



Expression Analysis of *BMPR1B*, *BMP15*, *GDF9*, *Smad1*, *Smad5*, and *Smad9* in Rams with Different Fecundity

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ABSTRACT

To elucidate the tissue expression levels of *BMPR1B*, *BMP15*, *GDF9*, *Smad1*, *Smad5*, and *Smad9* genes in rams with different fecundity, quantitative real-time polymerase chain reaction was used to investigate the expression level of six genes in the brain, cerebellum, hypothalamus, pituitary, testis, epididymis, vas deferens, and adrenal gland in high fecundity (Small Tail Han sheep) and low fecundity (Sunite sheep) rams. The results were as follows: *BMPR1B*, *GDF9*, *Smad1*, *Smad5* and *Smad9* were expressed in all selected tissues, but *BMP15* was specifically expressed in the epididymis. Further study indicated that the expression of *BMPR1B* in the brain, hypothalamus, pituitary, epididymis, and adrenal gland was significantly higher in Sunite sheep than in Small Tail Han sheep ($p < 0.05$, $p < 0.01$); the expression of *BMP15* in the epididymis was significantly higher in Sunite sheep than in Small Tail Han sheep ($p < 0.01$); the expression of *GDF9* in the cerebellum and vas deferens was significantly higher in Small Tail Han sheep than in Sunite sheep ($p < 0.05$); the expression of *GDF9* in the adrenal gland was significantly higher in Sunite sheep than in Small Tail Han sheep ($p < 0.01$); the expression of *Smad1* in the brain and adrenal gland was significantly higher in Small Tail Han sheep than in Sunite sheep ($p < 0.05$); the expression of *Smad1* in vas deferens was significantly higher in Sunite sheep than in Small Tail Han sheep ($p < 0.01$); the expression of *Smad5* in the adrenal gland was significantly higher in Small Tail Han sheep than in Sunite sheep ($p < 0.05$); the expression of *Smad9* in the brain and epididymis was significantly higher in Sunite sheep than in Small Tail Han sheep ($p < 0.05$, $p < 0.01$); and the expression of *Smad9* in the cerebellum and hypothalamus was significantly higher in Small Tail Han sheep than in Sunite sheep ($p < 0.05$). This is the first study to systematically analyze the *BMPR1B*, *BMP15*, *GDF9*, *Smad1*, *Smad5*, and *Smad9* genes' tissue expression pattern in rams.

INTRODUCTION

Lambing is one of the most economically important traits of sheep, being closely related to the economic benefits of sheep breeding. The major genes related to the prolificacy of sheep have received much attention from researchers since the 1980s (Tang *et al.*, 2018). To date, three major candidate genes for the prolificacy of sheep have been found: bone morphogenetic protein receptor 1B

(*BMPR1B*), bone morphogenetic protein 15 (*BMP15*), and growth differentiation factor-9 (*GDF9*) (Pan *et al.*, 2018; Zhao *et al.*, 2019).

BMPR1B is the first major candidate gene found to be related to the prolificacy of sheep (Shokrollahi *et al.*, 2018) and possesses a mutation (A to G), known as FecB, which results in one amino acid substitution (Q to R) increasing the ovulation rate of Booroola ewes (Montgomery *et al.*, 1994). The FecB mutation has an additive effect on ovine ovulation number and litter size, so that one copy of the FecB mutation can increase the ovulation number by 1.3–1.6-fold and the litter size by 0.9–1.2-fold, and two copies by 2.73 and 1.1–1.7, respectively (El-Seedy *et al.*, 2017).

BMP15 also known as growth differentiation factor-

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9B (*GDF9B*), and *GDF9*, which both belong to the transforming growth factor- β (TGF- β) superfamily and are recognized as major candidate genes for the prolificacy of sheep—were found to regulate the growth and differentiation of follicles, the secretion of reproductive hormones, and the growth of germ cells (Belli *et al.*, 2018). To date, *FecX^I*, *FecX^H* (Galloway *et al.*, 2000), *FecX^G* (Hanrahan *et al.*, 2004), *FecX^L*, *FecX^B* (Bodin *et al.*, 2007), *FecX^R* (Monteagudo *et al.*, 2009), *FecX^{Gr}*, *FecX^O* (Julie *et al.*, 2013) and *FecX^{Bar}* (Lassoued *et al.*, 2017) mutations have been found on the sheep *BMP15* gene; they strongly affect the ovulation rate and prolificacy, similar to the effect of G1 (Wang *et al.*, 2018), G4 (Alam *et al.*, 2018), G6, *FecG^F* (G7) (Våge *et al.*, 2013), *FecG^H* (G8) (Shafieyan *et al.*, 2013), *FecG^E* (Silva *et al.*, 2011), *FecG^T* (*FecI*) (Braun *et al.*, 2003), *FecG^V* (Souza *et al.*, 2014), *FecG^T* (Nicol *et al.*, 2009), and *FecG^{SI}* (Mullen *et al.*, 2014) mutations on the sheep *GDF9* gene.

Many studies revealed that the prolificacy of sheep is closely related to bone morphogenetic proteins (BMPs), BMP receptors (BMPR), and Smads, a downstream signaling molecule of the TGF- β /Smad signaling pathway (Lin *et al.*, 2018). Members of the BMPs initiate signaling from the cell membrane by interacting with two distinct serine/threonine kinase receptors. Ligand binding induces the formation of a complex in which the type II receptor phosphorylates and activates the type I receptor. This protein then propagates the signal by phosphorylating the Smad proteins such as *Smad1*, *Smad5*, and *Smad9* (Song *et al.*, 2018). Phosphorylated *Smad1/5/9* can form a complex by interacting with *Smad4*, which can further activate or inhibit the expression of target genes (Rol *et al.*, 2018). Given its interaction with BMPs, *Smad1/5/9* might be related to the prolificacy of sheep.

Small Tail Han sheep (STH) and Sunite sheep (SNT) are two Chinese local sheep (*Ovis aries*) breeds with different estrous modes (year-round and seasonal, respectively). Both are known for their excellent meat production performance (Tang *et al.*, 2018). Significant differences between the two sheep breeds in fecundity have resulted in increasing interest in the expression pattern of major prolificacy genes in these sheep.

BMPR1B, *BMP15*, *GDF9*, *Smad1*, *Smad5*, and *Smad9* are important in prolificacy. Many studies on the expression of these six genes in the tissues of ewes have been reported; however, no research has yet been reported about these genes in rams. To explore the potential role of these six genes in rams, we analyzed the tissue expression profile and the mRNA expression levels in eight prolificacy-related tissues between high fecundity sheep breed (STH) and low fecundity sheep breed (SNT)

rams. Our study helps elucidate the genetic mechanism controlling high fecundity in rams.

MATERIALS AND METHODS

Selection of experimental sheep and sample collection

The 3 Small Tail Han rams and 3 Sunite rams were supplied by Yuncheng Breeding Sheep Farm (Yuncheng County, China) and Sheep and Goat Breeding Farm of Tianjin, Institute of Animal Sciences (Tianjin, China). All rams were healthy, approximately 2.5 years old, and were kept in a sheltered outdoor paddock and were provided with alfalfa hay and concentrate, with clear water available ad libitum. Eight tissues (brain, cerebellum, hypothalamus, pituitary, testis, epididymis, vas deferens, and adrenal gland) were collected from each animal. All tissues were snap-frozen in liquid nitrogen and then stored at -80°C to be used for RNA extraction.

All experimental procedures mentioned in the present study were approved by the Science Research Department (in charge of animal welfare issue) of the Institute of Animal Sciences, Chinese Academy of Agricultural Sciences (IAS-CAAS) (Beijing, China). Ethical approval was provided by the animal ethics committee of IAS-CAAS (No. IASCAAS-AE-03, December 12, 2016).

Total RNA extraction and cDNA synthesis

Tissue RNA was extracted from the 8 tissues using a total RNA extraction kit for animal tissue (Tiangen, Beijing, China) and Trizol (Invitrogen Inc., Carlsbad, CA, USA) was used to dissolve the tissues (each tissue smashed, mixed, and 50–100 mg used for RNA extraction). The quantity and quality of total RNA were monitored using 1.5% agarose gel electrophoresis and ultraviolet spectrophotometry (UV-1201, Shimadzu, Kyoto, Japan), respectively. Then, the RNA was stored at -80°C until use.

The first strand of cDNA was prepared following the instructions of the PrimeScript™ RT Reagent Kit (TaKaRa Bio Inc., Dalian, China). The reaction program was as follows: 37°C for 15 min, followed by 85°C for 5 sec, with a total volume of 20 μL that contained PrimeScript RT Enzyme 1.0 μL , Random 6 mers 1.0 μL , 5 \times PrimeScript Buffer (for Real Time) 4.0 μL , total RNA 1.0 μL and RNase-free ddH₂O 13 μL . Prior to storage at -80°C , the standard working concentration of cDNA is 200ng/ μl , the cDNA quality was evaluated by housekeeping gene (*RPL-19*) amplification, and then the reverse products were stored at -20°C until use.

RNA extraction and cDNA synthesis

Total RNA samples were analyzed using 1.5% agarose gel electrophoresis ($U = 150$ V; $I = 240$ mA).

Three bands were detected (28S, 18S, and 5S)—the 28S band was brighter than the 18S band, and the 5S band was unclear. The OD260 nm/OD280 nm ratios (1.8–2.0) of the RNA samples were all 1.9 to 2.0, which showed that the extracted total RNA was of sufficient purity with no contamination or degradation. Therefore, these tissue RNAs were appropriate for use in the follow-up experiment.

Primer design

A total of 7 primers were designed using the Primer Premier (version 5.0, PREMIER Biosoft Co., Palo Alto, CA, USA) software to amplify different fragments of ovine *BMPR1B*, *BMP15*, *GDF9*, *Smad1*, *Smad5*, *Smad9*, and *RPL-19* genes based on their assembled sequences in GenBank. All primers were synthesized by Beijing Tianyi Biotechnology Co., Ltd. (Beijing, China). The housekeeping gene (*RPL-19*, accession number: XM_012186026.1) was used as an internal control to normalize the threshold cycle (Ct) values. The primers are detailed in Table I.

qPCR

Real-time polymerase chain reaction (PCR) amplification was performed in 20 μ L of reaction mixture that contained 10 μ L SYBR Premix EX Taq II (TaKaRa Bio Inc., Dalian, China), 0.4 μ L of each forward and reverse primer, 6.4 μ L RNase-Free ddH₂O, and 2 μ L cDNA. PCR amplification was performed in triplicate wells using the following conditions: initial denaturation at 95°C for 5 min, followed by 40 cycles of 95°C for 10 sec, and 60°C for 30 sec. The dissociation curve was analyzed after amplification. A melting temperature (*Tm*) peak at 85±0.8°C on the dissociation curve was used to determine the specificity of PCR amplification.

Table I.- Primers of studied genes.

Gene Name	Primer sequence (5'→3')	Length (bp)	Tm (°C)	Accession No.
BMPR1B	F: 5'-TGACGGACCTATAACACCACA-3' R: 5'-GTACCGAGGTCTGGCTCTT-3'	121	60	NM_001142888.2
BMP15	F: 5'-TGGTGGCAAAAGCTCTGGA-3' R: 5'-GCCATGCCACCAGAACTCAA-3'	106	60	NM_001114767.1
GDF9	F: 5'-AACAGACGCCACCTCTACAA-3' R: 5'-CACGATCCAGGTTAACAGCA-3'	124	60	NM_001009431.1
Smad1	F: 5'-TGGTTCCAAGACACAGCGAATA-3' R: 5'-GGTGTATCTGCTGGCATCTGAA-3'	252	60	XM_015101506.1
Smad5	F: 5'-GCACAGCCTCTGGTCA-3' R: 5'-GGGTAGGGACTATTGGAG-3'	132	60	XM_012115987.1
Smad9	F: 5'-CCAGCACTCAGATTTCGGC-3' R: 5'-GCACTCGGCATAGACCTCTC-3'	147	60	XM_015098108.1
RPL-19	F: 5'-ATCGCCAATGCCAACTC-3' R: 5'-CCTTCGCTTACCTATAACC-3'	154	60	XM_012186026.1

Data

The $2^{-\Delta\Delta Ct}$ method (Guo *et al.* 2018) was used to process the real-time PCR results. Statistical analyses were carried out using SPSS 19.0 software (IBM, Armonk, NY, USA). The levels of gene expression were analyzed for significant differences with one-way analysis of variance (ANOVA), followed by the Fisher's least significant difference (LSD) test as a multiple comparison test. All experimental data are shown as mean±SEM. A probability of $p \leq 0.05$ was considered statistically significant, and a probability of $p \leq 0.01$ was considered to be extremely statistically significant.

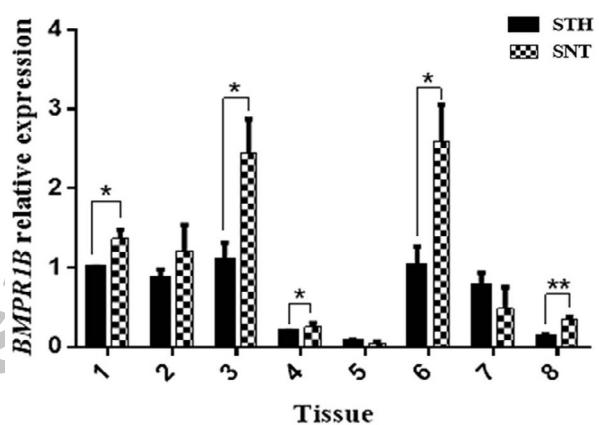


Fig. 1. Comparison of the expression of *BMPR1B* in eight tissues (Tissues 1–8: brain, cerebellum, hypothalamus, pituitary, testis, epididymis, vas deferens, and adrenal gland, respectively) between Small Tail Han sheep (STH) and Sunnite sheep (SNT). Means with different superscripts are significantly different. The significant results with a *p*-values lower than 0.01 and 0.05 are given two asterisks (**) and one asterisk (*), respectively.

RESULTS

Expression levels of BMPR1B, BMP15, GDF9, Smad1, Smad5, and Smad9

The expression levels of *BMPR1B*, *BMP15*, *GDF9*, *Smad1*, *Smad5*, and *Smad9* in eight tissues (brain, cerebellum, hypothalamus, pituitary, testis, epididymis, vas deferens, and adrenal gland) in both high fecundity breed Small Tail Han sheep and low fecundity breed Sunit sheep were measured by qPCR in this study.

As shown in Figure 1, *BMPR1B* is expressed in all tissues with the highest level in the epididymis, followed by the hypothalamus, brain, and cerebellum. The expression of *BMPR1B* in the brain, hypothalamus, pituitary, epididymis, and adrenal gland is significantly higher in SNT than in STH ($p < 0.01$, $p < 0.05$).

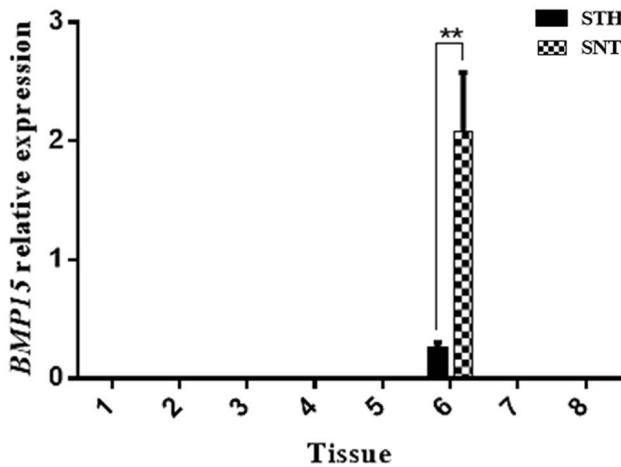


Fig. 2. Comparison of the expression of *BMP15* in eight tissues (Tissues 1–8: brain, cerebellum, hypothalamus, pituitary, testis, epididymis, vas deferens, and adrenal gland, respectively) between STH and SNT. Means with different superscripts are significantly different. The significant results with a p -values lower than 0.01 is given two asterisks (**).

As shown in Figure 2, *BMP15* is specifically expressed in the epididymis and the expression of *BMP15* in the epididymis is significantly higher in SNT than in STH ($p < 0.01$).

As shown in Figure 3, *GDF9* is expressed in all tissues with the highest level in the brain, followed by the testis, cerebellum, and hypothalamus. The expression of *GDF9* in the cerebellum and vas deferens is significantly higher in STH than in SNT ($p < 0.05$). The expression of *GDF9* in the adrenal gland is significantly higher in SNT than in STH ($p < 0.01$).

As shown in Figure 4, *Smad1* is expressed in all tissues

with the highest level in the epididymis, followed by the cerebellum, brain, and hypothalamus. The expression of *Smad1* in the brain and adrenal gland is significantly higher in STH than in SNT ($p < 0.05$). The expression of *Smad1* in the vas deferens is significantly higher in SNT than in STH ($p < 0.01$).

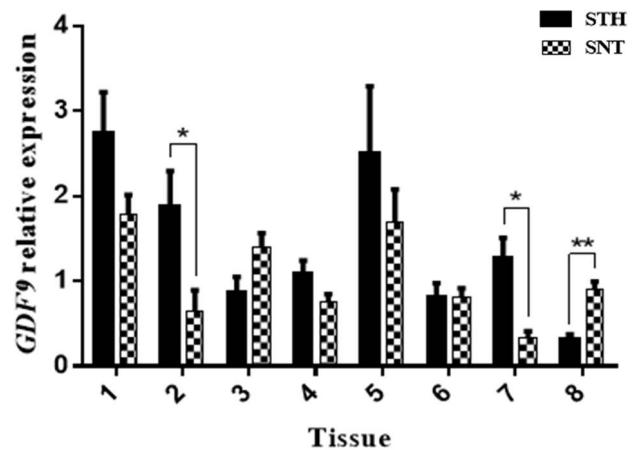


Fig. 3. Comparison of the expression of *GDF9* in eight tissues (Tissues 1–8: brain, cerebellum, hypothalamus, pituitary, testis, epididymis, vas deferens, and adrenal gland, respectively) between STH and SNT. Means with different superscripts are significantly different. The significant results with a p -values lower than 0.01 and 0.05 are given two asterisks (**) and one asterisk (*), respectively.

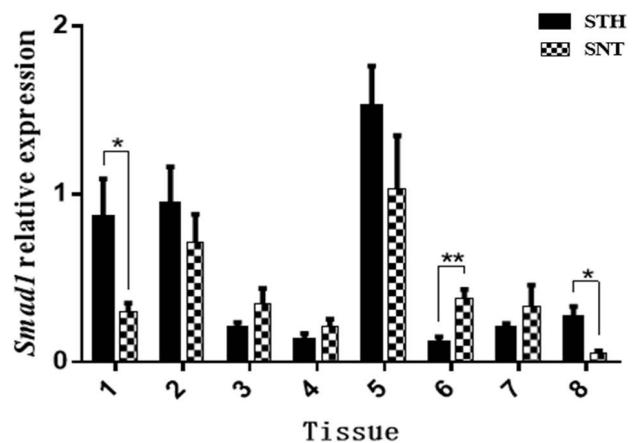


Fig. 4. Comparison of the expression of *Smad1* in eight tissues (Tissues 1–8: brain, cerebellum, hypothalamus, pituitary, testis, epididymis, vas deferens, and adrenal gland, respectively) between STH and SNT. Means with different superscripts are significantly different. The significant results with a p -values lower than 0.01 and 0.05 are given two asterisks (**) and one asterisk (*), respectively.

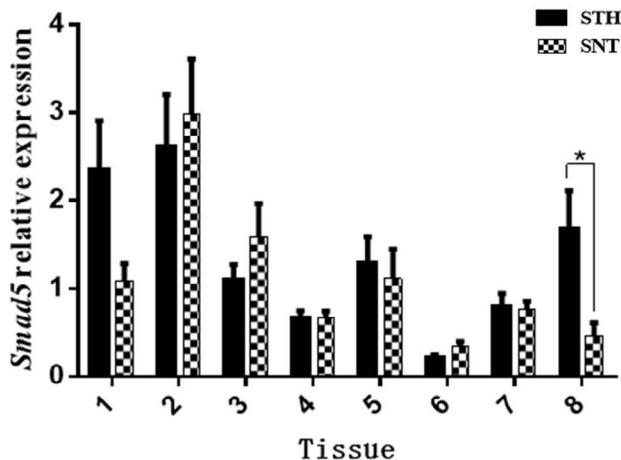


Fig. 5. Comparison of the expression of *Smad5* in eight tissues (Tissues 1–8: brain, cerebellum, hypothalamus, pituitary, testis, epididymis, vas deferens, and adrenal gland, respectively) between STH and SNT. Means with different superscripts are significantly different. The significant results with a *p*-values lower than 0.05 is given one asterisk (*).

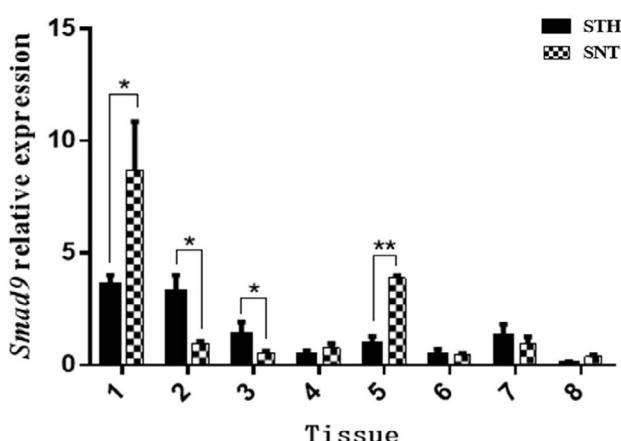


Fig. 6. Comparison of the expression of *Smad9* in eight tissues (Tissues 1–8: brain, cerebellum, hypothalamus, pituitary, testis, epididymis, vas deferens, and adrenal gland, respectively) between STH and SNT. Means with different superscripts are significantly different. The significant results with a *p*-values lower than 0.01 and 0.05 are given two asterisks (**) and one asterisk (*), respectively.

As shown in Figure 5, *Smad5* is expressed in all tissues with the highest level in cerebellum, followed by the brain, hypothalamus, and adrenal gland. The expression of *Smad5* in the adrenal gland is significantly higher in STH than in SNT (*p* < 0.05).

As shown in Figure 6, *Smad9* is expressed in all

tissues with the highest level in the brain, followed by the cerebellum, epididymis, and hypothalamus. The expression of *Smad9* in the brain and epididymis is significantly higher in SNT than in STH (*p* < 0.05, *p* < 0.01, respectively). The expression of *Smad9* in the cerebellum and hypothalamus is significantly higher in STH than in SNT (*p* < 0.05).

DISCUSSION

BMPR1B

BMPR1B (*FecB* gene) is one of the major fecundity genes in female reproduction; however, not much is known about the reproductive role of the *BMPR1B* gene in male reproduction. Previous reports have found that *BMPR1B* belongs to the type I receptors of BMPs (Aquino *et al.*, 2017; Kaivo-oja *et al.*, 2006), which figures prominently in the directional migration and proliferation of primordial germ cells (PGCs) (Dudley *et al.*, 2007; Okamura *et al.*, 2005) and precursor cells of sperm (Hammoud *et al.*, 2014; Larriba *et al.*, 2018). Therefore, the *BMPR1B* gene has a certain impact on male reproduction.

Studies have found that *BMPR1B* is widely expressed in the ovary, liver, hypothalamus, pituitary, uterus, heart, and muscle of mammals (Valdecantos *et al.*, 2017; Goyal *et al.*, 2017; Foroughinia *et al.*, 2017). In ewes, *BMPR1B* is highly expressed in the reproductive tissues and moderately expressed in the brain, skeletal muscle, and kidney (Ciller *et al.*, 2016; Tang *et al.*, 2018; Wilson *et al.*, 2001). In this research, *BMPR1B* was found to be expressed in all selected tissues and highly expressed in the brain, cerebellum, hypothalamus, and epididymis, which indicated that *BMPR1B* may have a role in both ewe and ram reproduction.

The expression of *BMPR1B* in the brain, hypothalamus, pituitary, epididymis, and adrenal gland is significantly higher in SNT than in STH. This observation was different from a previous study comparing prolific and non-prolific ewes (Xu *et al.*, 2010), in which ewes with high fecundity were reported to have a higher expression of *BMPR1B* in the reproductive tissues, which implies that ram may have a different regulation mechanism in reproduction when compared to ewe. Considering the function of *BMPR1B* in the proliferation of PGCs, it seems plausible that *BMPR1B* may have a certain inhibitory effect on ram reproduction. Of course, further studies should be performed deeply to investigate the relationship between *BMPR1B* and ram reproduction.

BMP15

BMP15 is an important regulator of male germ stem cell (mGSC) proliferation and differentiation (Liu *et al.*, 2018). Hu *et al.* (2017) reported that over-expression of

BMP15 in goat mGSCs leads to the increased expression level of c-Kit, a gene that promotes spermatogonial differentiation and the proliferation of mGSCs. Thus, *BMP15* is an important candidate gene in male fertility.

In 1998, Dube *et al.* (1998) explored the expression of *BMP15* in several tissues, including ovary and testis in mice, and found that *BMP15* is specifically expressed in the ovary. The expression of *BMP15* in goats (Silva *et al.*, 2005) and pigs (Li *et al.*, 2008) is similar to that in mice. In contrast, Aaltonen *et al.* (1999) reported the expression of *BMP15* in the testis and ovary in humans. Similarly, Pennetier *et al.* (2004) reported the expression of *BMP15* in the testis and ovary in cattle. This study suggested that *BMP15* is specifically expressed in the epididymis in rams. One potential explanation is that the differences in the genetic models led to these results.

It is known that *BMP15* exerts its biological effects by initially interacting with a type II receptors of BMPs, which results in the activation and phosphorylation of *BMPR1B* (Moore *et al.*, 2003). We compared the expression level of *BMP15* and *BMPR1B*, the expression of *BMP15* and *BMPR1B* in the epididymis in SNT was found to be significantly higher than in STH which implies that the expression level of *BMP15* and *BMPR1B* may be negatively correlated with the fecundity of rams.

GDF9

For a long time, *GDF9* was considered to be specifically expressed in the ovaries of animals, until Fitzpatrick *et al.* (1998) reported the expression of *GDF9* in non-ovarian tissues including the testis, brain, pituitary and bone marrow. Earlier, the *GDF9* expression was detected in the testis in rats, mice, humans, cattle (Tang *et al.*, 2017), alpacas (Guo *et al.*, 2013) and cats (Zhao *et al.*, 2011). To our knowledge, no research on the expression of *GDF9* in rams has ever been reported. In the present study, *GDF9* was detected in all 8 tissues in rams, which implies that it plays a role in promoting the differentiation of many tissues. The highest expression of *GDF9* in the epididymis further confirmed that *GDF9* is associated with the epididymal function.

Because numerous studies revealed that *GDF9* has promoting effects on genetic and cellular signaling levels and the mitosis of germ cells (Tang *et al.*, 2013; He *et al.*, 2012), we compared the expression level of *GDF9* in the testis, epididymis, and vas deferens between two sheep breeds. We found no significant difference between the expression level of *GDF9* in the testis and epididymis of SNT and STH, but the expression level of *GDF9* in the vas deferens of STH is significantly higher than in SNT. Our findings are in agreement with Tang *et al.* (2017) who

found exogenous *GDF9* significantly promoted Sertoli cells (SCs) proliferation and inhibited the apoptosis of SCs which suggested *GDF9* to have a supporting role for the maintenance of spermatogenesis. Therefore, we concluded that the prolificacy of ram might be due to the high expression level of the *GDF9* gene.

Smad

Smad1/5/9 is widely expressed in mammals, especially in the brain and hypothalamus–pituitary–gonadal (HPG) axis (Wang *et al.*, 2018a, b; Ohyama *et al.*, 2015). The present study found that *Smad1/5/9* is detectable in eight tissues of rams; meanwhile, the expression level of *Smad1/5/9* in the brain, cerebellum, and HPG axis is higher than in other tissues, which is consistent with our expectation.

BMP type I receptors are transphosphorylated by type II receptors, resulting in cascades of *Smad* signaling (Aquino *et al.*, 2017). Shi *et al.* (2016) found that the deletion of *BMPR1B* leads to an increased phosphorylation level and decreased expression level of *Smad1/5/9* in male mice. To explore the expression pattern of *BMPR1B* in rams, we compared the expression level of *BMPR1B* and *Smad1/5/9*. We surprisingly found that the expression level of *BMPR1B* in the brain, epididymis, and adrenal gland is significantly higher in SNT than in STH; meanwhile, the expression level of *Smad1/5* in the brain and adrenal gland was higher in STH than in SNT. The results show that *BMPR1B* signaling may be involved in some of those mechanisms in the brain and adrenal gland of rams; however, the expression level of *Smad1* in the epididymis is significantly higher in SNT than in STH. We speculate that *BMPR1B* may have a certain inhibitory effect on the spermatogenesis of rams; however, the degree of the function of *BMPR1B* in rams remains to be investigated.

Studies have demonstrated that the BMP/*Smad* signaling pathway may be associated with the spermatogenesis process (Itman *et al.*, 2010). Research on mice (Mendis *et al.*, 2011) and rats (Chan *et al.*, 2017) may provide an insight into the synergistic effect of *BMP15* and *Smad1/5/9* in male animals: the expression level of *Smad1/5/9* is positively regulated by BMP signaling (Katakawa *et al.*, 2016). Additionally, we compared the expression level of *BMP15* and *Smad1/5/9*. The expression of *BMP15* and *Smad1* in the epididymis of SNT is significantly higher than in STH, but there is no significant difference between the expression of *Smad5* and *Smad9* in the epididymis of two sheep breeds. We provide evidence that the BMP/*Smad* signaling pathway may be associated with the spermatogenesis process in rams to some degree. Further research is necessary to draw conclusions.

CONCLUSIONS

In conclusion, we found that *BMPR1B*, *GDF9*, *Smad1*, *Smad5*, and *Smad9* are expressed in all selected tissues and are highly expressed in the epididymis, whereas *BMP15* is specifically expressed in the epididymis, which indicates that they may play important roles in the ovine epididymis and promote spermatogenesis. Our findings of ovine *BMPR1B*, *BMP15*, *GDF9*, *Smad1*, *Smad5*, and *Smad9* will help to further understand their expression and function, and may contribute to exploring their role in the ram reproduction system. This is the first study of the six genes tissue expression pattern in rams.

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Statement of conflicts of interest

All authors declare no conflicts of interest.

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