest.* 117:13–23. doi:10.1172/JCI30227.
- Damak, S., M. Rong, K. Yasumatsu, Z. Kokrashvili, V. Varadarajan, S. Zou, P. Jiang, Y. Ninomiya, and R. F. Margolskee. 2003. Detection of sweet and umami taste in the absence of taste receptor T1r3. *Science*. 301:850–853. doi:10.1126/science.1087155.
- Depoortere, I. 2014. Taste receptors of the gut: emerging roles in health and disease. *Gut*. 63:179–190. doi:10.1136/gutjnl-2013-305112.
- Diao, Q., R. Zhang, and T. Fu. 2019. Review of strategies to promote rumen development in calves. *Animals*. 9:490. doi:10.3390/ani9080490.
- DiRienzo, D. B. 1992. Free and peptide amino acid fluxes across the mesenteric and non-mesenteric viscera of sheep and calves [PhD dissertation]: Virginia Tech.
- Fei, Y. J., Y. Kanai, S. Nussberger, V. Ganapathy, F. H. Leibach, M. F. Romero, S. K. Singh, W. F. Boron, and M. A. Hediger. 1994. Expression cloning of a mammalian proton-coupled oligopeptide transporter. *Nature*. 368:563–566. doi:10.1038/368563a0.
- Gerber, P. J., A. Mottet, C. I. Opio, A. Falcucci, and F. Teillard. 2015. Environmental impacts of beef production: review of challenges and perspectives for durability. *Meat Sci.* 109:2–12. doi:10.1016/j.meatsci.2015.05.013.
- Gilbert, E. R., E. A. Wong, and K. E. Webb, Jr. 2008. Board-invited review: peptide absorption and utilization: implications for animal nutrition and health. *J. Anim. Sci.* 86:2135–2155. doi:10.2527/jas.2007-0826.
- Gonda, S., S. Matsumura, S. Saito, Y. Go, and H. Imai. 2013. Expression of taste signal transduction molecules in the

- caecum of common marmosets. *Biol. Lett.* 9:20130409. doi:10.1098/rsbl.2013.0409.
- Gotoh, T., T. Nishimura, K. Kuchida, and H. Mannen. 2018. The Japanese Wagyu beef industry: current situation and future prospects—a review. *Asian Australas. J. Anim. Sci.* 31:933–950. doi:10.5713/ajas.18.0333.
- Groneberg, D. A., F. Döring, P. R. Eynott, A. Fischer, and H. Daniel. 2001. Intestinal peptide transport: ex vivo uptake studies and localization of peptide carrier PEPT1. *Am. J. Physiol. Gastrointest. Liver Physiol.* 281:G697–G704. doi:10.1152/ajpgi.2001.281.3.G697.
- Gruninger, R. J., G. O. Ribeiro, A. Cameron, and T. A. McAllister. 2019. Invited review: application of meta-omics to understand the dynamic nature of the rumen microbiome and how it responds to diet in ruminants. *Animal.* 13:1843–1854. doi:10.1017/S1751731119000752.
- Hass, N., K. Schwarzenbacher, and H. Breer. 2010. T1R3 is expressed in brush cells and ghrelin-producing cells of murine stomach. *Cell. Tissue. Res.* 339:493–504. doi:10.1007/s00441-009-0907-6.
- Janssen, S., and I. Depoortere. 2013. Nutrient sensing in the gut: new roads to therapeutics? *Trends Endocrinol. Metab.* 24:92–100. doi:10.1016/j.tem.2012.11.006.
- Kunishima, N., Y. Shimada, Y. Tsuji, T. Sato, M. Yamamoto, T. Kumasaka, S. Nakanishi, H. Jingami, and K. Morikawa. 2000. Structural basis of glutamate recognition by a dimeric metabotropic glutamate receptor. *Nature.* 407:971–977. doi:10.1038/35039564.
- Laffitte, A., F. Neiers, and L. Briand. 2014. Functional roles of the sweet taste receptor in oral and extraoral tissues. *Curr. Opin. Clin. Nutr. Metab. Care.* 17:379–385. doi:10.1097/MCO.0000000000000058.
- Li, X. 2009. T1R receptors mediate mammalian sweet and umami taste. *Am. J. Clin. Nutr.* 90:733S–737S. doi:10.3945/ajcn.2009.27462G.
- Livak, K. J., and T. D. Schmittgen. 2001. Analysis of relative gene expression data using real-time quantitative PCR and the 2⁻(-Delta Delta C(T)) Method. *Methods.* 25:402–408. doi:10.1006/meth.2001.1262.
- Lu, H., and C. Klaassen. 2006. Tissue distribution and thyroid hormone regulation of *Pept1* and *Pept2* mRNA in rodents. *Peptides.* 27:850–857. doi:10.1016/j.peptides.2005.08.012.
- Matsuishi, M., M. Fujimori, and A. Okitani. 2001. Wagyu beef aroma in wagyu (Japanese Black cattle) beef preferred by the Japanese over imported beef. *Anim. Sci. J.* 72:498–504. doi:10.2508/chikusan.72.498.
- Moran, A. W., M. Al-Rammahi, C. Zhang, D. Bravo, S. Calsamiglia, and S. P. Shirazi-Beechey. 2014. Sweet taste receptor expression in ruminant intestine and its activation by artificial sweeteners to regulate glucose absorption. *J. Dairy Sci.* 97:4955–4972. doi:10.3168/jds.2014-8004.
- Nelson, G., J. Chandrashekar, M. A. Hoon, L. Feng, G. Zhao, N. J. Ryba, and C. S. Zuker. 2002. An amino-acid taste receptor. *Nature.* 416:199–202. doi:10.1038/nature726.
- Prandi, S., M. Bromke, S. Hübner, A. Voigt, U. Boehm, W. Meyerhof, and M. Behrens. 2013. A subset of mouse colonic goblet cells expresses the bitter taste receptor *Tas2r131*. *PLoS One.* 8:e82820. doi:10.1371/journal.pone.0082820.
- Rémond, D., L. Bernard, and C. Poncet. 2000. Free and peptide amino acid net flux across the rumen and the mesenteric- and portal-drained viscera of sheep. *J. Anim. Sci.* 78:1960–1972. doi:10.2527/2000.7871960x.
- Salter, A. M. 2017. Improving the sustainability of global meat and milk production. *Proc. Nutr. Soc.* 76:22–27. doi:10.1017/S0029665116000276.
- Spanier, B., and F. Rohm. 2018. Proton coupled oligopeptide transporter 1 (PepT1) function, regulation, and influence on the intestinal homeostasis. *Compr. Physiol.* 8:843–869. doi:10.1002/cphy.c170038.
- Sternini, C. 2007. Taste receptors in the gastrointestinal tract. IV. Functional implications of bitter taste receptors in gastrointestinal chemosensing. *Am. J. Physiol. Gastrointest. Liver Physiol.* 292:G457–G461. doi:10.1152/ajpgi.00411.2006.
- Tagari, H., K. Webb, Jr, B. Theurer, T. Huber, D. DeYoung, P. Cuneo, J. E. Santos, J. Simas, M. Sadik, A. Alio, et al. 2004. Portal drained visceral flux, hepatic metabolism, and mammary uptake of free and peptide-bound amino acids and milk amino acid output in dairy cows fed diets containing corn grain steam flaked at 360 or steam rolled at 490 g/L. *J. Dairy Sci.* 87:413–430. doi:10.3168/jds.S0022-0302(04)73181-1.
- Tagari, H., K. Webb, Jr, B. Theurer, T. Huber, D. DeYoung, P. Cuneo, J. E. Santos, J. Simas, M. Sadik, A. Alio, et al. 2008. Mammary uptake, portal-drained visceral flux, and hepatic metabolism of free and peptide-bound amino acids in cows fed steam-flaked or dry-rolled sorghum grain diets. *J. Dairy Sci.* 91:679–697. doi:10.3168/jds.2007-0629.
- Terada, T., Y. Shimada, X. Pan, K. Kishimoto, T. Sakurai, R. Doi, H. Onodera, T. Katsura, M. Imamura, and K. Inui. 2005. Expression profiles of various transporters for oligopeptides, amino acids and organic ions along the human digestive tract. *Biochem. Pharmacol.* 70:1756–1763. doi:10.1016/j.bcp.2005.09.027.
- Thornton, B., and C. Basu. 2011. Real-time PCR (qPCR) primer design using free online software. *Biochem. Mol. Biol. Educ.* 39:145–154. doi:10.1002/bmb.20461.
- Vancleef, L., T. Van Den Broeck, T. Thijs, S. Steensels, L. Briand, J. Tack, and I. Depoortere. 2015. Chemosensory signalling pathways involved in sensing of amino acids by the ghrelin cell. *Sci. Rep.* 5:15725. doi:10.1038/srep15725.
- Viennois, E., A. Pujada, J. Zen, and D. Merlin. 2018. Function, regulation, and pathophysiological relevance of the POT superfamily, specifically *PepT1* in inflammatory bowel disease. *Compr. Physiol.* 8:731–760. doi:10.1002/cphy.c170032.
- Wang, H. C., F. Y. Shi, M. J. Hou, X. Y. Fu, and R. J. Long. 2016. Cloning of oligopeptide transport carrier *PepT1* and comparative analysis of *PepT1* messenger ribonucleic acid expression in response to dietary nitrogen levels in yak and indigenous cattle on the Qinghai-Tibetan plateau. *J. Anim. Sci.* 94:3431–3340. doi:10.2527/jas.2016-0501.
- Zhang, Y., E. Viennois, M. Zhang, B. Xiao, M. K. Han, L. Walter, P. Garg, and D. Merlin. 2016. *PepT1* expression helps maintain intestinal homeostasis by mediating the differential expression of miRNAs along the crypt-villus axis. *Sci. Rep.* 6:27119. doi:10.1038/srep27119.
- Zhao, G. Q., Y. Zhang, M. A. Hoon, J. Chandrashekar, I. Erlenbach, N. J. Ryba, and C. S. Zuker. 2003. The receptors for mammalian sweet and umami taste. *Cell.* 115:255–266. doi:10.1016/s0092-8674(03)00844-4.