**ABSTRACT.** To achieve a better understanding of rabbit large intestinal functions, such as production of hard and soft feces and cecal fermentation, knowledge of the intestinal wall structure is essential. However, such knowledge is far from complete. Therefore, the aims of this study were to measure the thickness of the wall and its constituent layers and describe distribution of mucous cells in each segment of the large intestine in New Zealand White rabbits. Results showed that the cecum had the thinnest entire wall throughout the large intestine, and the fusus coli and rectum had a thicker entire wall in comparison to the cecum, the first segment of the proximal colon, the second segment of the proximal colon, and the distal colon. Moreover, the thickness of the mucosa in the fusus coli and that of the inner and outer layers of the tunica muscularis in the rectum were greater than that of the other segments. Mucous cells in the mucosa were the fewest in the cecum and most numerous in the fusus coli. This study provides detailed knowledge of the wall thickness and distribution of mucous cells in the large intestine of the rabbit. These findings are important for improving our understanding of rabbit intestinal physiology and pathology.

**KEY WORDS:** large intestine, mucous cell distribution, rabbit, wall thickness

Rabbits (*Oryctolagus cuniculus*) are classified as hindgut fermenters and have a highly developed and well-differentiated large intestine. The rabbit cecum is a large organ having a wide bulge and a narrow constriction, both of which run spirally from the base of the cecum toward the appendix [19]. The colon is divided into the proximal and distal parts. The proximal colon consists of three segments; the first segment (P1) has three teniae that separate the rows of haustra, the second segment (P2) has a single tenia and one row of haustra, and the third segment called the fusus coli has no teniae and haustra [8, 14, 19, 20]. The mucosal surface of P1 and P2 exhibits a pattern of protrusions into the intestinal lumen, which are somewhat larger and more regularly arranged in P1 than P2 [19]. P1 also has more numerous tubular glands as compared to P2 [19]. The mucosal surface of the fusus coli is characterized by permanent longitudinally running folds [19], and displays the greatest density of tubular glands in its thick epithelium [19, 20]. The distal colon has a smooth mucosal surface without any surface specialization and abundant mucous cells along short crypts, and continues to the rectum [19].

Extensive physiological studies have shown that each segment of the rabbit large intestine displays unique and different functions during the formation of hard and soft feces [3, 7, 8, 10, 19]. The cecum receives the ileal contents and mixes them with the contents of the cecum for microbial fermentation [4]. P1 secretes fluid from the mucosa in order to dilute the digesta, and creates muscular contractions to facilitate mechanical separation of the digesta into small fermentable particles that are transported back to the cecum, and large unfermentable fibers that are passed toward P2 [4, 15]. P2 continues further mechanical separation of 2 kinds of particles, sending the small particles back to P1, and dividing the unfermentable particles into pellets to pass them anally [10]. During hard feces production, the fusus coli creates powerful contractions using the thick tunica muscularis in order to remove water out of the fecal pellets. The distal colon gradually absorbs water further from the fecal pellets and transfers them as hard, dry, round dark pellets to the rectum to be expelled from the anus [19, 20]. In contrast, during soft feces production, the fusus coli produces soft pellets by breaking down large boluses made by P2 that receives fermented cecal contents via P1 [10]. The fusus coli also envelopes soft pellets with a thick mucus sheath secreted by its mucosa in order to facilitate their speedy transport and to protect them against gastric digestion after cecotrophy [10]. The distal colon adds extra mucus to the envelope of the soft feces [5] and rapidly transfers them through the rectum to the anus [8, 20]. These studies suggest that each segment of the rabbit large intestine, particularly the mucous and muscular layers, is important for the hard and soft feces formation. Although there have been...
a few fragmentary descriptions reported on the thickness of the entire wall in the cecum and of the mucosa in some segments [18, 19], there have been few comprehensive studies that have described in detail the thickness of the entire wall and constituent layers in each large intestinal segment. Similarly, elaborate descriptions of the distribution of the mucous cells in each segment are still insufficient. Therefore, the present study examined the wall and its constituent laminar thickness and the distribution of mucous cells in each segment of the rabbit large intestine.

MATERIALS AND METHODS

All animal experimental procedures in the present study were approved (Approval No. H28-23) by the Research Ethics Committee for Animal Experimentation of the Tokyo University of Agriculture and Technology.

In this study, 3 healthy male New Zealand White rabbits (Tokyo Laboratory Animal Science Co., Tokyo, Japan) aged 10–11 weeks (2.0–2.1 kg) were used. Until the day of sampling, all animals were kept in individual cages in an air-conditioned room with controlled temperature and free access to food and water. Prior to sacrifice, the animals were sedated with xylazine (10 mg/kg, intramuscular) and then euthanized with sodium pentobarbital (100 mg/kg, intravenous or intraperitoneal). The abdominal cavity was opened along the linea alba in order to carefully separate the large intestine from the mesentery. Samples with 2 cm length along the oro-anal axis were collected at the following sites for each segment; for the cecum, the bulging wall in the middle part along the proximodistal axis of the cecal second gyrus; for P1, 4 cm anal to the ceco-colic border; for P2, 9 cm anal to the P1-P2 border; for fusus coli, 2 cm anal to the P2-fusus coli border; for the distal colon, 40 cm anal to the fusus coli-distal colon border; and finally, for the rectum, 4 cm oral to the anus (Fig. 1). Immediately after the samples were gently washed with cold saline to remove the residues of intestinal contents, they were fixed by immersion into 4% paraformaldehyde in 0.1 M phosphate buffer for 24 hr at 4°C. Subsequently, the samples were then embedded in paraffin and cut transversely at a 6–8 µm thickness. The sections were processed with conventional hematoxylin and eosin (HE) to observe the general laminar structure and measure the thickness. Other sections were stained with Alcian blue (AB) pH 2.5, or periodic acid Schiff (PAS) or combined AB and PAS, followed by hematoxylin, to observe the distribution of mucous cells. Digital images of representative sections were taken using a DS-Ri1 camera (Nikon, Tokyo, Japan) attached to an Eclipse Ni-U microscope (Nikon). The images were adjusted to obtain adequate color, sharpness, brightness, and resolution by Adobe Photoshop (Adobe Systems, San Jose, CA, U.S.A.), and assembled into Adobe Illustrator CC (Adobe Systems).

For quantitative analysis, measurements were obtained from HE-stained sections of the bulging part in the cecum, of the tenial and haustral parts in P1 and P2, and of the non-folded part in the fusus coli, distal colon and rectum. From each of these sections, 6 sites were randomly selected to measure the thickness of the entire intestinal wall and its constituent layers (mucosa, lamina muscularis mucosae, submucosa, inner circular and outer longitudinal layer of the tunica muscularis, and serosa or adventitia), using ImageJ version 1.15s software (NIH, Bethesda, MD, U.S.A.). The thickness of the mucosa included the lamina propria. To compare the differences in the thickness between each part of the segment across 3 rabbits, the data were assembled, and the means and standard deviations were calculated using Excel version 16.20 software (Microsoft, Redmond, WA, U.S.A.). Subsequently, the difference in the total and laminar thicknesses was statistically tested by either a one-way analysis of variance and the Tukey’s test for comparison of more than 2 means or Student’s t test for comparison of 2 means, using Prism 8 version 8.1.1 software.

Fig. 1. Photograph showing approximate sampling sites (magenta bars) depicted in an unfolded rabbit large intestine. Arrowheads indicate the borders of each segment.
significantly thinner lamina muscularis mucosae than the rectum (114.6 ± 20.9 µm) compared to all of the segments other than the fusus coli (Fig. 3b). However, the fusus coli (Fig. 2f), the mucosa was the thickest throughout the large intestine and the tunica muscularis was relatively well-developed. The distal colon, particularly its mucosa, was thinner than the fusus coli (Fig. 2g), and the rectum had the thickest entire wall, lamina muscularis mucosae and tunica muscularis across the large intestine (Fig. 2h).

Quantitative analysis more clearly showed the difference in the thickness of the entire wall and the constituent layers of each segment. The entire intestinal wall in the cecum was significantly thinner (299.7 ± 32.8 µm) and that in the fusus coli (996.9 ± 168.0 µm) and rectum (1,048.1 ± 108.4 µm) was significantly thicker, compared to the other segments (Fig. 3a and Table 1). In-between were P1 (680.6 ± 117.0 µm), P2 (676.7 ± 123.4 µm) and the distal colon (626.4 ± 115.9 µm), whose thickness of the entire wall did not significantly differ from each other (Fig. 3a and Table 1).

For the thickness of the mucosa, the cecum was significantly thinner (136.8 ± 23.0) and the fusus coli was significantly thicker (627.7 ± 129.9 µm) as compared to the other segments. Moreover, P1 had a significantly thicker mucosa, compared to all of the segments other than the fusus coli (Fig. 3b).

For the lamina muscularis mucosae (Fig. 3c and Table 1), the thickness did not differ among the cecum (7.5 ± 2.4 µm), P1 (11.5 ± 6.1 µm) and P2 (12.3 ± 4.3 µm), all of which had a significantly thinner lamina muscularis mucosae than the distal colon and rectum. The mucosa and P1 also had a significantly thinner lamina muscularis mucosae than the fusus coli (Fig. 3c). The fusus coli (20.8 ± 16.2 µm) had a significantly thinner lamina muscularis mucosae than the distal colon (54.2 ± 9.8 µm), which, in turn, had a significantly thinner lamina muscularis mucosae than the rectum (114.6 ± 20.9 µm) (Fig. 3c and Table 1).

The thickness of the submucosa did not have any significant difference among others in the cecum (33.1 ± 13.5 µm), P1 (34.5 ± 17.9 µm), P2 (37.1 ± 28.7 µm), the fusus coli (41.5 ± 23.0 µm) and the distal colon (49.2 ± 12.0 µm) (Fig. 3d and Table 1). However, the rectum (72.3 ± 31.3 µm) had a significantly thicker submucosa (Fig. 3d and Table 1), compared to all of the other segments.

For both layers of the tunica muscularis, the cecum was significantly thinner (63.8 ± 18.9 µm) for the inner circular and 41.9 ± 10.4 µm for the outer longitudinal layer) in comparison to the other segments except for the outer longitudinal layer of the distal colon (Fig. 3e, 3f and Table 1). On the contrary, both layers of the rectum were significantly thicker (350.8 ± 80.7 µm for the inner circular and 208.6 ± 38.5 µm for the outer longitudinal layer), in comparison to those of the other segments (Fig. 3e, 3f and Table 1). There was a second peak of the thickness of the tunica muscularis in the proximal colon along the oro-anal axis of the large intestine (Fig. 3e, 3f and Table 1), where P2 (255.5 ± 72.4 µm) was the second thickest for the inner circular layer (Fig. 3e and Table 1), whereas P1 (117.7 ± 64.2 µm) was the second thickest for the outer longitudinal layer (Fig. 3f and Table 1).

Finally, the cecum (16.6 ± 6.7 µm), P1 (18.0 ± 7.7 µm) and P2 (14.9 ± 9.6 µm) had significantly or tended to have a thicker serosa than the fusus coli (9.6 ± 9.5 µm) and distal colon (7.1 ± 4.3 µm) (Fig. 3g and Table 1).

The thickness of the entire wall and its constituent layers was also compared between the tenial and hastral part of P1 (Fig. 4 and Table 2) and between the tenial and hastral part of P2 (Fig. 5 and Table 3). For P1, the tenial part (765.1 ± 92.2 µm) had a significantly thicker entire wall than the hastral part (596.1 ± 67.7 µm) (Fig. 4a and Table 2). The inner circular and outer longitudinal layers were significantly thicker in the tenial (143.9 ± 22.9 µm and 152.3 ± 22.9 µm, respectively) than in the hastral part (120.3 ± 31.9 µm and 14.2 ± 4.3 µm, respectively) (Fig. 4e, 4f, and Table 2), whereas the serosa was significantly thicker in the hastral (19.9 ± 9.8 µm) than the tenial part (16.0 ± 4.2 µm) (Fig. 4g and Table 2). The thickness of the mucosa, lamina muscularis mucosae and submucosa did not have any significant difference between the tenial and hastral part (Fig. 4b–d, and Table 2).

For P2, there was no significant difference in the thickness of the entire wall between the tenial (715.0 ± 132.0 µm) and hastral part (638.4 ± 104.0 µm) (Fig. 5a and Table 3). However, the mucosa was significantly thicker in the tenial part (333.2 ± 70.8 µm) than in the hastral part (282.4 ± 46.4 µm) (Fig. 5b and Table 3), whereas the inner circular layer was significantly thinner in the tenial part (219.6 ± 53.5 µm) than in the hastral part (291.5 ± 72.1 µm) (Fig. 5c and Table 3). The outer longitudinal layer of the tunica muscularis was observed only at 1 out of 18 sites in the hastral part, therefore, statistical test was not able to be performed. There were no significant differences in the thickness of the lamina muscularis mucosae, submucosa and serosa between the tenial and hastral part (Fig. 5c, 5d, 5f, and Table 3).

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Fig. 2. Cross sections of the cecum (a), tenial part of P1 (b), haustral part of P1 (c), tenial part of P2 (d), haustral part of P2 (e), fusus coli (f), distal colon (g) and rectum (h), demonstrating the difference in the thickness of the entire wall and the constituent layers between each segment. HE staining. Scale bar, which is applicable to all panels, equals 100 μm.
Sections stained with AB, PAS and AB-PAS showed that the distribution of mucous cells was remarkably different along the oro-anal axis of the large intestine (Figs. 6 and 7). In general, mucous cells were the fewest in the cecum and most numerous in the fusus coli. Most of the individual mucous cells were stained with both AB and PAS, unless otherwise mentioned below.

The cecum had a few mucous cells, which rarely made clusters and resided in the epithelium facing the intestinal lumen and lining the shallow intestinal crypt (arrowheads in Fig. 6a, a’ and a”). In P1, the staining pattern was similar in the tenial and haustral parts.

Numerous mucous cells were present in the epithelium lining the crypt, particularly in the deep half (Fig. 6b, b’ and b”).

### Table 1. Laminar and entire wall thickness (mean ± SD in µm) of each segment of the rabbit large intestines

| Layer                                | Cecum     | P1        | P2        | Fusus coli | Distal colon | Rectum   |
|--------------------------------------|-----------|-----------|-----------|------------|--------------|----------|
| Mucosa                               | 136.8 ± 23.0 | 406.2 ± 59.6 | 307.8 ± 64.4 | 627.7 ± 129.9 | 313.3 ± 59.4 | 279.6 ± 38.8 |
| Lamina muscularis mucosae            | 7.5 ± 2.4  | 11.5 ± 6.1  | 12.3 ± 4.3  | 20.8 ± 16.2  | 54.2 ± 9.8  | 114.6 ± 20.9 |
| Submucosa                            | 33.1 ± 13.5 | 34.5 ± 17.9  | 37.1 ± 28.7  | 41.5 ± 23.0  | 49.2 ± 12.0  | 72.3 ± 31.3 |
| Tunica muscularis, inner circular    | 63.8 ± 18.9 | 132.1 ± 29.9 | 255.5 ± 72.4 | 203.9 ± 52.2 | 157.2 ± 52.8 | 350.8 ± 80.7 |
| Tunica muscularis, outer longitudinal| 41.9 ± 10.4 | 117.7 ± 64.2 | 92.7 ± 28.5  | 93.7 ± 30.8  | 45.4 ± 14.2  | 208.6 ± 38.5 |
| Serosa                               | 16.6 ± 6.7  | 18.0 ± 7.7  | 14.9 ± 9.6  | 9.6 ± 9.5   | 7.1 ± 4.3  | 22.4 ± 8.8 |
| Total                                | 299.7 ± 32.8 | 680.6 ± 117.0 | 676.7 ± 123.4 | 996.9 ± 168.0 | 626.4 ± 115.9 | 1,048.1 ± 108.4 |

n=18 for each value. For P1 and P2, n=36, because their values are calculated across the tenial and haustral parts. For the outer longitudinal layer of the tunica muscularis in P1, n=24 and that in P2, n=19, because of the occasional absence of the longitudinal muscles. Note that for the serosa in the rectum, the value is of the adventitia.

### Mucous cell distribution

Sections stained with AB, PAS and AB-PAS showed that the distribution of mucous cells was remarkably different along the oro-anal axis of the large intestine (Figs. 6 and 7). In general, mucous cells were the fewest in the cecum and most numerous in the fusus coli. Most of the individual mucous cells were stained with both AB and PAS, unless otherwise mentioned below.

The cecum had a few mucous cells, which rarely made clusters and resided in the epithelium facing the intestinal lumen and lining the shallow intestinal crypt (arrowheads in Fig. 6a, a’ and a”). In P1, the staining pattern was similar in the tenial and haustral parts. Numerous mucous cells were present in the epithelium lining the crypt, particularly in the deep half (Fig. 6b, b’ and b”), and a small
number of mucous cells were also located in the surface epithelium. There was a tendency that cells stained with AB alone were distributed more in the vicinity of the neck of the crypt (Fig. 6b and b’’), while those stained with PAS alone were distributed more in the bottom of the crypt (Fig. 6b’ and b’’). There was another tendency that cells stained with PAS were darker in the deep crypt than in the superficial crypt and surface epithelium. In P2, the staining pattern was similar in the tenial and haustral parts. Mucous cells in P2 were also numerous and were distributed in a manner similar to those in P1 (Fig. 6c, c’ and c’’). In the thick mucosa of the fusus coli, mucous cells were the most numerous and tended to be aggregated in the epithelium that lined the deep half of the intestinal crypt, while they were somewhat more diffusely distributed in the epithelium that lined the superficial half of the crypt.

Table 2. Laminar and entire wall thickness (mean ± SD in µm) of the tenial and haustral part of P1

| Layer                              | Tenial part     | Haustral part    |
|------------------------------------|-----------------|------------------|
| Mucosa                             | 401.8 ± 51.9    | 410.6 ± 67.7     |
| Lamina muscularis mucosae          | 11.7 ± 6.1      | 11.3 ± 6.2       |
| Submucosa                          | 39.5 ± 21.9     | 29.5 ± 11.2      |
| Tunica muscularis, inner circular  | 143.9 ± 22.9    | 120.3 ± 31.9     |
| Tunica muscularis, outer longitudinal | 152.3 ± 22.9   | 14.2 ± 4.3       |
| Serosa                             | 16.0 ± 4.2      | 19.9 ± 9.8       |
| Total                              | 765.1 ± 92.2    | 596.1 ± 67.7     |

n=18 for each value except for the outer longitudinal layer of the tunica muscularis of the haustral part, where n=6. This is due to the occasional absence of the longitudinal muscles.

Table 3. Laminar and entire wall thickness (mean ± SD in µm) of the tenial and haustral part of P2

| Layer                              | Tenial part     | Haustral part    |
|------------------------------------|-----------------|------------------|
| Mucosa                             | 333.2 ± 70.8    | 282.4 ± 46.4     |
| Lamina muscularis mucosae          | 12.3 ± 3.2      | 12.4 ± 5.2       |
| Submucosa                          | 41.3 ± 13.1     | 32.9 ± 38.6      |
| Tunica muscularis, inner circular  | 219.6 ± 53.5    | 291.5 ± 72.1     |
| Tunica muscularis, outer longitudinal | 95.9 ± 25.3    | NA               |
| Serosa                             | 12.5 ± 4.1      | 17.4 ± 12.7      |
| Total                              | 715.0 ± 132.0   | 638.4 ± 104.0    |

n=18 for each value except for the outer longitudinal layer of the tunica muscularis in the haustral part, where the longitudinal muscles were observed at only 1 site (33.5 µm). Therefore, the value is not applicable (NA) for calculation.

Fig. 4. Columnar graphs showing the statistical difference in the thickness of the entire wall (a), mucosa (b), lamina muscularis mucosae (c), submucosa (d), inner circular layer of the tunica muscularis (e), outer longitudinal layer of the tunica muscularis (f) and serosa (g) between the tenial and haustral part of P1 across 3 rabbits. The height and whisker of each column represent the mean ± SD. n=18 except for the outer longitudinal layer of the haustral part, where n=6. The X-axis indicates the intestinal segment, whereas the Y-axis indicates the thickness in µm. *, P<0.05. P1-t, the tenial part of P1 and P1-h, haustral part of P1.
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(Fig. 7a, a’ and a”). A small number of mucous cells were also located in the surface epithelium. The distal colon and rectum (Fig. 7b, b’, b”, 7c, c’ and c”) had a moderate number of mucous cells that were aggregated in the epithelium lining the crypt, while the surface epithelium had fewer, scattered mucous cells. Mucous cells stained with PAS were distributed more densely in the distal colon than the rectum (Fig. 7b’ and c’), however, these cells were fewer than the mucous cells stained with AB in both the distal colon and rectum (Fig. 7b, b’, 7c and c’).

DISCUSSION

This is the first systematic study documenting the thickness of the wall and the mucous cell distribution throughout all segments of the rabbit large intestine. Our findings demonstrated that the cecum had the thinnest entire wall across the large intestine. The thickness that we observed (299.7 ± 32.8 µm) was remarkably thinner as compared to the 554.5 µm thickness reported by Snipes [18]. The reason for the difference between the findings of Snipes [18] and our current study may be attributable to several factors such as differences in the strain, age, and numbers of the rabbit, in addition to the sampling sites, tissue processing, and measurement methods. This may also explain the difference in the thickness values of other segments between the findings of Snipes and the present study [18, 19]. The cecum has the thinnest mucosa in agreement with a previous study [18], and the mucosal epithelium contains only a few mucous cells. This indicates that most of the epithelial cells may consist of absorptive cells, which are advantageous for absorption of electrolytes and volatile fatty acids through the cecal mucosa [6, 8, 14]. The cecum also has the thin tunica muscularis, which does not so often create the wall contraction [17]. It is, therefore, suggested that the ingesta in the cecum may be gently mixed for fermentation that takes place within the cecum.

The present findings showed that the entire wall of P1 was as thick as that of P2 and the distal colon. Within P1, the tenial wall is thicker than the haustral wall. This is attributable to the thicker inner circular and outer longitudinal layers of the tunica muscularis in the tenial part and the occasional absence of the outer longitudinal layer in the haustral part. The mucosa of P1 was the second thickest across all of the segments, in agreement with the results reported by Snipes et al. [19]. In the mucosa, numerous mucous cells located in the crypt may add mucus to lubricate the transport of intestinal contents going back to the cecum.
and forward to P2 [2, 10]. The mucous cells stained with AB tended to be arranged more superficially in the crypt than those stained with PAS. This arrangement appears beneficial to resist against microbial invasion through the epithelium, since acidic mucins secreted by AB-stained mucous cells may be resistant to microbial degradation of the intestinal mucosa [9, 16]. The paucity of mucous cells in the surface and superficial crypt epithelium, that is, rich absorptive cells in these sites, may be advantageous for effective electrolyte absorption [19]. Also of particular interest is that P1 has a well-developed longitudinal outer muscular layer of the tunica muscularis, but has a poorly developed circular layer, particularly in the haustral part. This muscular arrangement may be important to facilitate separation of digesta into small fermentable particles and unfermentable fibers for soft and hard feces formation, although the exact mechanisms of such separation are still unknown.

The mucosa in P2 was somewhat thinner than that in P1. Within P2, the mucosa of the tenial part was thicker than that of the haustral part. The distribution of mucous cells in both parts of P2 was essentially similar to that in P1. For the tunica muscularis, the inner circular layer in P2 was the second thickest across the large intestine. Since P2 is the only segment where the formation of hard fecal pellets takes place [10], it is considered that the presence of the thick circular layer in P2 is important for creating

Fig. 6. Cross sections of the mucosa of the cecum (a, a' and a''), tenial part of P1 (b, b' and b'') and tenial part of P2 (c, c' and c''), demonstrating the difference in the distribution of mucous cells in each segment. Alcian blue (a, b and c), periodic acid Schiff (a', b' and c') and combined Alcian blue-periodic acid Schiff staining (a'', b'' and c''). Arrowheads in (a), (a') and (a'') indicate a few mucous cells. Scale bar, which is applicable to all panels, equals 100 µm.
strong segmental contractions in order to separate the solid contents into pellets, and for carrying the liquid contents back towards the cecum [10].

The fusus coli had the thickest entire wall among each segment of the large intestine, except for the rectum that was as thick as

Fig. 7. Cross sections of the fusus coli (a, a' and a''), distal colon (b, b' and b'') and rectum (c, c' and c''), demonstrating the difference in the distribution of mucous cells in each segment. Alcian blue (a, b and c), periodic acid Schiff (a', b' and c') and combined Alcian blue-periodic acid Schiff staining (a'', b'' and c''). Scale bar, which is applicable to all panels, equals 100 μm.
the fusus coli. The thickness of the fusus coli was due to the thickest mucosa across the large intestine. Our data of mucous cell-staining demonstrated that the fusus coli had the most numerous mucous cells compared to the other segments. The mucus secreted by these cells may lubricate the mucosal surface for facilitation of speedy transport of soft fecal pellets and to envelop them to be protected against digestion after cecotrophy [19, 20]. The paucity of the mucous cells in the superficial part of the crypt epithelium indicates that this part may contain more absorptive cells, which may work to remove water out of the fecal pellets [19, 20]. In the fusus coli, the circular and longitudinal layers of the tunica muscularis were relatively well-developed. It is assumed that the contraction of the tunica muscularis of the fusus coli forms the small soft fecal pellets. In addition, it has also been postulated that during the hard feces formation, the contraction of the tunica muscularis may extrude water out of the hard pellets [10, 19].

The entire wall of the distal colon is relatively well-developed. In the mucosa, the mucous cells stained with AB was more numerous than those stained with PAS. As discussed above, this may be beneficial to protect the mucosa against microbial invasion [9, 16]. The distal colon has a thick inner circular layer of the tunica muscularis relative to the outer longitudinal layer. This may be advantageous for speedy transport of soft feces towards the anus and extrusion of water from hard feces, as the fusus coli may also do [10, 20].

The rectum was shown to have the thickest entire wall across all of the other segments except for the fusus coli. In the mucosa, the distribution of the mucous cells in the rectum was similar to that in the distal colon. The rectum had the thickest lamina muscularis mucosae and circular and longitudinal layers of the tunica muscularis across all of the other segments. In rabbits, soft feces are expelled once or twice a day while hard pellets are often expelled throughout the day [20]. Therefore, the presence of a thick lamina muscularis mucosae and tunica muscularis in this segment may be an advantageous benefit for adapting to frequent defecation.

The results of our current study demonstrate the anatomical difference in the thickness of the wall and its layers as well as the distribution of mucous cells in the mucosa in each segment of the rabbit large intestine. Such structural difference is implicated in different functional roles played by each segment for formation of soft and hard feces. The present findings are important as a basis for understanding physiology of the rabbit large intestine and its pathological conditions, such as bowel inflammatory diseases [1, 11] and intestinal microbial infections [12, 13], where the normal structure of intestine may be altered.

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