

Molecular Cloning, Sequence Identification and Tissue Expression Profile of Three Novel Sheep (*Ovis aries*) Genes-*RAB1A*, *RAB4A* and *RAB5A*

¹Jiang Caode, ²Liu Yonggang and ³Xia Xueshan

¹School of Life Science, Southwest University, 400715 Chongqing, China

²College of Animal Science and Technology, Yunnan Agricultural University,
650201 Kunming, China

³Faculty of Life Science and Technology, Kunming University of Science and Technology,
650224 Kunming, China

Abstract: The complete coding sequences of three sheep genes-*RAB1A*, *RAB4A* and *RAB5A* were amplified using the Reverse Transcriptase Polymerase Chain Reaction (RT-PCR). Sequence analysis revealed that the sheep *RAB1A* gene encodes a protein of 204 amino acids that shares high homology with the RAB1A, member RAS oncogene family (RAB1A) proteins of nine species-human (98%), mouse (98%), rat (98%), pig (98%), zebrafish (98%), Western clawed frog (97%), Atlantic salmon (96%), disc abalone (92%) and great pond snail (87%). The sheep *RAB4A* gene encodes a protein of 218 amino acids that shares high homology with the RAB4A, member RAS oncogene family (RAB4A) proteins of eight species-cattle (100%), dog (99%), human (99%), rhesus monkey (99%), horse (99%), chicken (99%), rat (98%) and mouse (98%). The sheep *RAB5A* gene encodes a protein of 215 amino acids that shares high homology with the RAB5A, member RAS oncogene family (RAB5A) proteins of fifteen species-cattle (98%), rabbit (98%), dog (98%), horse (98%), pig (98%), sumatran orangutan (98%), human (97%), mouse (97%), rat (97%), chimpanzee (97%), chicken (95%), rhesus monkey (93%), African clawed frog (93%), Western clawed frog (93%) and zebrafish (92%). Finally, these three novel sheep genes were assigned to GeneIDs: 100302065, 100302066 and 100302084. Phylogenetic analysis indicated that the sheep *RAB1A* gene has a closer genetic relationship with the *RAB1A* genes of human, mouse and rat. The sheep *RAB4A* and *RAB5A* genes both have closer genetic relationships with the *RAB4A* and *RAB5A* genes of cattle. Tissue expression profile analysis was also carried out and results demonstrated that sheep *RAB1A*, *RAB4A* and *RAB5A* genes were all generally but differentially expressed in detected tissues.

Key words: Sheep, RAB1A, RAB4A, RAB5A, tissue expression, China

INTRODUCTION

RAB1A, member RAS oncogene family (RAB1A) is a member of Rab1 subfamily. Rab1 is found in eukaryote and is an important regulatory factor for the transport of vesicles from the ER to the Golgi apparatus. Latest researches suggested a novel function for Rab1a in the regulation of cell migration through controlling integrin beta1 recycling and localization to lipid rafts via a specific downstream effector pathway. Researches also revealed that the Rab1a plays a crucial role in mammalian autophagy (Huang *et al.*, 2011; Wang *et al.*, 2010; Diao *et al.*, 2008).

RAB4A, member RAS oncogene family (RAB4A) is a member of Rab4 subfamily. Rab4 has been implicated in numerous functions within the cell. Experimental data revealed that overexpression of Rab4 regulates

angiotensin II type I receptor phosphorylation and sensitization. Rab4A had also been identified to be a critical effector of VEGFR1 during branching morphogenesis of the vasculature (Esseltine *et al.*, 2011; Schonhoff *et al.*, 2009; Kachhap *et al.*, 2007).

RAB5A, member RAS oncogene family (RAB5A) is a member of Rab5-related subfamily. This subfamily includes Rab5 and Rab22 of mammals, Ypt51/Ypt52/Ypt53 of yeast and RabF of plants. Recent researches showed that selective upregulation of Rab5 level is associated with mild cognitive impairment, Alzheimer's disease and sporadic motor neuron disease. Rab5a can promote proliferation of ovarian cancer cells (Ginsberg *et al.*, 2010; Zhao *et al.*, 2010; Matej *et al.*, 2010).

As mentioned before, *RAB1A*, *RAB4A* and *RAB5A* genes are three genes which have important functions. Until today, *RAB1A*, *RAB4A* and *RAB5A* genes had

been reported in human and other animals but the sheep *RAB1A*, *RAB4A* and *RAB5A* genes have not been reported yet.

In present experiment, the researchers will isolate the coding sequences of sheep *RAB1A*, *RAB4A* and *RAB5A* genes based on the coding sequence information of *RAB1A*, *RAB4A* and *RAB5A* genes from human or other mammals and their highly homologous sheep ESTs sequence information, subsequently perform some necessary sequence analysis and tissue expression profile analysis for these genes. These will establish the primary foundation of understanding these three sheep genes.

MATERIALS AND METHODS

Animals and sample preparation: Five adult Yunnan local sheep were slaughtered. Spleen, skin, lung, fat, muscle, heart, liver, kidney and ovary samples were collected, frozen in liquid nitrogen and then stored at -80°C. The total RNA was extracted using the total RNA extraction kit (Gibco, USA). First-strand cDNA synthesis was performed as that described by Liu *et al.* (2004). These first-strand cDNA samples were used to perform RT-PCR for the isolation of sheep *RAB1A*, *RAB4A* and *RAB5A* genes and for the tissue expression profile analysis.

Isolation of the sheep *RAB1A*, *RAB4A* and *RAB5A* genes: The primers for sheep *RAB1A* gene isolation were designed based on the coding sequence information of human *RAB1A* gene and its highly homologous sheep EST sequences: EE751788 and EE747095. Similarly, the primers for sheep *RAB4A* gene isolation were designed based on the coding sequence information from human *RAB4A* gene and its highly homologous sheep EST sequence: EE794089. The primers for sheep *RAB5A* gene isolation were designed based on the coding sequence information from human and mouse *RAB5A* genes and their highly homologous sheep EST sequences: EE806405 and EE791858. These primer sequences and their annealing temperature for RT-PCR reaction were shown in Table 1. The RT-PCR was performed to isolate these three sheep genes using the pooled cDNAs from different tissues above. The 25 µL reaction system was 2.0 µL cDNA, 2.5 µL 2 mM mixed dNTPs, 2.5 µL 10×Taq DNA polymerase buffer, 2.5 µL 25 mM MgCl₂, 2.0 µL 10 µM forward primer, 2.0 µL 10 µM reverse primer, 2.0 units of Taq DNA polymerase (1 U/1 µL) and 9.5 µL sterile water. The PCR program initially started with a 94°C denaturation for 4 min followed by 35 cycles of 94°C/50 sec, Ta°C/50 sec, 72°C/50 sec then 72°C extension for 10 min, finally 4°C to terminate the reaction.

Table 1: Primers for sheep *RAB1A*, *RAB4A*, *RAB5A* and *Beta-actin* genes and their annealing temperatures

Genes	Primer sequence	Ta/°C
<i>RAB1A</i>	Forward 5'-ATGTCCAGCATGAATCCC-3'	56
	Reverse:5'-TTAGCAGCACCTCCACTT-3'	
<i>RAB4A</i>	Forward :5'-ATGTCGCAGACGCCATGT-3	62
	Reverse: 5'-TCAGCAGCCGCACTCCTG-3	
<i>RAB5A</i>	Forward5'-ATGGCTAATCGAGGAGCAG-3'	57
	Reverse:5'-TCAGTTGCTGCAGCACTG-3'	
<i>Beta-actin</i>	Forward: 5'-CTTGATGTACGGACGATTT -3'	56
	Reverse: 5'-CACGGCATTGTCACCAACT-3'	

These PCR products for sheep *RAB1A*, *RAB4A* and *RAB5A* genes were then cloned into PMD18-T vector and sequenced bidirectionally with the commercial fluorometric method. At least five independent clones were sequenced for every gene.

RT-PCR for tissue expression profile analysis: RT-PCR for tissue expression profile analysis was performed as previously described elsewhere (Liu and Gao, 2009; Yonggang and Shizheng, 2009; Liu, 2009). We selected the housekeeping gene *beta-actin* (Accession No.: NM_001009784) as a positive control. The primers of sheep *RAB1A*, *RAB4A* and *RAB5A* genes which were used to perform the RT-PCR for tissue expression profile analysis were same as the primers for isolation RT-PCR above. The PCR reactions were optimized for a number of cycles to ensure product intensity within the linear phase of amplification. The 25 µL reaction system was 1 µL cDNA (100 ng µL⁻¹), 5 pmoles each oligonucleotide primer, 2.5 µL 2 mmolL⁻¹ mixed dNTPs, 2.5 µL 10×Taq DNA polymerase buffer, 2.5 µL 25 mmol L⁻¹ MgCl₂, 1.0 unit of Taq DNA polymerase and finally add sterile water to volume 25 µL. The PCR program initially started with a 94°C denaturation for 4 min followed by 25 cycles of 94°C/50 sec, Ta°C/50 sec, 72°C/50 sec then 72°C extension for 10 min, finally 4°C to terminate the reaction.

Sequence analysis: The cDNA sequence prediction was conducted using GenScan software (<http://genes.mit.edu/GENSCAN.html>). The protein prediction and analysis were performed using BLAST tool at the National Center for Biotechnology Information (NCBI) server (<http://www.ncbi.nlm.nih.gov/BLAST>) and the ClustalW software (<http://www.ebi.ac.uk/clustalw>).

RESULTS AND DISCUSSION

RT-PCR results for sheep *RAB1A*, *RAB4A* and *RAB5A* genes: Through RT-PCR with pooled tissue cDNAs for sheep *RAB1A*, *RAB4A* and *RAB5A* genes, the resulting PCR products were 615, 657 and 648 bp (Fig. 1).


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Dog      MSQTAMSETYDFLKFVIGNAGTGKSCLLHQFIEKKFKDDSNHTIGVEFGSKIINVGGK
Chicken  MSQAMSETYDFLKFVIGNAGTGKSCLLHQFIEKKFKDDSNHTIGVEFGSKIINVGGK
Sheep_Cattle MSQTAMSETYDFLKFVIGNAGTGKSCLLHQFIEKKFKDDSNHTIGVEFGSKIINVGGK
Human_Rhesus monkey MSQTAMSETYDFLKFVIGNAGTGKSCLLHQFIEKKFKDDSNHTIGVEFGSKIINVGGK
Horse    MSQTAMSETYDFLKFVIGNAGTGKSCLLHQFIEKKFKDDSNHTIGVEFGSKIINVGGK
Rat      MAQTAMSETYDFLKFVIGNAGTGKSCLLHQFIEKKFKDDSNHTIGVEFGSKIINVGGK
Mouse    MAQTAMSETYDFLKFVIGNAGTGKSCLLHQFIEKKFKDDSNHTIGVEFGSKIINVGGK
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Dog      YVKLQIWDTAGQERFRSVTRSYYRGAAGALLVVDITSRETYNALTINWLTARDMLASQNI
Chicken  YVKLQIWDTAGQERFRSVTRSYYRGAAGALLVVDITSRETYNALTINWLTARDMLASQNI
Sheep_Cattle YVKLQIWDTAGQERFRSVTRSYYRGAAGALLVVDITSRETYNALTINWLTARDMLASQNI
Human_Rhesus monkey YVKLQIWDTAGQERFRSVTRSYYRGAAGALLVVDITSRETYNALTINWLTARDMLASQNI
Horse    YVKLQIWDTAGQERFRSVTRSYYRGAAGALLVVDITSRETYNALTINWLTARDMLASQNI
Rat      YVKLQIWDTAGQERFRSVTRSYYRGAAGALLVVDITSRETYNALTINWLTARDMLASQNI
Mouse    YVKLQIWDTAGQERFRSVTRSYYRGAAGALLVVDITSRETYNALTINWLTARDMLASQNI
*****

Dog      IILCGNKKDLADREVTFLASRFAQENELMFLETSAITGENVEEAFVQCARKILNKIES
Chicken  IILCGNKKDLADREVTFLASRFAQENELMFLETSAITGENVEEAFVQCARKILNKIES
Sheep_Cattle IILCGNKKDLADREVTFLASRFAQENELMFLETSAITGENVEEAFVQCARKILNKIES
Human_Rhesus monkey IILCGNKKDLADREVTFLASRFAQENELMFLETSAITGENVEEAFVQCARKILNKIES
Horse    IILCGNKKDLADREVTFLASRFAQENELMFLETSAITGENVEEAFVQCARKILNKIES
Rat      IILCGNKKDLADREVTFLASRFAQENELMFLETSAITGENVEEAFVQCARKILNKIES
Mouse    IILCGNKKDLADREVTFLASRFAQENELMFLETSAITGENVEEAFVQCARKILNKIES
:*****:*****:*****

Dog      GELDPERMGSGIQYGDAAALRQLRSPPRAQAQPSAQECGC
Chicken  GELDPERMGSGIQYGDAAALRQLRSPPRAQAQPSAQECGC
Sheep_Cattle GELDPERMGSGIQYGDAAALRQLRSPPRAQAQPSAQECGC
Human_Rhesus monkey GELDPERMGSGIQYGDAAALRQLRSPPRAQAQPSAQECGC
Horse    GELDPERMGSGIQYGDAAALRQLRSPPRTQAPNAQECGC
Rat      GELDPERMGSGIQYGDAAALRQLRSPPRTQAPSAQECGC
Mouse    GELDPERMGSGIQYGDAAALRQLRSPPRTQAPSAQECGC
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Fig. 3: The alignment of the protein encoded by sheep *RAB4A* gene and eight other kinds of *RAB4A* proteins

NP_037151; 98%), mouse (accession number: NP_033029; 98%) (Fig. 3). The sheep *RAB5A* protein has high homology with the *RAB5A*, member RAS oncogene family (*RAB5A*) proteins of fifteen species-cattle (Accession No.: NP_001069654; 98%), rabbit (Accession No.: XP_002716250; 98%), dog (Accession No.: NP_001003317; 98%), pig (Accession No.: NP_001116652; 98%), horse (Accession No.: XP_001495368; 98%), sumatran orangutan (Accession No.: XP_002814060; 98%), human (Accession No.: NP_004153; 97%), mouse (Accession No.: NP_080163; 97%), rat (Accession No.: NP_073183; 97%), chimpanzee (Accession No.: XP_516319; 97%), chicken (Accession No.: NP_001006363; 95%), rhesus monkey (Accession No.: XP_001086669; 93%), African clawed frog (Accession No.: NP_001080535; 93%), Western clawed frog (Accession No.: NP_001008068; 93%) and zebrafish (Accession No.: NP_958893; 92%) (Fig. 4).

Based on the results of the alignment of *RAB1A*, *RAB4A* and *RAB5A* proteins, three phylogenetic trees were constructed using the Dendrogram procedure of ClustalW software (<http://align.genome.jp/>) as shown in Fig. 5-7.

The phylogenetic analysis revealed that the sheep *RAB1A* gene has a closer genetic relationship with the *RAB1A* genes of human, mouse and rat. The sheep

RAB4A and *RAB5A* genes both have closer genetic relationships with the *RAB4A* and *RAB5A* genes of cattle.

Tissue expression profile: Tissue expression profile analysis was carried out and results revealed that the sheep *RAB1A*, *RAB4A* and *RAB5A* genes are all generally but differentially expressed in tissues including spleen, lung, muscle, kidney, ovary, skin, liver, heart and fat (Fig. 8).

In the current study, the researchers firstly get the coding sequences of sheep *RAB1A*, *RAB4A* and *RAB5A* genes by RT-PCR. With the development of modern bioinformatics and establishment of specific sheep NCBI EST database, researchers can easily find the useful ESTs which were highly homologous to the coding sequences of human genes. Based on these sheep EST sequences, the researchers can obtain the complete coding sequences of some novel sheep genes through the some experimental methods such as RT-PCR. From the clone and sequence analysis of sheep *RAB1A*, *RAB4A* and *RAB5A* genes, it could be seen that this is an effective method to isolate some novel sheep genes.

Through sequence analysis, the researchers found that the encoding protein of the sheep *RAB1A*, *RAB4A* and *RAB5A* genes are highly homologous with *RAB1A*, *RAB4A* and *RAB5A* proteins of human and some other

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Mouse      -MANRGATRPNPNTGNKICQFKLVLLGESAVGKSSLVLRVFKGQFHEFQESTIGAAFLT
Rat        -MANRGATRPNPNTGNKICQFKLVLLGESAVGKSSLVLRVFKGQFHEFQESTIGAAFLT
Cattle     -MANRGATRPNPNTGNKICQFKLVLLGESAVGKSSLVLRVFKGQFHEFQESTIGAAFLT
Pig        -MANRGATRPNPNTGNKICQFKLVLLGESAVGKSSLVLRVFKGQFHEFQESTIGAAFLT
Sheep      -MANRGAARPNPNTGNKICQFKLVLLGESAVGKSSLVLRVFKGQFHEFQESTIGAAFLT
Rabbit     -MANRGATRPNPNTGNKICQFKLVLLGESAVGKSSLVLRVFKGQFHEFQESTIGAAFLT
Human      -MASRGATRPNPNTGNKICQFKLVLLGESAVGKSSLVLRVFKGQFHEFQESTIGAAFLT
Chimpanzee -MASRGATRPNPNTGNKICQFKLVLLGESAVGKSSLVLRVFKGQFHEFQESTIGAAFLT
Sumatran orangutan -MANRGATRPNPNTGNKICQFKLVLLGESAVGKSSLVLRVFKGQFHEFQESTIGAAFLT
Dog_Horse  -MANRGATRPNPNTGNKICQFKLVLLGESAVGKSSLVLRVFKGQFHEFQESTIGAAFLT
Rhesus monkey -MANRGATRPNPNTGNKICQFKLVLLGESAVGKSSLVLRVFKGQFHEFQESTIGAAFLT
Chicken    -MANRGATRPNPNTGNKICQFKLVLLGESAVGKSSLVLRVFKGQFHEFQESTIGAAFLT
African clawed frog MANRGGATRPNPNTGNKICQFKLVLLGESAVGKSSLVLRVFKGQFHEFQESTIGAAFLT
Western clawed frog MANRGGATRPNPNTGNKICQFKLVLLGESAVGKSSLVLRVFKGQFHEFQESTIGAAFLT
Zebrafish  MANRGGATRPNPNTGNKICQFKLVLLGESAVGKSSLVLRVFKGQFHEFQESTIGAAFLT
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Mouse      QTVCLDDITVKFEIWDTAGQERYHSLAPMYRGAQAIVVYDITNEESFARAKNWKELQ
Rat        QTVCLDDITVKFEIWDTAGQERYHSLAPMYRGAQAIVVYDITNEESFARAKNWKELQ
Cattle     QTVCLDDITVKFEIWDTAGQERYHSLAPMYRGAQAIVVYDITNEESFARAKNWKELQ
Pig        QTVCLDDITVKFEIWDTAGQERYHSLAPMYRGAQAIVVYDITNEESFARAKNWKELQ
Sheep      QTVCLDDITVKFEIWDTAGQERYHSLAPMYRGAQAIVVYDITNEESFARAKNWKELQ
Rabbit     QTVCLDDITVKFEIWDTAGQERYHSLAPMYRGAQAIVVYDITNEESFARAKNWKELQ
Human      QTVCLDDITVKFEIWDTAGQERYHSLAPMYRGAQAIVVYDITNEESFARAKNWKELQ
Chimpanzee QTVCLDDITVKFEIWDTAGQERYHSLAPMYRGAQAIVVYDITNEESFARAKNWKELQ
Sumatran orangutan QTVCLDDITVKFEIWDTAGQERYHSLAPMYRGAQAIVVYDITNEESFARAKNWKELQ
Dog_Horse  QTVCLDDITVKFEIWDTAGQERYHSLAPMYRGAQAIVVYDITNEESFARAKNWKELQ
Rhesus monkey QTVCLDDITVKFEIWDTAGQERYHSLAPMYRGAQAIVVYDITNEESFARAKNWKELQ
Chicken    QTVCLDDITVKFEIWDTAGQERYHSLAPMYRGAQAIVVYDITNEESFARAKNWKELQ
African clawed frog QTVCLDDITVKFEIWDTAGQERYHSLAPMYRGAQAIVVYDITNEESFARAKNWKELQ
Western clawed frog QTVCLDDITVKFEIWDTAGQERYHSLAPMYRGAQAIVVYDITNEESFARAKNWKELQ
Zebrafish  QTVCLDDITVKFEIWDTAGQERYHSLAPMYRGAQAIVVYDITNEESFARAKNWKELQ
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Mouse      RQASPNIVIALSGNKADLANKRAVDFQEAQSYADDNSLLFMETSAKTSMNVNEIFMAIAK
Rat        RQASPNIVIALSGNKADLANKRAVDFQEAQSYADDNSLLFMETSAKTSMNVNEIFMAIAK
Cattle     RQASPNIVIALSGNKADLANKRAVDFQEAQSYADDNSLLFMETSAKTSMNVNEIFMAIAK
Pig        RQASPNIVIALSGNKADLANKRAVDFQEAQSYADDNSLLFMETSAKTSMNVNEIFMAIAK
Sheep      RQASPNIVIALSGNKADLANKRAVDFQEAQSYADDNSLLFMETSAKTSMNVNEIFMAIAK
Rabbit     RQASPNIVIALSGNKADLANKRAVDFQEAQSYADDNSLLFMETSAKTSMNVNEIFMAIAK
Human      RQASPNIVIALSGNKADLANKRAVDFQEAQSYADDNSLLFMETSAKTSMNVNEIFMAIAK
Chimpanzee RQASPNIVIALSGNKADLANKRAVDFQEAQSYADDNSLLFMETSAKTSMNVNEIFMAIAK
Sumatran orangutan RQASPNIVIALSGNKADLANKRAVDFQEAQSYADDNSLLFMETSAKTSMNVNEIFMAIAK
Dog_Horse  RQASPNIVIALSGNKADLANKRAVDFQEAQSYADDNSLLFMETSAKTSMNVNEIFMAIAK
Rhesus monkey RQASPNIVIALSGNKADLANKRAVDFQEAQSYADDNSLLFMETSAKTSMNVNEIFMAIAK
Chicken    RQASPNIVIALSGNKADLANKRAVDFQEAQSYADDNSLLFMETSAKTSMNVNEIFMAIAK
African clawed frog RQASPNIVIALSGNKADLANKRAVDFQEAQSYADDNSLLFMETSAKTSMNVNEIFMAIAK
Western clawed frog RQASPNIVIALSGNKADLANKRAVDFQEAQSYADDNSLLFMETSAKTSMNVNEIFMAIAK
Zebrafish  RQASPNIVIALSGNKADLANKRAVDFQEAQSYADDNSLLFMETSAKTSMNVNEIFMAIAK
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Mouse      KLPKNEPQNPGANSARGRG-VDLTEPAQPARSQCCSN
Rat        KLPKNEPQNPGANSARGRG-VDLTEPAQPARSQCCSN
Cattle     KLPKNEPQNPGANSTRGRG-VDLTEPTQPTRSQCCSN
Pig        KLPKNEPQNPGINCTRGRG-VDLTEPTQPTRSQCCSN
Sheep      KLPKNEPQNPGAIPPRGRG-VDLTEPTQPTRSQCCSN
Rabbit     KLPKNEPQNPGGNSARGRG-VDLTEPTQPTRSQCCSN
Human      KLPKNEPQNPGANSARGRG-VDLTEPTQPTRSQCCSN
Chimpanzee KLPKNEPQNPGANSARGRGVLDLTEPTQPTRSQCCSN
Sumatran orangutan KLPKNEPQNPGANSARGRG-VDLTEPTQPTRSQCCSN
Dog_Horse  KLPKNEPQNPGANSARGRG-VDLTEPTQPTRSQCCSN
Rhesus monkey KLEKDEPQNPGANSARGRG-VDLTEPTRSQPTRSQCCSN
Chicken    KLPKNEPQNTGASSARGRG-VDLTEPTQPPKSQCCSN
African clawed frog KLPKTEPQAGASNTIRGRG-VDLTETAQPTKSQCCSN
Western clawed frog KLPKTEPQAGGSNTIRGRG-VDLTETAQPTKSQCCSN
Zebrafish  KLPKSEPQAAGANSGRSRG-VDLTETAQPTKAPCCSN
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Fig. 4: The alignment of the protein encoded by sheep *RAB5A* gene and fifteen other kinds of *RAB5A* proteins

animals. This implied that the *RAB1A*, *RAB4A* and *RAB5A* genes were highly conserved in some species and the sheep *RAB1A*, *RAB4A* and *RAB5A* genes will have similar functions as the *RAB1A*, *RAB4A* and *RAB5A* genes of human and other animals. The researchers also found that

the sheep *RAB1A*, *RAB4A* and *RAB5A* proteins do not show complete identity to human or other animals. This implied that the sheep *RAB1A*, *RAB4A* and *RAB5A* genes will have some differences in functions to those of human or other mammals.

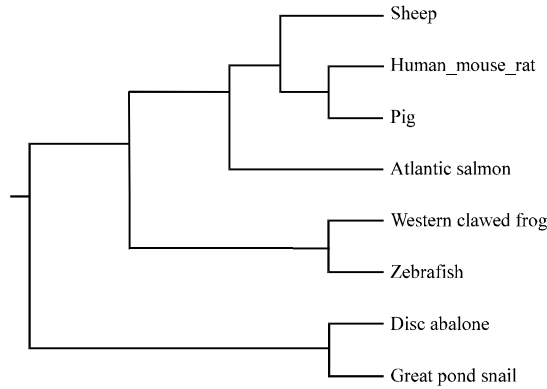


Fig. 5: The phylogenetic analysis for ten kinds of *RAB1A* genes

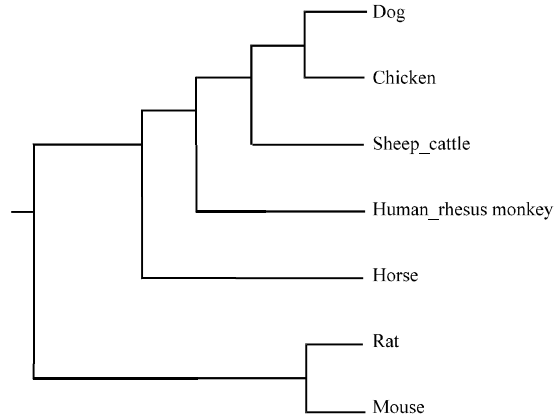


Fig. 6: The phylogenetic analysis for nine kinds of *RAB4A* genes

The phylogenetic analysis revealed that the sheep *RAB1A* gene has a closer genetic relationship with the *RAB1A* genes of human, mouse and rat. This implied that the researchers can use human, mouse and rat as model organisms to study the sheep *RAB1A* gene or use sheep as model organism to study the human, mouse and rat *RAB1A* genes.

The sheep *RAB4A* and *RAB5A* genes both have closer genetic relationships with the *RAB4A* and *RAB5A* genes of cattle. Similarly, we can use cattle as a model organism to study the sheep *RAB4A* and *RAB5A* genes or use sheep as a model organism to study the cattle *RAB4A* and *RAB5A* genes.

From the tissue distribution analysis in the experiment it can be seen that the sheep *RAB1A*, *RAB4A* and *RAB5A* genes were obviously differentially expressed in some tissues. As the researchers did not study functions at protein levels yet, there might be many possible reasons for differential expression of sheep *RAB1A*, *RAB4A* and *RAB5A* genes. The suitable explanation for this under current conditions is that at

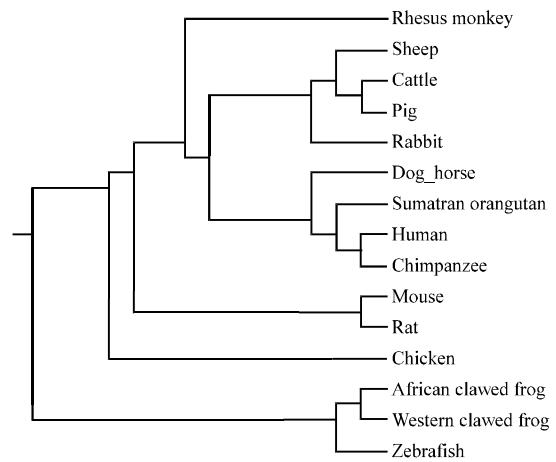


Fig. 7: The phylogenetic analysis for sixteen kinds of *RAB5A* genes

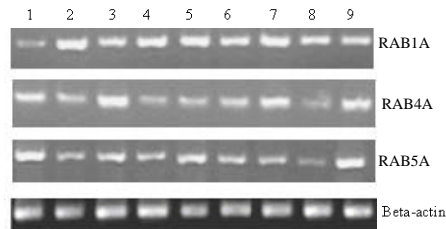


Fig. 8: Tissue expression distribution of sheep *RAB1A*, *RAB4A* and *RAB5A* genes. The beta-actin expression is the internal control. 1: Spleen; 2: Skin; 3: Lung; 4: Muscle; 5: Heart; 6: Fat; 7: Liver; 8: Kidney and 9: Ovary

the same time those biological activities related to the mRNA expression of sheep *RAB1A*, *RAB4A* and *RAB5A* genes were presented diversely in different tissues.

CONCLUSION

In this study, the researchers first isolated the sheep *RAB1A*, *RAB4A* and *RAB5A* genes and performed necessary sequence analysis and tissue transcription profile analysis. This established the primary foundation for further insight into these novel sheep genes.

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REFERENCES

- Diao, A., L. Frost, Y. Morohashi and M. Lowe, 2008. Coordination of golgin tethering and SNARE assembly: GM130 binds syntaxin 5 in a p115-regulated manner. *J. Biol. Chem.*, 283: 6957-6967.
- Esseltine, J.L., L.B. Dale and S.S. Ferguson, 2011. Rab GTPases bind at a common site within the angiotensin II type I receptor carboxyl-terminal tail: Evidence that Rab4 regulates receptor phosphorylation, desensitization and resensitization. *Mol. Pharmacol.*, 79: 175-184.
- Ginsberg, S.D., E.J. Mufson, S.E. Counts, J. Wu, M.J. Alldred, R.A. Nixon and S. Che, 2010. Regional selectivity of rab5 and rab7 protein upregulation in mild cognitive impairment and Alzheimer's disease. *J. Alzheimers Dis.*, 22: 631-639.
- Huang, J., C.L. Birmingham, S. Shahnazari, J. Shiu and Y.T. Zheng *et al.*, 2011. Antibacterial autophagy occurs at PI(3)P-enriched domains of the endoplasmic reticulum and requires Rab1 GTPase. *Autophagy*, 7: 17-26.
- Kachhap, S.K., D. Faith, D.Z. Qian, S. Shabbeer and N.L. Galloway *et al.*, 2007. The N-Myc down regulated Gen1 (NDRG1) Is a Rab4a effector involved in vesicular recycling of E-cadherin. *PloS One*, 2: e844-e844.
- Liu, G.Y. and S.Z. Gao, 2009. Molecular cloning, sequence identification and tissue expression profile of three novel sheep (*Ovis aries*) genes-BCKDHA, NAGA and HEXA. *Biol. Res.*, 42: 69-77.
- Liu, G.Y., 2009. A novel HADHA gene differentially expressed in muscle and other tissues from black-boned vs. ordinary sheep. *Anim. Sci. Pap. Rep.*, 27: 127-137.
- Liu, G.Y., Y.Z. Xiong, C.Y. Deng, B. Zuo and J.H. Zhang, 2004. Comparison of gene expression patterns in Longissimus dorsi of pigs between the high-parent heterosis cross combination Landrace x Large White and the mid-parent heterosis cross combination Large White x Meishan. *Asian-Aust. J. Anim. Sci.*, 17: 1192-1196.
- Matej, R., G. Botond, L. Laszlo, N. Kopitar-Jerala, R. Rusina, H. Budka and G.G. Kovacs, 2010. Increased neuronal Rab5 immunoreactive endosomes do not colocalize with TDP-43 in motor neuron disease. *Exp. Neurol.*, 225: 133-139.
- Schonhoff, C.M., K. Thankey, C.R. Webster, Y. Wakabayashi, A.W. Wolkoff and M.S. Anwer, 2009. Rab4 facilitates cyclic adenosine monophosphate-stimulated bile acid uptake and Na⁺-taurocholate cotransporting polypeptide translocation. *Hepatology*, 48: 1665-1670.
- Wang, C., Y. Yoo, H. Fan, E. Kim, K.L. Guan and J.L. Guan, 2010. Regulation of Integrin β 1 recycling to lipid rafts by Rab1a to promote cell migration. *Biol. Chem.*, 285: 29398-29405.
- Yonggang, L. and G. Shizheng, 2009. A novel sheep gene, MMP7, differentially expressed in muscles from black-boned sheep and local common sheep. *J. Applied Genet.*, 50: 253-256.
- Zhao, Z., X.F. Liu, H.C. Wu, S.B. Zou and J.Y. Wang *et al.*, 2010. Rab5a overexpression promoting ovarian cancer cell proliferation may be associated with APPL1-related epidermal growth factor signaling pathway. *Cancer Sci.*, 101: 1454-1462.