

Expressional Patterns of Connexin Isoforms in the Rat Epididymal Fat during Postnatal Development

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ABSTRACT : In the multicellular tissue, cell-cell interaction is important for a precise control of its function. The exchange of signaling molecules between adjacent cells via connexon allows the functional harmony of cells in the tissue. The present research was to determine the presence and expressional patterns of connexin (Cx) isoforms in the rat epididymal fat during postnatal development using quantitative real-time polymerase chain reaction (PCR) analysis. Of 13 Cx isoforms examined, expression of 11 Cx isoforms in the epididymal fat during postnatal development was detected. These Cx isoforms include Cx26, Cx31, Cx31.1, Cx32, Cx33, Cx36, Cx37, Cx40, Cx43, Cx45, and Cx50. Expressional levels of all Cx isoforms at 1 and 2 years of age were significantly higher than those at the early postnatal ages, such as 7 days, 14 days, and 24 days of ages. Except Cx33 and Cx43, the transcript levels of rest Cx isoforms at 1 year of age were significantly lower than that at 2 years of age. In addition, expressional patterns of Cx isoforms between 7 days and 5 months of ages generally varied according to the isoform. The existence of various Cx isoforms in the rat epididymal fat has been identified and expression of each Cx isoform in the epididymal fat during postnatal development has shown a particular pattern, distinguishable from the others. To our knowledges, this is the first report showing expressional patterns of Cx isoforms at transcript level in the epididymal fat at various postnatal ages.

Key words : Connexin, Quantitative real-time polymerase chain reaction (PCR), Epididymal fat, Postnatal development, Gene expression

INTRODUCTION

Adipose tissue is composed of various cell types, including adipocyte (fat cell), macrophages, endothelial cells, fibroblasts, and adipose precursor cells (Frühbeck, 2008). Adipose tissue is commonly found in all mammalian species as well as a number of non-mammalian species (Hausman et al., 2001). Formation of adipose tissue in mammals begins *in utero* and continues throughout life time. Based on histological and molecular characteristics, mammalian adipose tissue is separated into two types, white adipose

tissue (WAT) and brown adipose tissue (BAT) (Berry et al., 2013). It is generally considered that WAT is specialized for energy storage and BAT is chiefly involved in energy consumption and heat generation (Berry et al., 2013). In spite of the presence of functional and morphological differences, adipocytes in WAT and BAT are originated from a common stellate or fusiform precursor cells derived from mesenchymal cells (Frühbeck, 2008).

Adipose tissue usually exists in a depot form, and there are two classes of WAT, subcutaneous and visceral WAT, according to regional, developmental timing, and molecu-

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lar features, and biological functions (Berry et al., 2013). Visceral WAT is further divided into retroperitoneal, perigonadal, and mesenteric WATs, which have specific and distinct morphology and texture (Berry et al., 2013). The epididymal fat depot is included in perigonadal WAT and embeds the efferent ductules located between the testis and the epididymis. In the rat, the weight of epididymal fat during postnatal development becomes remarkably increased in several hundred-fold (Hubbard & Matthew, 1971; Cleary et al., 1977). The increase of epididymal fat weight with aging is due to not only an increase of fat cell number but also an increase of fat cell size (Cleary et al., 1977). In addition to the changes in fat cell, weight of non-lipid filled cells in the epididymal fat during postnatal development is also increased at a lower rate than that of fat cell (Cleary et al., 1977). Even though the gross change of epididymal fat from the early pubertal age to the adult age has been briefly studied, the change of epididymal fat at the prepubertal and/or the elderly age has not been examined yet.

Cell-cell communication within multicellular tissue is an important biological phenomenon to permit a functional accordance of tissue. Cellular communication is usually accomplished by three junctional complexes, including tight junction, adherens junction, and gap junction (Cyr, 2011). Of these junctional complexes present within multicellular tissue, gap junction plays various important roles by mediating direct communication between adjacent cells (Cyr, 2011). Gap junction is consisted of homomeric or heteromeric hemichannel, called connexon, and each connexon is made of six connexin (Cx) molecules (Cyr, 2011). Direct exchanges of small molecules, such as signaling particles, metabolites, and even ribonucleic acid (RNAs), between neighboring cells are permitted through connexon (Goodenough et al., 1996). There are 20 Cx isoforms identified in mammals to date, and expression of Cx isoforms in various tissues has been examined from a number of researches (Goodenough et al., 1996; Willecke et al., 2002). Some Cx isoforms are present in a broad range of tissues

and/or cell types, while expression of certain Cx isoforms is restricted into specific cell types (Pointis et al., 2005). Even though it is not largely examined, expression of Cx isoforms and possible role of Cx isoforms in fat tissues have been studied (Burke et al., 2014; Kim et al., 2017). Among the male reproductive tract, expression of Cx isoforms in the testis and epididymis has been extensively evaluated by other and our previous researches (Hejmej et al., 2007; Han & Lee, 2013). However, it is hard to find any report on the presence and/or expression of Cx isoforms in the epididymal fat.

Thus, the present research was focused to examine the presence of Cx isoforms in the rat epididymal fat at messenger RNA (mRNA) level and, if so, expressional patterns of Cx isoforms during postnatal development from the neonatal age to extremely old age.

MATERIALS AND METHODS

1. Experimental animals and collection of the tissue

Male pups of Sprague Dawley rat were obtained from pregnant female rats (n=10) caged individually upon the delivery (Samtako, OSan, Korea). The neonatal and young male rats at 1 week (n=10), 14 days (n=10), and 24 days (n=8) were used for the present research. Male rats at 44 days of age (n=7) were purchased from Samtako. These animals were allowed free access to food and water throughout the experiment. Male Sprague Dawley rats at 5 months (n=5), 1 year (n=3), and 2 years (n=3) of ages were kindly supported by Aging Tissue Bank (Department of Pharmacology, Pusan National University, S. Korea).

To collect the epididymal fat, the experimental animal was anesthetized by CO₂ stunning, and an opening was made on lower abdominal area by a pair of scissors. The entire reproductive tract, including epididymal fat, testis, and epididymis, was dissected out and rapidly placed in cold-PBS (phosphate-buffered saline) containing dish. The epididymal fat was separated from the rest of reproductive

tract, and the efferent ductules embedded within the epididymal fat were carefully removed. The fat was quickly washed with a fresh cold-PBS and frozen in liquid nitrogen. The fat tissue stored in -80°C was used for total RNA isolation within a week after the tissue isolation.

2. Isolation of total RNA from the epididymal fat and construction of the first-stranded complementary DNA (cDNA)

About 20 mg of the epididymal fat tissue was homogenized in easy-Blue total RNA extract solution (iNtRON Biotech, Sungnam, S. Korea) by using a polytron homogenizer (Fisher Scientific, Pittsburgh, USA). Total RNA pellet was collected by the sequential addition of chloroform and isopropanol, and an air-dried pellet was resuspended in DEPC-dH₂O. The qualitative and quantitative evaluations of isolated total RNA were carried out by 2% agarose gel electrophoresis and a NanoDrop Lite spectrophotometry (Thermo Scientific, Wilmington, DE), respectively. The total RNA was stored in -80°C until used for reverse transcription (RT) reaction.

The ImProm-IITM reverse transcription system (Promega, Madison, USA) was used to perform the RT reaction with 1 μg of total RNA. A RT mixture having total RNA, oligo-dT primer, dNTPs, RTase, and buffer in a total volume of 20 μL was exposed to 25°C for 5 min, followed by an incubation at 42°C for 90 min and 70°C for 15 min. Generated cDNA was immediately used for quantitative real-time polymerase chain reaction (PCR) analysis.

3. Quantitative real-time polymerase chain reaction analysis and data analysis

Table 1 shows the oligonucleotide primers examined in the present research for real-time PCR analysis. The primer design was performed with Primer 3 software (<http://www.bioinformatics.org/cgi-bin/primer3.cgi>: Whitehead Institute/MIT Center for Genomes Research, USA). Initially, oligonucleotide primers of 13 *Cx* isoforms were designed and

tested for the expression in the epididymal fat. However, expression of *Cx30* and *Cx30.1* in the epididymal fat was not detected in the epididymal fat throughout the postnatal development. Thus, the sequence information of *Cx30* and *Cx30.1* was excluded from Table 1.

To perform quantitative real-time PCR, 1 μL of cDNA was mixed with 10 pmol of each primer, 7 μL of iQTM SYBR[®] Green Supermix (Bio-Rad Laboratories, Hercules, CA), and DNase (deoxyribonuclease)-free dH₂O to make a final volume of 20 μL . The mixture was allowed for a pre-denaturation step at 95°C for 5 min. Then, the PCR procedure was carried out in a thermocycler (Bio-Rad Laboratories) with cycles of denaturation at 95°C for 30 sec, annealing at T_m for 30 sec, and extension at 72°C for 30 sec. At the last, an extension step at 72°C for 10 min was added to each PCR. The size of expected PCR product was checked by 1.2% agarose gel electrophoresis. Beta-actin (*Actb*) was included as an internal PCR control.

Independent quadruplication of RT reaction and PCR for each postnatal age was executed to obtain a mean and a standard error. The PCR result was presented in the relative ratio of expressional level between *Actb* and *Cx* isoform. Statistical comparison among the transcript level of different postnatal ages for each *Cx* isoform was achieved by one-way ANOVA, followed by Duncan's test, a post-hoc analysis. If $p < 0.05$, means between age groups were judged as statistically significant.

RESULTS

1. Expressional patterns of *Cx26* and *Cx31* expression in the epididymal fat of rat during postnatal period

Expressional level of *Cx26* in the epididymal fat of rat at 7 days of postnatal age was not significantly changed until 45 days of age, while an expressional surge of *Cx26* was detected at 5 months of age (Fig. 1A). The expressional level of *Cx26* at 1 year of age was about 4 times higher than that at 5 months of age (Fig. 1A). Compared with the

Table 1. Information for oligonucleotide primers utilized for quantitative real-time PCR analysis

Gene	Primer sequence (5' → 3')	T _m (°C)	PCR product size (bps)
<i>Gjb2</i> (Cx26) (NM_001004099)	(F) TCCTCTTCATCTTCCGCATC (R) CCGTTTCTTTTCGTGTCTCC	55	233
<i>Gjb3</i> (Cx31) (NM_019240)	(F) TTGAGCGGTGTGAACCAGTA (R) TGTTGGAGATGGGGAAGAAG	58	193
<i>Gjb5</i> (Cx31.1) (NM_019241)	(F) CATCGTCTGCATCCTGCTTA (R) ATGAGGTTCGCTTGAGAGGAA	55	165
<i>Gjb1</i> (Cx32) (NM_017251)	(F) AGAATCATGGTGCTGGTGGT (R) CCTCAAGCCGTAGCATTTTC	58	235
<i>Gja6</i> (Cx33) (NM_019308)	(F) TGAGAGGCAGATTGCTGCTA (R) AGACACCATTGACACCACCA	58	221
<i>Gjd2</i> (Cx36) (NM_019281)	(F) TCTGGAGATTGGGTTTCTGG (R) CGGACAGCCAGTTTGATCTT	58	231
<i>Gja4</i> (Cx37) (NM_021654)	(F) AGTGTCTGTACCTTGATGCC (R) CAGCACACTTAGCCAAGAGC	55	223
<i>Gja5</i> (Cx40) (NM_019280)	(F) ATACCATTGAGCCTGGTTGC (R) CGGCCTCTTTAGCTTTCTCA	58	189
<i>Gja1</i> (Cx43) (NM_012567)	(F) AGCAAGCTAGCGAGCAAAC (R) GAGTTCATGTCCAGCAGCAA	55	151
<i>Gja7</i> (Cx45) (NM_001085381)	(F) GATCATCCTGGTTGCTACTC (R) GATCCTCTTCATGGTCCTCT	58	173
<i>Gja8</i> (Cx50) (NM_153465.2)	(F) CCACTCCATTGCAGTTTCCT (R) AGAAGGCAGGGTTTCTTGGT	58	211
<i>Actb</i> (NM_031144.3)	(F) ACCTTCAACACCCCAGCCATGTACG (R) CTGATCCACATCTGCTGGAAGGTGG	55	698

PCR, polymerase chain reaction, *Cx*, connexin; *Gja*, gap junction protein, alpha; *Gjb*, gap junction protein, beta; *Gjd*, gap junction protein, delta; *Actb*, actin, beta.

level of *Cx26* mRNA at 1 year of age, over 7.5-fold increase of *Cx26* transcript level was observed at 2 years of postnatal age (Fig. 1A).

Expression of *Cx31* was first significantly increased at 14 days of age, and the level of *Cx31* transcript was remained

in steady until 24 days of age (Fig. 1B). However, the level of *Cx31* mRNA was significantly decreased and became the lowest level during postnatal development (Fig. 1B).

Expressional level of *Cx31* at 5 months of age was higher than 14 and 24 days of ages, and a huge increase of *Cx31*

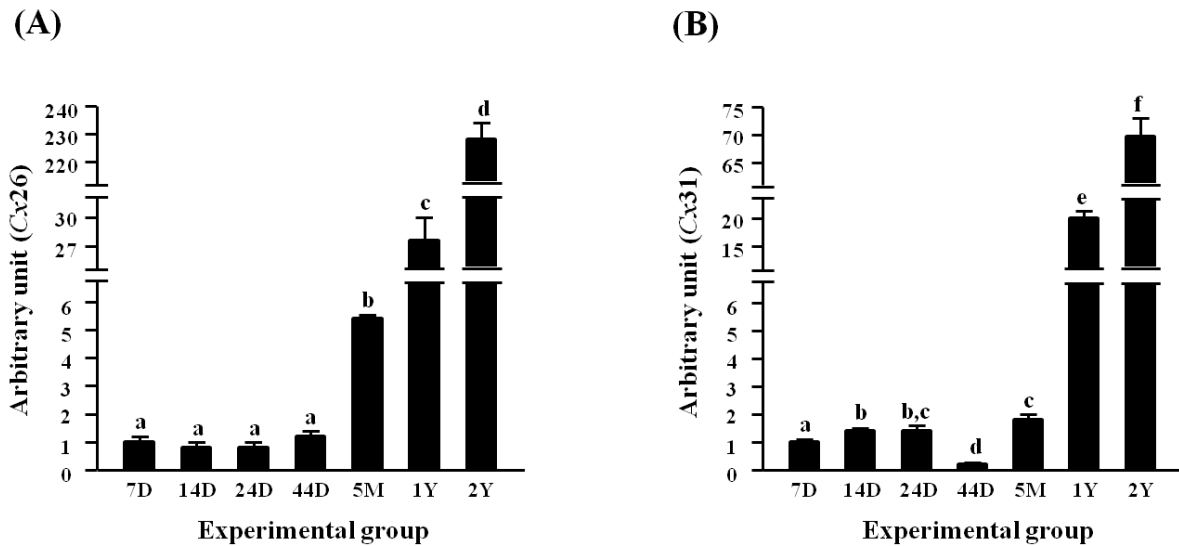


Fig. 1. Expressional patterns of Cx26 and Cx31 in the rat epididymal fat during postnatal period. The relative expressional levels of Cx26 (A) and Cx31 (B) are shown here. Different letters indicate statistical significances at $p < 0.05$. Cx, connexin; D, day; M, month; Y, year.

expression was detected at 1 year of age (Fig. 1B). The level of Cx31 transcript at 2 years of age was more than 3 times higher than that at 1 year of age (Fig. 1B).

2 Expressional patterns of Cx31.1 and Cx32 expression in the epididymal fat of rat during postnatal period

There was no significant change of Cx31.1 transcript level until 5 months of age (Fig. 2A). However, a tremendous increase, about 30 times, of Cx31.1 mRNA level was detected at 1 year of age, followed by an additional increase of the transcript level at 2 years of age (Fig. 2A).

A significant increase of Cx32 transcript level was observed at 14 days of age, followed by a drop at 24 days of age to the level of Cx32 mRNA at 7 days of postnatal age (Fig. 2B). No significant change of Cx32 transcript level was detected until 44 days of age (Fig. 2B). However, the level of Cx32 mRNA was significantly increased at 5 months of age (Fig. 2B). Expression of Cx32 in the epididymal fat was significantly increased at 1 year of age, compared with that at 5 months of age (Fig. 2B). A vast increase, over 25-fold, of Cx32 transcript level at 2 years of age was followed (Fig. 2B).

3. Expressional patterns of Cx33 and Cx36 expression in the epididymal fat of rat during postnatal period

Expressional level of Cx33 in the rat epididymal fat was first significantly increased at 14 days of age, and an additional increase of Cx33 transcript level was observed at 44 days of postnatal age (Fig. 3A). A further induction of Cx33 expression was found at 5 months of age, followed by mostly 5-fold increase of Cx33 transcript level at 1 year of age (Fig. 3A). But, the level of Cx33 transcript at 2 years of postnatal age was significantly decreased (Fig. 3A).

There was no significant change of Cx36 transcript level until 24 days of age (Fig. 3B). However, an increase of Cx36 mRNA level was observed at 44 days of age, and the level of Cx36 transcript at 5 months of age was significantly higher than that at 44 days of age (Fig. 3B). A great jump of Cx36 transcript level was detected at 1 year of age, followed by a further significant increase of the level at 2 years of age (Fig. 3B).

4. Expressional patterns of Cx37 and Cx40 expression in the epididymal fat of rat during postnatal period

Fig. 4A shows the expressional pattern of Cx37 in the rat

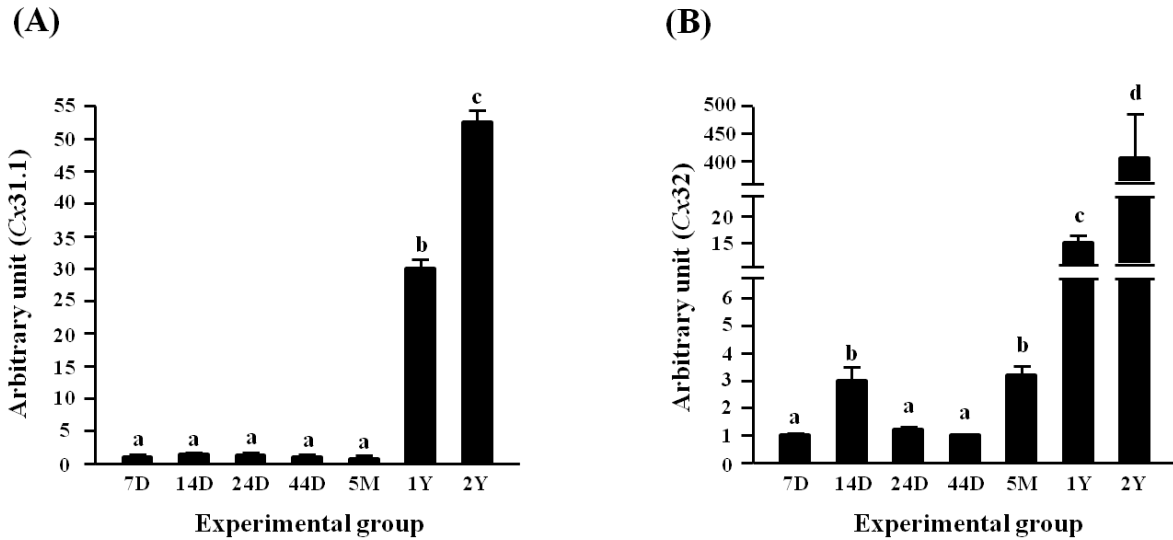


Fig. 2. Expressional patterns of Cx31.1 and Cx32 in the rat epididymal fat during postnatal period. The relative expressional levels of Cx31.1 (A) and Cx32 (B) are shown here. Different letters indicate statistical significances at $p < 0.05$. Cx, connexin; D, day; M, month; Y, year.

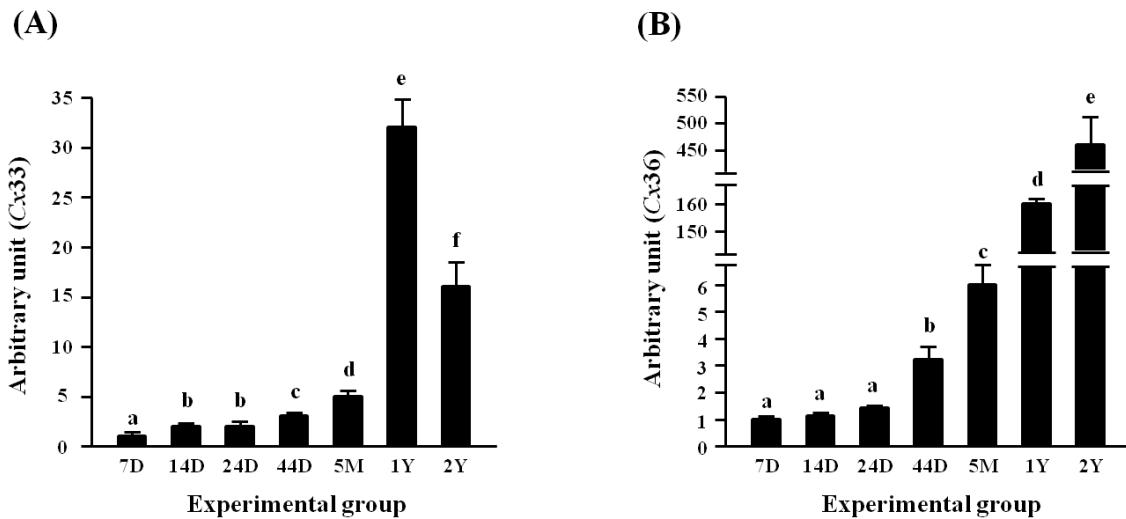


Fig. 3. Expressional patterns of Cx33 and Cx36 in the rat epididymal fat during postnatal period. The relative expressional levels of Cx33 (A) and Cx36 (B) are shown here. Different letters indicate statistical significances at $p < 0.05$. Cx, connexin; D, day; M, month; Y, year.

epididymal fat during postnatal development. The levels of Cx37 transcript at 14 and 24 days of age were significantly higher than that at 7 days of age (Fig. 4A). But, transcript level of Cx37 at 44 days and 5 months of age was lower than those at 14 and 24 days of age but still higher than that at 7 days of age (Fig. 4A). An increase of Cx37 mRNA level was found at 1 year of age, and a rise of Cx37 tran-

script level was additionally detected at 2 years of age (Fig. 4A).

The level of Cx40 transcript in the rat epididymal fat was significantly induced at 14 days of age, compared with that at 7 days of age (Fig. 4B). Another significant increase of Cx40 mRNA level was detected at 24 days of age, followed by a quick drop of the level at 44 days of age (Fig.

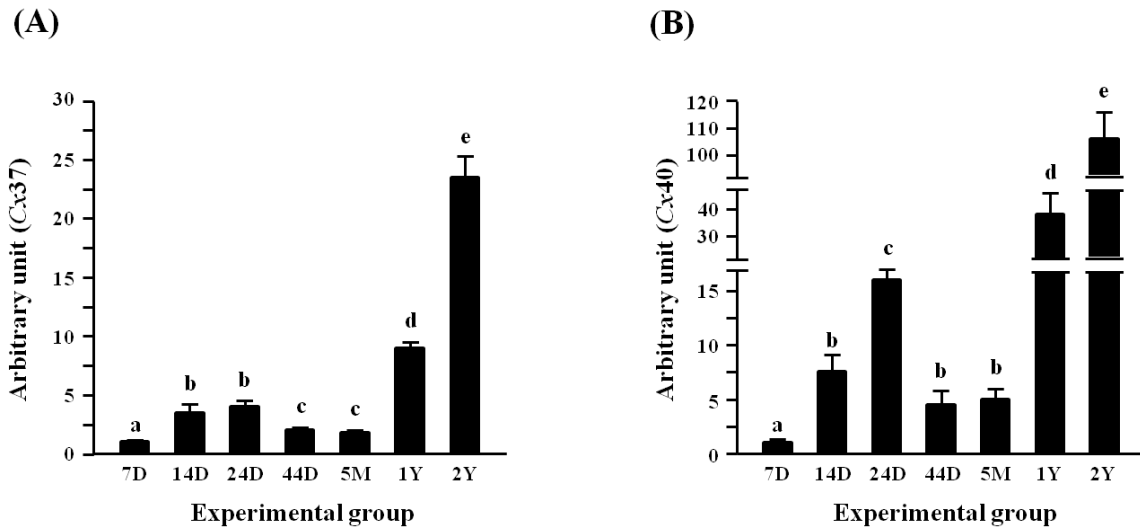


Fig. 4. Expressional patterns of Cx37 and Cx40 in the rat epididymal fat during postnatal period. The relative expressional levels of Cx37 (A) and Cx40 (B) are shown here. Different letters indicate statistical significances at $p < 0.05$. Cx, connexin; D, day; M, month; Y, year.

4B). Compared with the level of Cx40 transcript at 5 months of age, over 7-fold increase of Cx40 transcript level was found at 1 year of age (Fig. 4B). Expressional level of Cx40 in the rat epididymal fat at 2 years of age was 2 times higher than that at 1 year of age (Fig. 4B).

5. Expressional patterns of Cx43, Cx45, and Cx50 expression in the epididymal fat of rat during postnatal period

Expressional level of Cx43 in the rat epididymal fat

was not significantly changed until 24 days of age (Fig. 5A). There was the first significant increase of Cx43 transcript level at 44 days of age, followed by a drop of the level at 5 months of age (Fig. 5A). However, another significant surge of Cx43 mRNA abundance was detected at 1 year of postnatal age (Fig. 5A).

The transcript level of Cx45 was significantly increased at 24 days of age, and a further increase of Cx45 mRNA level was followed at 44 days of age (Fig. 5B). The level of Cx45 transcript was significantly decreased at 5

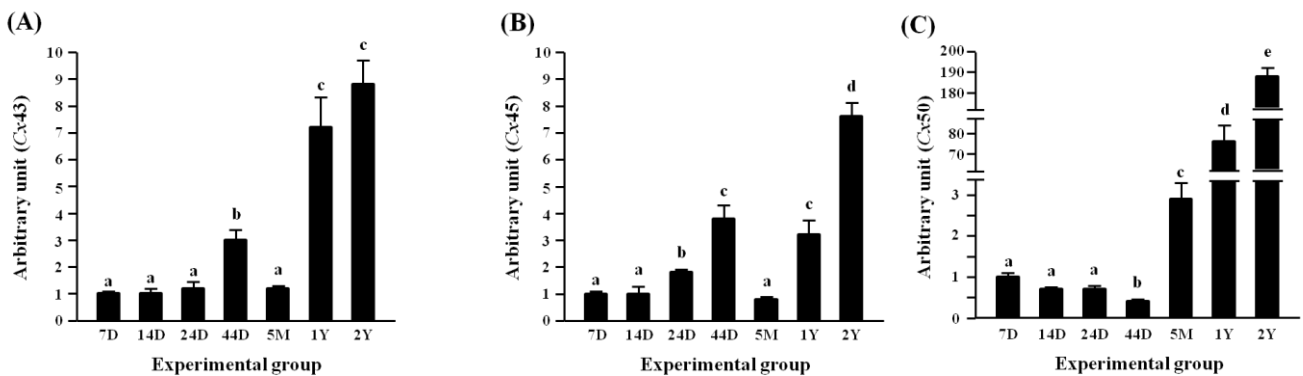


Fig. 5. Expressional patterns of Cx43, Cx45 and Cx50 in the rat epididymal fat during postnatal period. The relative expressional levels of Cx43 (A), Cx45 (B), and Cx50 (C) are shown here. Different letters indicate statistical significances at $p < 0.05$. Cx, connexin; D, day; M, month; Y, year.

months of age, but was again increased at 1 year of age (Fig. 5B). The highest expression level of *Cx45* was observed at 2 years of age (Fig. 5B).

The transcript level of *Cx50* was not changed until 24 days of age, followed by a significant decrease at 44 days of postnatal age (Fig. 5C). However, a surge of *Cx50* transcript level was found at 5 months of age (Fig. 5C). The level of *Cx50* transcript at 1 year of age was about 25 times higher than that at 5 months of age (Fig. 5C). The transcript level of *Cx50* at 2 years of postnatal age was over 1-fold higher than that at 1 year of age (Fig. 5C).

DISCUSSION

The present research shows the presence of several *Cx* isoforms in the epididymal fat and different expressional patterns of *Cx* isoforms during postnatal development. In general, compared with the transcript level of *Cx* isoforms at the neonatal age, the levels of all *Cx* isoforms at the old age, 1 year and 2 years of ages, are greatly increased in several folds, even hundreds folds. Even though expression of some *Cx* isoforms is steadily increased from the neonatal age to the adult age, expressional patterns of other *Cx* isoforms are fluctuated.

Unlike other tissues in the male reproductive tract, relatively little attention is paid to the function and necessity of the epididymal fat. There are some research reports showing a biological function of the epididymal fat on spermatogenesis. Surgical removal of the epididymal fat results in a decrease of seminiferous tubules in size and the cease of normal spermatogenesis in the testis (Chu et al., 2010). However, the removal of the epididymal fat dose not influence on testosterone production in the testis, indicating the existence of the epididymal fat-derived factor(s) involving in regulation of spermatogenesis in the testis (Chu et al., 2010). The epididymal fat expresses a number of adipose genes at a relatively high level, including leptin, adiponectin, and perilipin (Liu et al., 2011). In addition, it

has been observed that the aging could give an impact on global gene expression in the epididymal fat (Liu et al., 2011). Pinterova et al (2000) have found that the rat epididymal fat expresses all components of renin-angiotensin system, including renin, angiotensinogen, and angiotensin-converting enzyme, which is presumably involved in metabolic pathway or in the regulation of blood flow. These findings suggest that the epididymal fat would have a variety of physiological functions presumably involved in male reproduction.

The epididymal fat becomes greatly increased in weight with aging as a consequence of increases of fat cell size and number (Cleary et al., 1977). Not only the age but also other factors, such as sex and food intake, could influence on weight gain of the epididymal fat during postnatal development (Hubbard & Matthew, 1971). In addition, functional impairment of certain genes could induce abnormal hypertrophy of the epididymal fat. For example, leptin-deficient (*ob/ob*) or leptin receptor-deficient (*db/db*) mouse has extremely heavy epididymal fat with increases of fat cell size and number (Johnson & Hirsch, 1972). Interestingly, the epididymal fat of estrogen receptor α (*ER α*)-knockout (*α ERKO*) male mouse is heavier than that of wild-type mouse throughout entire postnatal period, and difference of the epididymal fat weight between *α ERKO* mouse and wild-type mouse becomes more clear as ages (Heine et al., 2000). Moreover, the exposure to exogenous compounds also gives an impact on weight of the epididymal fat. The oral administration of genistein, an estrogenic phytoestrogen, results in increases of epididymal fat weight and fat cell size (Penza et al., 2006). The presence and expression of *ER α* and androgen receptor in the epididymal fat have been reported (Dieudonne et al., 1995; Metz et al., 2016). Together, these observations suggest that the development of epididymal fat during postnatal period could be influenced by various intrinsic factor(s) as well as exogenous compounds.

A limited number of researches have demonstrated the presence and possible biological function of *Cx* isoforms in the epididymal fat (Zhu et al., 2016; Kim et al., 2017). Yanagiya et al (2007) have demonstrated that functional inhibition of gap junctional complex by 18 α -glycyrrhetic acid results in expressional reduction of several molecules involving in mitotic clonal expansion during adipogenesis. In addition, down-regulation of *Cx43* by small interfering RNA in 3T3-L1 cells causes a decrease of CCAAT/ enhancer-binding protein β (C/EBP β) expression (Yanagiya et al., 2007). The *Cx* isoforms function not only as a type of cell junctions but also as a hemichannel allowing direct exchange of small molecules between neighboring cells (Goodenough et al., 1996). Thus, it is speculated that expressional changes of *Cx* isoforms in the rat epididymal fat during postnatal development would relate with modification of adipogenic capacity and function of epididymal fat cells with aging. Expressional alteration of *Cx* isoforms is closely associated with the development of diseases, including atherosclerotic plaques (Kwak et al., 2002). This information suggests that increases of *Cx* isoform expression in epididymal fat at the old age would associate with onset and/or development of diverse age-related diseases and symptoms.

To our knowledge, the present research is the first report showing the presence and expressional patterns of multi-*Cx* isoforms in the rat epididymal fat during postnatal development. Based on current and other's findings, expression of *Cx* isoforms in the rat epididymal fat seems to be closely associated with the change of epididymal fat weight during postnatal development. However, there are some questions on expression of *Cx* isoforms in the epididymal fat to be answered. For examples, which cell type(s) in the epididymal fat tissue does express *Cx* isoforms? Which factor(s) causes differential expression of *Cx* isoforms during postnatal period and what are regulatory mechanisms on expression of *Cx* isoforms in the epididymal fat tissue? What is the biological function(s) of *Cx*

isoforms in the epididymal fat?

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