



## RESEARCH ARTICLE

# Differential gene expression in porcine SK6 cells infected with wild-type and SAP domain-mutant foot-and-mouth disease virus

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**Foot-and-mouth disease virus (FMDV) is the causative agent of a highly contagious disease in livestock. The viral proteinase L<sup>pro</sup> of FMDV is involved in pathogenicity, and mutation of the L<sup>pro</sup> SAP domain reduces FMDV pathogenicity in pigs. To determine the gene expression profiles associated with decreased pathogenicity in porcine cells, we performed transcriptome analysis using next-generation sequencing technology and compared differentially expressed genes in SK6 cells infected with FMDV containing L<sup>pro</sup> with either a wild-type or mutated version of the SAP domain. This analysis yielded 1,853 genes that exhibited a  $\geq 2$ -fold change in expression and was validated by real-time quantitative PCR detection of several differentially expressed genes. Many of the differentially expressed genes correlated with antiviral responses corresponded to genes associated with transcription factors, immune regulation, cytokine production, inflammatory response, and apoptosis. Alterations in gene expression profiles may be responsible for the variations in pathogenicity observed between the two FMDV variants. Our results provided genes of interest for the further study of antiviral pathways and pathogenic mechanisms related to FMDV L<sup>pro</sup>.**

**KEYWORDS** Foot-and-mouth disease virus (FMDV); leader protein; SAP region; transcriptome analysis

## INTRODUCTION

Foot-and-mouth disease virus (FMDV) is a positive-stranded RNA virus capable of infecting a variety of domestic and wild biungulate species (Pega et al., 2013). The highly contagious foot-and-mouth disease (FMD) is caused by FMDV and is perhaps the most important lim-

iting factor in the trade of animals and animal products (Barasa et al., 2008; Perry et al., 2007; Rufael et al., 2008), with outbreaks usually resulting in large economic losses for the local livestock industry. FMDV belongs to the *Aphthovirus* genus of the *Picornaviridae* family, and its genome is a single-stranded, positive-sense RNA that encodes a polyprotein. The polyprotein is post-translationally cleaved by three viral proteinases, leader (L<sup>pro</sup>), 2A, and 3C<sup>pro</sup>, into precursors and mature viral structural (VP4, VP2, VP3, and VP1) and nonstructural proteins (L<sup>pro</sup>, 2A, 2B, 2C, 3A, 3B, 3C<sup>pro</sup>, and 3D) (Racaniello, 2007).

Translation of the FMDV polyprotein begins at two different AUG start codons separated by 84 nucleotides, resulting in two alternative forms of L<sup>pro</sup> designated as Lab<sup>pro</sup> and Lb<sup>pro</sup> (Clarke et al., 1985; Piccone et al.,

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1995). During its evolution, FMDV has consistently counteracted against host immune systems to facilitate its survival and replication; several mechanisms have evolved to antagonize host immune responses, with L<sup>pro</sup> reported to play significant pathogenic roles (Grubman et al., 2008). L<sup>pro</sup> is a well-characterized, papain-like proteinase (Medina et al., 1993; Piccone et al., 1995) that can self-cleave from the nascent polyprotein. Host translation-initiation factor eIF-4G can also be cleaved by L<sup>pro</sup>, greatly reducing host cap-dependent mRNA translation without affecting viral cap-independent protein synthesis, which is a characteristic of most picornavirus infections (de Los Santos et al., 2009; Devaney et al., 1988; Kirchweger et al., 1994; Zhu et al., 2010). Additionally, L<sup>pro</sup> inhibits dsRNA-induced type I interferon (IFN) transcription by inhibiting the expression of IFN-regulatory factor 3/7 (Wang et al., 2010).

In eukaryotic cells, the SAP domain (scaffold-attachment factors A and B, apoptotic chromatin-condensation inducer in the nucleus, and protein inhibitor of activated STAT proteins) is a putative DNA-binding domain found in diverse nuclear proteins (Aravind et al., 2000). The SAP domain consists of 35 amino acids, including conserved hydrophobic and charged residues (Aravind et al., 2000), and is found in a number of chromatin-associating proteins, such as scaffold-attachment factors, DNA-repair proteins, RNA-processing complexes, and proto-oncogene proteins (Ahn et al., 2003; Aravind et al., 2000; Bohm et al., 2005; Kipp et al., 2000).

A conserved SAP domain was also identified in the L<sup>pro</sup>-coding region of FMDV (de Los Santos et al., 2009). Genetically engineered FMDV strains lacking the L<sup>pro</sup>-coding region (leaderless viruses) or possessing a mutated L<sup>pro</sup> SAP domain exhibited attenuated viral replication in infected cattle and swine (Chinsangaram et al., 1998; Diaz-San Segundo et al., 2012; Zhu et al., 2010). SAP-domain mutants carrying I55A and L58A substitutions abolish L<sup>pro</sup> retention in the nuclei of FMDV-infected cells and subsequently prevent FMDV-related degradation of nuclear factor-kappa B (NF-κB) (de Los Santos et al., 2007), resulting in upregulation of several cytokines, chemokines, and IFN-stimulated genes (ISGs) (de Los Santos et al., 2009). Additionally, inoculation of swine with FMDV containing a mutated SAP domain induced early protection against disease (Diaz-San Segundo et al., 2012), and transcriptome analysis of embryonic bovine kidney cells (EBKs) infected with SAP-mutated FMDV showed enhanced expression of various IFN-related genes as compared with EBK cells infected with FMDV containing a wild-type SAP domain (de Los Santos et al., 2009).

FMDV pathogenesis presents particular features depending on the host (Pega et al., 2013). In different species, the viral entry routes, primary infective and replic-

ative sites, and, consequently, the associated symptoms and immune responses elicited showed clear differences (Alexandersen et al., 2003). The different host responses triggered by FMDV also correlated with viral replication, thereby affecting viral propagation in different cells. In EBK cells, an intact L<sup>pro</sup> SAP domain was correlated with type I IFN responses (de Los et al., 2006; Zhu et al., 2010). Here, we compared the pathogenic characteristics of FMDV containing wild-type or mutant SAP domains in swine PK15 and SK6 cells and found that the SAP-mutant variant exhibited decreased pathogenicity in both cell lines, with a more pronounced decrease observed in SK6 cells, relative to the wild-type SAP variant.

To analyze the different transcription profiles induced by SAP-domain status, FMDV containing either wild-type or mutant SAP variants was used to infect SK6 cells, followed by comparative transcriptome analysis using next-generation sequencing (NGS) technology to systematically observe the differences in gene expression and host response. Of 20,421 genes detected, differentially enhanced or repressed expression was observed in 1,670 and 183 genes, respectively, with many associated with antiviral responses involving transcription, immune response, inflammation, apoptosis, and cytokines or chemokine production. Our results indicated that the FMDV L<sup>pro</sup> SAP domain was significantly correlated with FMDV viral pathogenicity in SK6 cells, and that the differential expression of various host genes is dependent upon infection with FMDV containing an intact L<sup>pro</sup> SAP domain.

## MATERIALS AND METHODS

### Viruses and cells

An engineered chimeric virus, rA-FMDV, which was previously constructed by Zheng *et al.*, was used as a candidate vaccine (Zheng et al., 2013). The rA-FMDV was constructed by replacing the *PI* gene in the O/CHA/99 strain (GenBank accession number: AF506822) with the *PI* gene from the A/HuBWH/CHA/2009 strain (GenBank accession number: JF792355). A SAP-mutant virus, rA-SAP-FMDV, was previously constructed by Zheng *et al.* by introducing the I55A and L58A mutations into the SAP domain of Lb<sup>pro</sup> in rA-FMDV (unpublished data). The schematic representation of the mutation information is shown in Figure 1A. SK6 cells were purchased from the Cell Bank of Type Culture Collection of the Chinese Academy of Sciences (Shanghai, China). The cells were maintained in Dulbecco's Modified Eagle's Medium (DMEM; Invitrogen, Carlsbad, USA) and cultured in the medium supplemented with 10% fetal bovine serum (FBS) at 37 °C under 5% CO<sub>2</sub>.

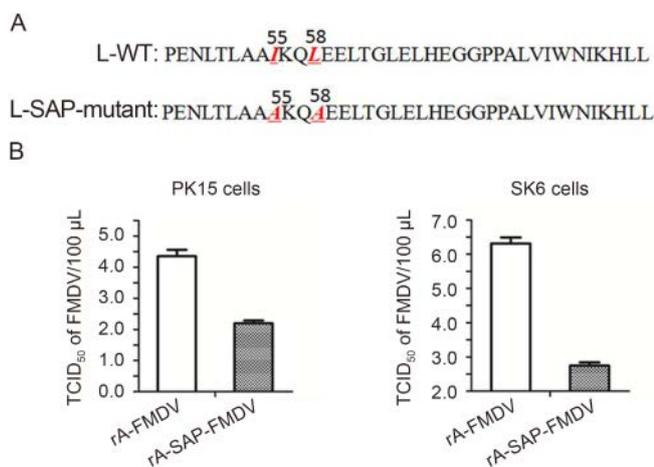


Figure 1. (A) Schematic of the I55A and L58A mutations (red, italics) in the L<sup>pro</sup> SAP domain. (B) Different pathogenicity observed between rA-FMDV and rA-SAP-FMDV in SK6 and PK15 cells. SK6 and PK15 cells were infected with rA-FMDV or rA-SAP-FMDV at similar MOIs, and the viral TCID<sub>50</sub> was detected and recorded. Results are presented as the mean  $\pm$  standard error from three independent experiments.

### Virus infection and 50% tissue culture infectious-dose (TCID<sub>50</sub>) assay

SK6 and PK15 cells were washed with phosphate-buffered saline and infected with FMDV at a multiplicity of infection (MOI) of 1 at 37 °C. After a 1-h adsorption period, the supernatant was removed, and the cells were incubated at 37 °C with DMEM containing 0.5% FBS. The cells used for transcription profile analyses were harvested at 6-h post-infection, because a minimal cytopathic effect (CPE) was observed at ~6 h, enabling a more complete identification of differentially expressed genes. The TCID<sub>50</sub> assay was performed in 96-well plates according to standard procedures, and cells were cultured until CPEs were clearly observed (3–5 days). The TCID<sub>50</sub> value was calculated using the Reed-Muench method (Reed et al., 1938).

### RNA extraction and real-time quantitative PCR (qPCR)

Total RNA was extracted from SK6 cells using TRIzol Reagent (Invitrogen) according to manufacturer protocol. Two micrograms of total RNA was used to synthesize the first strand of cDNA using M-MLV reverse transcriptase (Invitrogen), and the synthesized cDNA were subjected to qPCR analysis performed using SYBR Premix Ex Taq (Takara, Kyoto, Japan) according to manufacturer protocol. The housekeeping gene glyceraldehyde-3-phosphate dehydrogenase was used as an internal control. Primer sequences used in this study are listed

in Supplementary Table S1. The results were obtained from three independent experiments.

### RNA library construction and Solexa/Illumina sequencing

A sample-pooling strategy was performed in this study. The rA-FMDV- and rA-SAP-FMDV-infected samples were used as mixture samples, with each sample prepared by mixing four different dishes of virus-infected cells. After total-RNA extraction and DNase I treatment, magnetic beads conjugated with oligo (dT) were used to isolate mRNA. The mRNA was divided into short fragments, and cDNA was synthesized using the mRNA fragments as templates. The synthesized cDNA was purified and resolved with elution buffer [10 mmol/L Tris-Cl (pH 8.5)] for end repair and single-nucleotide (adenine) addition. Subsequently, the treated fragments were connected using adaptors and subjected to agarose gel electrophoresis, and suitable fragments were selected as templates for PCR amplification. The quality of the obtained library was verified using an Agilent 2100 Bioanalyzer (Agilent, Santa Clara, CA, USA) and an ABI StepOnePlus Real-Time PCR System (Applied Biosystems, Foster City, CA, USA) for quantification and qualification. The library was sequenced using an Illumina HiSeq™ 2000 (Illumina, San Diego, CA, USA), and the raw data was deposited as a National Center for Biotechnology Information (NCBI) BioProject (accession reference: PRJNA269140).

### Gene analysis

To identify the host genes associated with FMDV pathogenicity in SK6 cells, the original data were filtered and screened for differentially expressed genes. The NCBI database was selected as the analytical database, and genetic data containing adaptors and low-quality reads were excluded. The short oligonucleotide analysis package (SOAPaligner/SOAP2) (Li et al., 2008) was used to quickly and accurately align the reads generated by the Illumina/Solexa Genome Analyzer (Illumina); and the reads per kilobase transcriptome per million mapped reads [RPKM;  $RPKM = 10^6 C / (N \times L / 10^3)$ , where C represents the number of reads uniquely aligned to the gene of interest, N is the total number of reads that are uniquely aligned to all genes, and L is the number of bases in the gene of interest (Mortazavi et al., 2008)] was calculated. The RPKM method is able to eliminate the influence of different gene lengths and sequencing discrepancies on the calculation of gene expression. Therefore, the calculated gene expression can be used for comparing differences in gene expression among samples. A method to calculate the significance of digital gene-expression profiles was used for analysis of differentially expressed genes (Audic et al., 1997). We used a false discovery rate (FDR)  $\leq$  0.001 (mascot FDR calculation: <http://www>.

[matrixscience.com/help/decoy\\_help.html](http://matrixscience.com/help/decoy_help.html)) and an absolute value of the  $\text{Log}_2\text{Ratio} \geq 1$  as the thresholds to judge the significance of differences in gene expression. The screened differentially expressed genes were further analyzed by Gene Ontology (GO) and pathway-enrichment analysis. The GO database (<http://www.geneontology.org/>) and GO TermFinder software ([http://smd.stanford.edu/help/GO-TermFinder/GO\\_TermFinder\\_help.shtml/](http://smd.stanford.edu/help/GO-TermFinder/GO_TermFinder_help.shtml/)) were used to perform GO analysis, and the Kyoto Encyclopedia of Genes and Genomes database (Kanehisa et al., 2008) was used for pathway-enrichment analysis.

## RESULTS

### Viral replication in porcine cells differs between FMDV containing wild-type or mutant SAP domain

The mutated region in the SAP domain of rA-SAP-FMDV was confirmed by sequencing analysis. A low MOI leads to infection of a percentage of cells, resulting in these infected cells signaling adjacent, uninfected cells via cytokines in order to activate antiviral genes, including secreted innate immune proteins. To study the signal transduction pathways and proteins involved, infections in this study were performed at 1 MOI. To compare the replication status of rA-FMDV and rA-SAP-FMDV in porcine PK15 and SK6 cells, the cells were infected with equal concentrations of rA-FMDV or rA-SAP-FMDV. The samples were collected 12-h post-infection, and the titers determined by TCID<sub>50</sub> assay. The results showed that rA-FMDV replicated more quickly relative to rA-SAP-FMDV in both PK15 and SK6 cells (Figure 1B), indicating that the SAP mutation decreased FMDV replication in porcine PK15 and SK6 cells, with a larger decrease observed in SK6 cells.

### Differentially expressed genes between SK6 cells infected with FMDV containing wild-type or mutant SAP domain

To explore the differentially expressed genes involved in the altered pathogenicity observed in SK6 cells infected

with FMDV containing the SAP mutation, rA-FMDV-infected and rA-SAP-FMDV-infected SK6 cells were collected at 6-h post-infection, and transcriptome analysis was performed. After a stringent quality check and filtering of the data ( $\text{FDR} \leq 0.001$  and  $\text{fold-change} \geq 2$ ), 20,421 genes were detected, with 1,853 differentially expressed genes identified. A total of 1,670 and 183 genes were differentially upregulated and downregulated, respectively, between rA-SAP-FMDV- and rA-FMDV-infected SK6 cells (Supplementary Figure S1, S2). The expression of 117 transcription factor-related genes involved in 12 biological processes, 114 immune regulation-related genes participating in 40 immune-regulatory processes, 69 cytokine-related genes, including 20 involved in cytokine-production and -secretion processes, 12 inflammatory response-related genes, and 19 apoptosis-related genes were significantly altered (Table 1, Supplementary Table S2–S5). The distinctively different expression profiles of these genes may explain the decreased pathogenicity observed following infection with rA-SAP-FMDV. An analysis of the available literature indicated that the majority of the differentially expressed genes correlating with antiviral responses included (Table 2): 1) genes involved in transcriptional regulation (*EIF4A2*, *EIF5B*, *EIF3J*, *NFKBIA*, and *NFKBIZ*); 2) genes involved in the regulation of immune response (*IFIT1*, *ITCH*, *IL7R*, *JAK2*, *LTB*, *TNFSF10*, *IL7*, *BLM*, *IFIT1*, *IL18*, *IL6*, and *FOS*); 3) cytokine-related genes (*IL1*, *IL6*, *IL20*, *TNF*, *CCL2*, *CCL20*, *CXCL10*, *CXCL2*, *CCL3L1*, *CCL4*, *CCL5*, and *CXCL11*); 4) genes involved in the regulation of inflammation and chemokine production (*TNF*, *CCL5*, *IL1A*, *IL6*, *IL6ST*, *CCL2*, and *ITCH*); and 5) genes involved in apoptosis (*BLM*, *CASP3*, *BRCA2*, *PMAIP1*, *CD38*, *MAP3K5*, *CUL5*, *TNFSF10*, and *XIAP*).

### Validation of differentially expressed genes by qPCR

To further confirm and validate the transcriptome analysis results, we performed qPCR analysis to determine the reproducibility of the differential gene expression. A se-

Table 1. Summary of differential expressed genes

Function	Total gene number	Up- or down-regulated gene number	
		Up-regulated	Down-regulated
transcription factor-related genes	117	109	8
immune regulation-related genes	114	104	10
cytokine-related genes	69	62	7
inflammatory response-related genes	12	11	1
apoptosis-related genes	19	18	1

Table 2. List of genes that displayed significant differential expression at WT and SAP mutant FMDV- infected SK6 cells

Gene	Fold	Gene description	Function
<i>EIF4A2</i>	2.23	Eukaryotic initiation factor 4A-II	RNA helicase activity; adenylyl ribonucleotide binding
<i>EIF5B</i>	3.11	Eukaryotic translation initiation factor 5B	Translation factor activity, nucleic acid binding
<i>EIF3J</i>	2.47	Eukaryotic translation initiation factor 3 subunit J-like isoform 1	Translation factor activity, nucleic acid binding
<i>NFKBIA</i>	3.83	NF-kappa-B inhibitor alpha	Transcription factor binding
<i>NFKBIZ</i>	3.47	NF-kappa-B inhibitor zeta	Transcription cofactor activity, protein binding
<i>IFIT1</i>	3.74	Interferon induced protein with tetratricopeptide repeats 1	RNA binding, protein binding
<i>ITCH</i>	2.87	Itchy E3 ubiquitin protein ligase	Chemokine receptor binding, ubiquitin protein ligase activity
<i>IL7R</i>	4.32	Interleukin 7 receptor	Cytokine receptor activity
<i>JAK2</i>	2.27	Janus kinase 2	Kinase binding, cytokine receptor, protein kinase activity
<i>LTB</i>	13.15	Lymphotoxin-beta	Tumor necrosis factor receptor superfamily binding
<i>TNFSF10</i>	3.67	Tumor necrosis factor superfamily member 10	Cation binding, tumor necrosis factor receptor binding
<i>IL7</i>	2.31	PREDICTED: interleukin-7 isoform 3	Cytokine receptor binding
<i>BLM</i>	3.15	Bloom syndrome protein	ATP-dependent DNA helicase activity, double-stranded DNA binding
<i>IL18</i>	2.27	Interleukin 18	Receptor binding, cytokine activity
<i>IL6</i>	11.05	Interleukin 6	Cytokine receptor binding, cytokine activity
<i>FOS</i>	3.78	FBJ osteosarcoma oncogene	Nucleic acid binding transcription factor activity, protein dimerization activity
<i>IL1</i>	3.11	Interleukin 1	Cytokine activity
<i>IL20</i>	27.75	Interleukin 20	Cytokine receptor binding, cytokine activity
<i>TNF</i>	20.89	Tumor necrosis factor	Tumor necrosis factor receptor superfamily binding, sequence-specific DNA binding
<i>CCL2</i>	5.73	C-C motif chemokine ligand 2	Kinase activity, chemokine receptor binding
<i>CCL20</i>	7.62	C-C motif chemokine ligand 20	Cytokine activity, chemokine receptor binding
<i>CXCL10</i>	5.89	Chemokine (C-X-C motif) ligand 10	Protein kinase regulator activity, cytokine activity
<i>CXCL2</i>	2.38	Chemokine (C-X-C motif) ligand 2	Cytokine activity, chemokine activity
<i>CCL3L1</i>	6.17	C-C motif chemokine ligand 3 like 1	CCR chemokine receptor binding, chemokine activity
<i>CCL4</i>	15.34	C-C motif chemokine ligand 4	Cytokine activity, chemokine activity
<i>CCL5</i>	2.58	C-C motif chemokine ligand 5	CCR chemokine receptor binding, chemokine activity, protein tyrosine kinase activator activity
<i>CXCL11</i>	6.17	C-X-C motif chemokine ligand 11	Heparin binding, chemokine activity
<i>IL1A</i>	3.11	Interleukin-1 alpha precursor	Transition metal ion binding, cytokine receptor binding
<i>IL6ST</i>	3.58	Interleukin 6 signal transducer	Ciliary neurotrophic factor receptor activity, cytokine receptor binding
<i>CASP3</i>	2.08	Caspase 3	Endopeptidase activity, cyclin-dependent protein kinase regulator activity
<i>BRCA2</i>	5.42	Breast cancer 2	Structure-specific DNA binding, histone acetyltransferase activity
<i>PMAIP1</i>	2.78	Phorbol-12-myristate-13-acetate-induced protein 1	Protein binding
<i>CD38</i>	2.51	Cluster of differentiation 38	Transferase activity, NAD(P) <sup>+</sup> nucleosidase activity
<i>MAP3K5</i>	2.23	Mitogen-activated protein kinase kinase kinase 5	Metal ion binding, phosphatase binding, apoptotic protease activator activity
<i>CUL5</i>	3.23	Cullin 5	Signal transducer activity, enzyme binding
<i>XIAP</i>	2.59	X-linked inhibitor of apoptosis	Transition metal ion binding, cysteine-type endopeptidase inhibitor activity

Note: A minimum of twofold change ( $P < 0.0001$ ,  $Q < 0.0001$ ) was used as the standards for selecting genes of interest.

lected group of genes for which we had an established method available in our laboratory (with established primers and melting/annealing temperatures previously) were chosen for analysis. Six upregulated genes (*CCL4*, *CCL2*, *IL6*, *IL7*, *IL18*, and *EGR1*) and four downregulated genes (*SRPX2*, *CREB5*, *RASAL1*, and *RIN2*) were analyzed, and qPCR results confirmed the differential expression identified between rA-SAP-FMDV- and rA-FMDV-infected cells. As shown in Figure 2, the qPCR results corresponded with transcriptome analysis results and, while some fold-change differences were observed between results from each method, similarities in the overall expression profiles were revealed.

### Functional characterization of differentially expressed genes and pathways affected by infection with FMDV containing the SAP mutation

Infection with FMDV containing the SAP-domain mutation altered gene expression in SK6 cells, thereby affecting various biological processes and signal transduction pathways, and resulting in blocked viral replication and decreased pathogenicity. To systematically analyze the functional characterization of the differentially expressed genes and pathways associated with the FMDV L<sup>pro</sup> SAP domain, GO analysis and pathway annotation were conducted. The results indicated that the differentially expressed genes were involved in metabolism, cell cycle processes, and cellular component organization or biogenesis processes (Supplementary Figure S3), and functional analysis revealed that many of these genes were involved in nucleotide binding and functions asso-

ciated with nucleic acids (Supplementary Figure S4). Cellular component annotation results are shown in Supplementary Figure S5.

Pathway analysis indicated that 35 pathways were altered in rA-SAP-FMDV-infected cells as compared with rA-FMDV-infected cells, including regulation of actin cytoskeleton formation, endocytosis, phagosome formation, chemokine-signaling pathways, the cell cycle, the retinoic acid-inducible gene 1-like receptor signaling pathway, the NF- $\kappa$ B signaling pathway, and the nucleotide-binding oligomerization domain-like receptor signaling pathway (Supplementary Figure S6). The potential host targets for L<sup>pro</sup> and the protein-protein interaction pathways involved are shown in Figure 3. These findings suggested that expression of these genes might potentially result in decreased rA-SAP-FMDV replication.

### DISCUSSION

FMDV has the ability to manipulate various host-cell signal transduction pathways by subverting gene expression. FMDV L<sup>pro</sup> is a host-cell antagonist that interferes with host gene expression, promotes viral propagation, and subverts host immune systems by targeting eIF-4G, IRF3, IRF7, and NF- $\kappa$ B (de Los Santos et al., 2009; Kirchweiger et al., 1994; Wang et al., 2010). The FMDV L<sup>pro</sup> SAP domain is a putative DNA-binding domain found in diverse eukaryotic nuclear proteins (Aravind et al., 2000; de Los Santos et al., 2009) and inhibits host innate immune response (de Los Santos et al., 2009). The mutation of two amino acids (I55A and L58A) in the Lb<sup>pro</sup> SAP domain alters L<sup>pro</sup> subcellular localization and function (de Los Santos et al., 2009; Diaz-San Segundo et al., 2012).

In a previous study, we constructed a chimeric virus, rA-FMDV, as a candidate vaccine (Zheng et al., 2013). To decrease rA-FMDV pathogenicity and develop a potential live attenuated vaccine, we constructed rA-SAP-FMDV containing a mutated SAP domain (unpublished data). Here, we compared the different pathogenic characteristics of rA-FMDV and rA-SAP-FMDV in PK15 and SK6 cells, and showed that rA-SAP-FMDV was less pathogenic relative to rA-FMDV in both cell lines, suggesting that rA-SAP-FMDV may have potential as a vaccine strain based on its failure to disrupt cellular responses that inhibit viral replication.

Decreased rA-SAP-FMDV replication was more evident in SK6 cells as compared with PK15 cells. To analyze the differentially expressed genes correlated with the altered viral replication observed in SK6 cells, transcriptome analysis was performed using NGS technology. The results indicated differential expression of

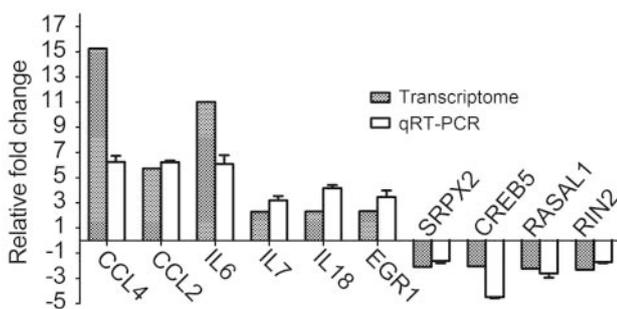


Figure 2. Validation of differentially expressed genes identified by transcriptome analysis through qPCR detection. Six upregulated and four downregulated genes were detected in an independent infection experiment undertaken in order to validate transcriptome analysis results. The expression profiles of the 10 selected genes were consistent between the transcriptome-analysis and qPCR-detection results. Results are presented as the mean  $\pm$  standard error from three independent experiments.

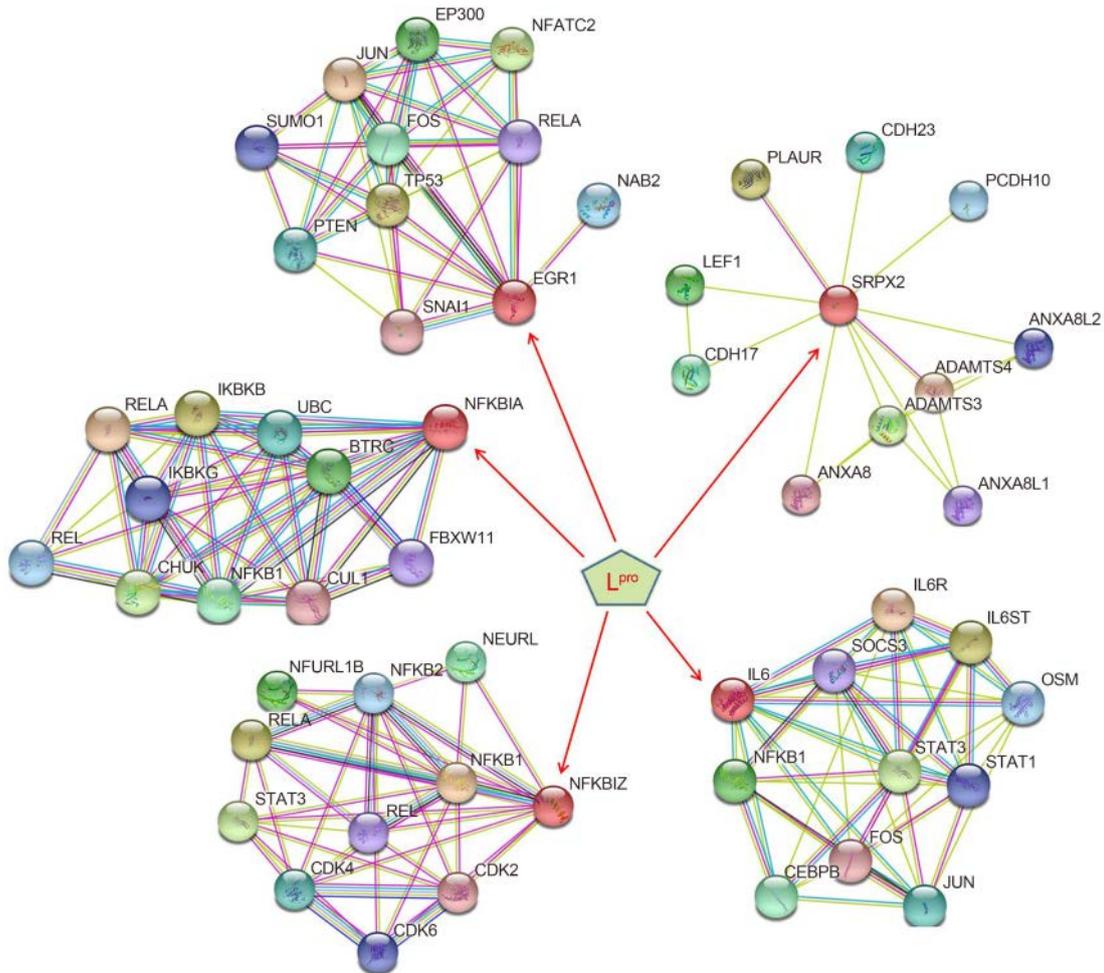


Figure 3. The potential host targets of  $L^{pro}$  and the involved protein-protein interaction pathways.

1, 853 genes between infected and non-infected SK6 cells, with these findings subsequently confirmed by qPCR analysis. These findings suggested that mutation of the FMDV  $L^{pro}$  SAP domain might adversely affect the ability of FMDV to inhibit host gene expression during infection, resulting in reduced viral pathogenicity.

Among the differentially expressed genes upregulated in rA-SAP-FMDV-infected cells, *EIF4A2*, *EIF5B*, and *EIF3J* are involved in the initiation of host translation by aiding in the recruitment of protein and mRNA components to ribosomes (Cheyssac et al., 2006; ElAntak et al., 2007; Kyono et al., 2002; Meijer et al., 2013; Unbehaun et al., 2007). Swine infected with FMDV containing a mutated SAP domain developed a strong neutralizing-antibody response as early as 2-days post-inoculation as compared with those infected with wild-type FMDV (Diaz-San Segundo et al., 2012). The upregulation of these translation factors possibly resulted in the enhancement of neutralizing-antibody production. Furthermore, our analysis revealed upregulation of other genes, including those involved in metabolic and cellular-response

processes (Supplementary Figure S3).

FMDV infection can induce degradation of NF- $\kappa$ B (de Los Santos et al., 2007); however, NF- $\kappa$ B activity was significantly enhanced in cells infected with FMDV containing a mutated SAP domain (Zhu et al., 2010). *NFKBIA* and *NFKBIZ* are involved cytokine production through NF- $\kappa$ B regulation (Ninomiya-Tsuji et al., 1999; Yamazaki et al., 2001). In this study, we found that the expression of *NFKBIA* and *NFKBIZ* was upregulated in rA-SAP-FMDV-infected cells, which may have altered the subsequent expression of NF- $\kappa$ B-induced cytokines to ensure a robust immune response (Figure 3). *CCL4* and *IL7*, both involved in cellular immune and inflammatory responses, were also upregulated in these cells. Additionally, the upregulation of many other genes involved in immune response, inflammation, chemokine production, and apoptosis was also observed.

Upregulation of *EGR1* and *IL6* expression was also observed in rA-SAP-FMDV-infected cells. *EGR1* mediates cell proliferation, differentiation, inflammation, and apoptosis (Han et al., 2015), and may interact with p53

or FOS to regulate apoptosis or transcription (Figure 3). IL6 is involved in inflammation, B cell maturation, and suppression of viral replication (Dienz et al., 2012). Here, upregulation of *IL6* expression possibly enhanced acute-phase response and suppressed rA-SAP-FMDV replication. Our results suggested that L<sup>pro</sup> may attenuate *EGR1* and *IL6* expression and adversely affect regulation of cell proliferation, transcription, differentiation, inflammation, immune response, and apoptosis to promote viral replication. Furthermore, mutation of the L<sup>pro</sup> SAP domain might impair this antagonistic effect, thereby inhibiting viral replication (Figure 3). The resulting upregulation of these genes likely reinforced the antiviral activity of the cells, directly resulting in the decreased pathogenicity observed following rA-SAP-FMDV infection. Conversely, downregulation of *SRPX2*, which is involved in anti-apoptotic activity, was observed in rA-SAP-FMDV-infected SK6 cells (Figures 2 and 3). This indicated an alternative pathway for suppressing viral replication through the promotion of cell death. Intact L<sup>pro</sup> likely inhibits the initiation of apoptosis, with our results suggesting that mutation of the L<sup>pro</sup> SAP domain impaired this inhibitory effect.

A previous study found that in EBK cells infected with a FMDV containing the SAP-domain mutation, NF- $\kappa$ B was the primary factor responsible for the differential transcription of many upregulated genes associated with innate immune response (Zhu et al., 2010). Here, we found that the differential gene expression observed in rA-SAP-FMDV-infected SK6 cells resulted from altered expression of genes involved in transcription and immune-related regulation. However, IFN-stimulated genes, such as *ISG15*, *ISG20*, *MX1*, *GBP1*, and *OAS1*, which were differentially expressed in EBK cells infected with a FMDV containing the SAP-domain mutation, were not observed in this study. This is possibly due to SK6 cells being deficient in type I IFN production (Ruggli et al., 2003), although the enhancement of various type I IFN-independent genes and pathways determined in this study were implied to perform crucial antiviral effects.

In summary, we reported that a FMDV L<sup>pro</sup> SAP-domain mutant exhibited decreased replication ability in SK6 cells as compared with wild-type FMDV. Transcriptome analysis suggested that the altered expression of genes involved in transcription, immune response, cytokine and chemokine production, inflammation, and apoptosis were the primary reasons for the observed decrease in pathogenicity. Our results provided insight into the pathogenic mechanisms associated with the FMDV L<sup>pro</sup> SAP-domain and suggested that mutation of region

provides a strategy for the development new FMDV-related vaccines having impaired host-antagonistic ability.

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## COMPLIANCE WITH ETHICS GUIDELINES

The authors declare that they have no conflict of interest. This article does not contain any studies with human or animal subjects performed by any of the authors.

## AUTHOR CONTRIBUTIONS

ZXN, FY, ZXZ and HXZ designed the research, ZXN, FY, ZXZ, XD, WWL and WJC performed the experiments. XLZ, YJ, JHG and XTL provided experiment support. ZXN, ZXZ and HXZ wrote the manuscript. All authors have read and approved the final manuscript for submission.

Supplementary figures/tables are available on the website of *Virologica Sinica*: [www.virosin.org](http://www.virosin.org); [link.springer.com/journal/12250](http://link.springer.com/journal/12250).

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