



Identification of New Isolates of Classical Swine Fever Virus Based on Untranslated Regions with Traditional and Machine Learning Classification

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ABSTRACT

Classical swine fever is a deadly and economically significant pig disease caused by the classical swine fever virus (CSFV). This disease is one of the most contagious viral infections affecting pig herds. Since late 2014 China has had a rising number of C-strain vaccinated swine farms and experienced substantial damages due to outbreaks of the recently discovered subgenotype 2.1d. We aimed to sequence and characterize untranslated regions (5'- and 3'-UTR) of eight new CSFV strains. For this purpose, we did a phylogenetic analysis of new CSFV strains and built neighborhood-based phylogenetic trees. We found that the Heilongjiang isolates HL2018-0494, HL2018-016 HL2016-0205 and Inner Mongolia isolates NM2016-0323 were grouped into subgenotype 2.1d, while the remaining isolates HL2018-0490, HL2018-0462, NM2016-0333 and (Shengdong) SD2018-0461 were grouped into subgenotype 2.1b. We used the support vector machine (artificial intelligence tool) method for more validation and obtained similar results. We demonstrated that monoclonal antibody (mAb) 5B8-2 strongly reacted with all CSFV strains. The new isolates showed molecular alterations in the 5'- and 3'-UTR regions. The findings exposed the genetic diversity and molecular features of CSFV, currently circulating in China, and provided insight into the new pattern of epidemiology. The present study could be useful for building new vaccines and updated CSF diagnostic strategies.

Article Information

Received 25 June 2024

Revised 09 September 2024

Accepted 23 September 2024

Available online 25 January 2025
(early access)

Published 29 December 2025

Authors' Contribution

MF, IN and YW conceived and designed the study. MF executed the experiment and analyzed the tissue samples and wrote the original draft preparation. IN and FR analyzed the data, reviewing and editing. YW, IA, KD, BF formal analysis. DF provides resources and funding. All authors interpreted the data, critically revised the manuscript for important intellectual contents and approved the final version.

Key words

Genome sequencing, Molecular characterization, Classical swine fever virus, subgenotype 2.1, Untranslated region (UTR)

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0030-9923/2026/0001-0371 \$ 9.00/0



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INTRODUCTION

Classical swine fever (CSF) is a communicable infectious disease of pigs, including both domestic and wild types, with a devastating impact on the swine industry, as it causes large economic losses (Postel *et al.*, 2012). China provides more than half the world's pigs, therefore being the largest producer in the world (Zhou, 2019). CSF is caused by the classical swine fever virus

(CSFV), which has a single-stranded, positive-sense RNA genome of 12.3 kb. It is a member of the *Pestivirus* genus and belongs to the *Flaviviridae* family (Zhou, 2019). On either side of the nucleic acid, there are 5' and 3' untranslated regions (UTRs) and a single open reading frame (ORF). It encodes eight nonstructural N^{pro}, p7, NS2, NS3, NS4A, NS4B, NS5A, and NS5B and four structural proteins i.e. C, E^{ms}, E1, and E2 (Rümenapf *et al.*, 1993; Paton *et al.*, 2000). The structural E2 protein aids in the generation of neutralizing antibodies through virus attachment and entrance into the cell, hence protecting pigs from a virulent challenge (Sánchez *et al.*, 2008). The 5'- and 3'-UTRs are important regulatory components of the genome. 5'-UTR is involved in polyprotein translation and replication (Hsu *et al.*, 2014). It has been shown that the 5'-terminal nucleotides (nt) 1–29 had an inhibitory effect on the translation of CSFV that is facilitated by IRES (the internal ribosome entry site) (Xiao *et al.*, 2009). The 3'-UTR is involved in negative-strand RNA synthesis by generating many structural elements (Deng and Brock, 1993). Furthermore, the 3'-UTR affects translation by interacting with the 5'-UTR or in certain other ways (Ito *et al.*, 1998). The conserved sequences at the UTRs suggest the signals of viral RNA replication (Fletcher and Jackson, 2002; Xiao *et al.*, 2004).

CSFV genotypes 1, 2, and 3 have a total of 11 sub-genotypes i.e., including 1.1, 1.2, 1.3, and 1.4; 2.1, 2.1a, 2.1b, 2.1c, 2.1d, 2.2, 2.3; and 3.1, 3.2, 3.3, and 3.4 (Beer *et al.*, 2015; Hu *et al.*, 2016). Since the 1980s, genotype 2.1 has been circulating, particularly the 2.1b strain was dominant and endemic in most regions of China (Zhang *et al.*, 2015).

All RNA viruses that include CSFV have higher mutation rates than DNA viruses, the reason behind this is that RNA-dependent RNA polymerase (RdRp) lacks proof-reading ability, resulting in poor fidelity replication (Drake and Holland, 1999; Hughes and Hughes, 2007). Nonetheless, the high mutation frequency might be advantageous for the viruses, allowing them to protect from the host immune responses (Sanjuán *et al.*, 2010).

Recently, sequencing based on untranslated and coding regions of CSFV has been a potent method to trace changes in the virus population over time (Postel *et al.*, 2012) or to characterize strains responsible for outbreaks in combination with epidemiological surveys (Greiser-Wilke *et al.*, 2000; Shen *et al.*, 2013). Moreover, several authors hypothesized that mutations (nucleotide substitutions or insertions/deletions) may accumulate in CSFV strains due to vaccination pressure and thus, in the near future, C-strain-based vaccines may lose the efficacy of protecting pigs against all strains of CSFV (Sanjuán *et al.*, 2010; Ji *et al.*, 2015; Yoo *et al.*, 2018).

Although extensive vaccination strategies have been implemented in China, CSF outbreaks are still common, as demonstrated by recent cases in pig herds vaccinated with C-strain (Luo *et al.*, 2014; Zhang *et al.*, 2018). Therefore, the present study aims to identify all CSFV genotypes and sub-genotypes circulating in different regions of China based on analysis of the untranslated regions.

MATERIALS AND METHODS

CSFV strains

The present study includes eight strains of CSFV Heilongjiang (HL2018-0494, HL2018-016, HL2016-0205, HL2018-0490, HL2018-0462), Inner Mongolia (NM2016-0323, NM2016-0333) and Shengdong (SH2018-0461), which have previously been separated from tissue samples (lymph node, spleen, kidney, and tonsil). These samples were taken from deceased or ill pigs suspected of having CSF at various swine farms in different locations in eastern China. Furthermore, these strains were also identified and characterized earlier by PCR.

RT-PCR, cloning and sequencing

Genomic material (RNA) of CSFV strains extracted from cell cultures and synthesis of cDNA performed and used to amplify the 5'-UTR and 3'-UTR regions by RT-PCR using two pairs of designed primers CSFV-FP-UTR (1-21)-5'-GTATACGAGGTTAGCTCATCC-3', CSFV-RP-UTR (988-1008)-3'-CTCTACCACAATCGTAGCATC-5'; CSFV-FP-UTR (11054-11075)-5'-CTATGCACATGTCAGAAGTACC-3', CSFV-RP-UTR (12259-12279)-3'-ACCTTAGTCCAACTATGGACG-5'. The resultant product was stained with gene SafeViewTM (Applied Biological Material, Richmond, BC, Canada) and electrophoresed on 1.5% gel of agarose before being seen using the Gel Doc XR+ system. The amplified product was cloned into the pMD^{18-T} vector before it was incorporated into *E. coli* (DH5 α) cells (TaKaRa, Dalian, China). The colonies were grown before being delivered to Comate Bioscience in Jiangsu, China, for sequencing.

Phylogenetic analysis and molecular characterization

Multiple sequence alignments of these eight strains of CSFV were generated using the Laser gene (Version 7.1., DNASTAR Inc., Madison, WI, USA) and MEGA6 Software (Center for Evolutionary Functional Genomics, the Biodesign Institute, Tempe, AZ, USA). The neighborhood-joining (NJ) based phylogenetic trees were constructed with MEGA6 (Tamura *et al.*, 2011).

Immunofluorescence assay

Virus confirmation and titration in the cell line were

determined by immunofluorescence assay (IFA) based on the EU Diagnostic Manual (EU Reference Laboratory for CSF (EURL), 2020).

Support vector machines and genetic type prediction

To further validate our results, we employed a Support Vector Machine (SVM) model (Shapshak *et al.*, 2019) to perform class predictions on 8 new chains. SVM is a supervised learning algorithm, which realizes classification by finding an optimal hyperplane. SVM is particularly suitable for complex datasets and high-dimensional feature spaces, making it an ideal choice for our research. Its ability to find the best decision boundary ensures accurate classification of the new chains. At first, we used k-mers as features to convert the CSFV gene sequence data into feature vector representations. This method, as described by Rahman *et al.* (2018), can transform sequence data (such as DNA sequences or protein sequences) into k consecutive character Subsequence, called k-mers. These k-mers can be used as feature representations for sequence data. By calculating the frequency of each k-mers in the sequence, a feature vector can be constructed, where each element represents a different k-mers and represents the number or frequency of occurrences of that k-mers in the sequence. Then, we trained on previous data sets using the SVM model and classified the new 8 CSFV chains. In the process of classifying CSFV, a linear kernel function is used for classification and a one-to-many strategy is used to handle multi-class classification problems. Furthermore, the performance of the model is evaluated through indicators such as Accuracy, Recall, and F1-score. Finally, to gain insights into the classification results, we employed PCA dimensionality reduction (Groth *et al.*, 2013) to visualize the distribution of the sequences additionally. PCA is a dimensionality reduction technique that allows us to represent the data in a lower-dimensional space while preserving its essential characteristics. In our study, we applied PCA to reduce the dimensionality of the feature vectors extracted from the CSFV gene sequences. By reducing the dimensionality to three components, we could visualize the classification results in a three-dimensional scatter plot.

RESULTS AND DISCUSSION

The samples were detected CSFV with IFA and processed next (Fig. 1). All selected monoclonal antibodies were specifically generated against the epitopes of the E2 protein. Epitopes of structural E2 proteins are important not only for the early stage of CSFV infection but also for serological differentiation of CSFV from other ruminant pestiviruses, given strong cross-reactivity with sheep and

goat sera in ELISA and neutralization assay (Huang *et al.*, 2021). The obtained sequence data sets of UTRs regions were subjected to phylogenetic analysis and investigated systematically to gain the discriminatory ability of the individual genomic regions. In total, 25 and 30 genome sequences including the 8 obtained strains together with 21 and 22 reference sequences were used in the construction of the phylogenetic trees based on the 5'- and 3'-UTRs. The isolates (HL2018-0494, NM2016-0323, HL2018-016 and HL2016-0205) were clustered into sub-genotype 2.1d, while the rest isolates (HL2018-0490, SD2018-0461, HL2018-0462 and NM2016-0333) were clustered into sub-genotype 2.1b (Fig. 2).

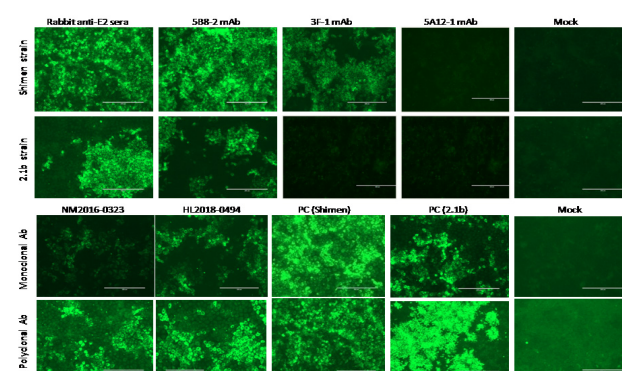


Fig. 1. Virus confirmation for reference control strains (Shimen and 2.1b) and representative new strains (NM2016-0323 and HL2018-0494) by indirect fluorescence assay (IFA) in PK-15 cells. The figure showed that mAb (monoclonal antibody) 5B8-2 was effective against both reference control strains. mAb 3F-1 was compatible with the Shimen strain but did not react appropriately with the 2.1b control strain. No fluorescence was observed for the 5A12-1 antibody with all strains. Mock= negative control; PC= positive control. The scale bar is 200 μ m.

Table I. Sample collection and GenBank database information of the UTRs regions.

No	Isolate name	Sample collection date	GenBank accession no. for 3'UTR	GenBank accession no. for 5'UTR
1	HL2016-0205	2018-6-30	OQ150757	OQ150765
2	HL2018-0416	2018-6-30	OQ150758	OQ150766
3	HL2018-0462	2016-6-16	OQ150759	OQ150767
4	HL2018-0490	2016-3-18	OQ150760	OQ150768
5	HL2018-0494	2018-6-29	OQ150761	OQ150769
6	NM2016-0323	2018-4-29	OQ150762	OQ150770
7	NM2016-0333	2018-6-30	OQ150763	OQ150771
8	SD2018-0461	2018-6-29	OQ150764	OQ150772

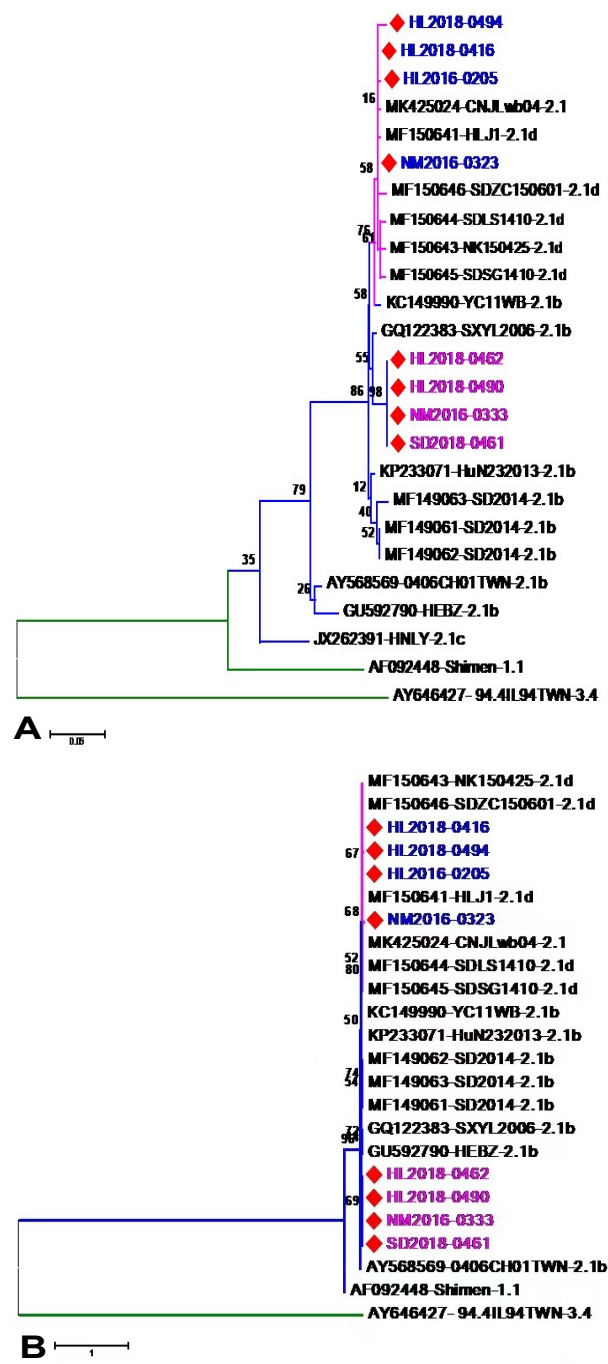


Fig. 2. Phylogenetic analysis of the novel CSFV strains based on 5'- 3'-UTRs regions. (A) Phylogenetic analysis of the 8 novel strains and 17 reference strains based on 5'-UTR. All novel CSFV isolates were designated by red diamonds. (B) Phylogenetic analysis of the 8 novel strains and 16 reference strains based on 3'-UTR. All novel CSFV isolates were designated by red diamonds. The phylogenetic tree is built using the neighbor-joining (NJ) method, with 1000 bootstrap replicates.

Table II. Results of SVM model predictions. The SVM model successfully predicted the category of each chain, regardless of whether the 5'-UTR or the 3'-UTR data was used for training.

Gene sequence	8 Strains ID	Prediction
5'-UTR/3'-UTR	HL2018-0462	Subgenotype 2.1b
	HL2018-0490	
	NM2016-0333	
	SD2018-0461	
	NM2016-0323	
	HL2018-0494	
	HL2018-0416	Subgenotype 2.1d
	HL2016-0205	

The classification results of the SVM model on our dataset achieved perfect Accuracy, Recall, and F1-score, as indicated in Table II. Whether using only the 5'-UTR or the 3'-UTR data for training the SVM model, we were able to successfully predict the categories of the new eight chains, and the prediction results were consistent with our phylogenetic tree analysis. That is, chains HL2018-0462, HL2018-0490, NM2016-0333, and SD2018-0461 were predicted as subgenotype 2.1b, while chains NM2016-0323, HL2018-0494, HL2018-0416, and HL2016-0205 were predicted as subgenotype 2.1d. Figure 3 illustrates the distinct separation of different categories in the PCA-reduced and visually represented classification results, indicating that the SVM model effectively captures

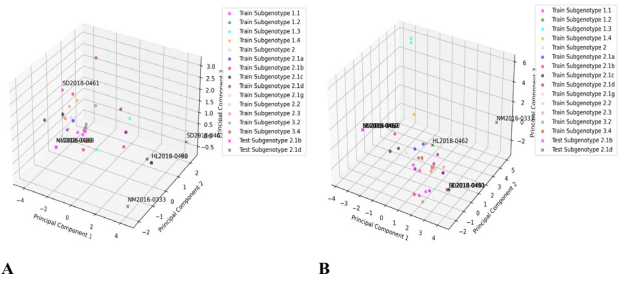


Fig. 3. PCA (Principal component analysis) visualization of sequence classification results. A, the SVM model predicts HL2018-0462, HL2018-0490, NM2016-0333, and SD2018-0461 as Subgenotype 2.1b, while NM2016-0323, HL2018-0494, HL2018-0416, and HL2016-0205 are predicted as Subgenotype 2.1d. This pattern is consistent with the predictions in B, where the classification results based on the 3'-UTR region also indicate HL2018-0462, HL2018-0490, NM2016-0333, and SD2018-0461 as Subgenotype 2.1b, and NM2016-0323, HL2018-0494, HL2018-0416, and HL2016-0205 as Subgenotype 2.1d. The overlapping labels or data points are again a result of the high sequence similarity.

and sub-genotype 2.1 strains exhibited some discontinuous nucleotide deletions compared with 1.1 isolates. Moreover, the interaction between 3'-UTR nucleotide deletions and other genes needs further investigation.

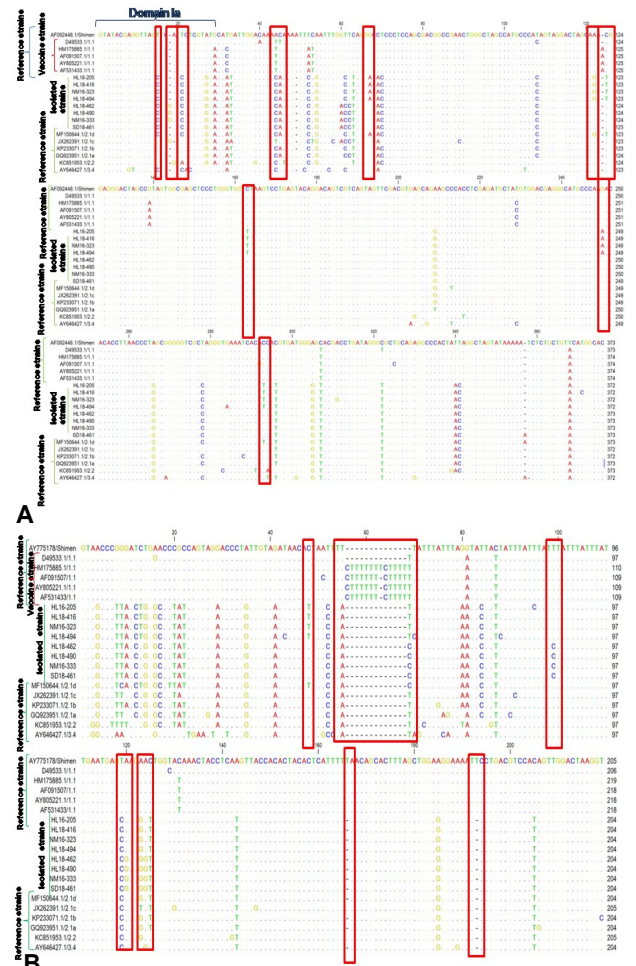


Fig. 4. Sequence alignment of isolated, reference and vaccine strains. A, 5'-UTR amino acid sequence alignment of the 8 studied novel CSFV strains and 12 reference isolates downloaded from NCBI (National Center for Biotechnology Information). B, 3'-UTR amino acid sequence alignment of the 8 studied novel CSFV strains and 12 reference isolates downloaded from NCBI. Most of the important mutations are highlighted in red boxes.

CONCLUSION

In conclusion, the present findings provide that genetic typing based on targeting UTRs of CSFV may be useful for evaluating methods to detect mutations in the virus population and characterize the strains involved in outbreaks. This could help to establish a national program

on prevention and control strategies to protect pigs against CSFV infection.

DECLARATIONS

Acknowledgments

Prof. John Morris (KMITL, Bangkok, Thailand) is thanked for his help in translating the manuscript and the authors extend their appreciation to Dalia Fouad for supporting this research through Researchers Supporting Project number (RSPD2024R965), King Saud University, Riyadh, Saudi Arabia.

Funding

This work was supported by the National Natural Science Foundation of China (Nos. 3157240 and 31402194), Natural Science Foundation of Heilongjiang Province of China (No. ZD201410), and the State Key Laboratory of Veterinary Biotechnology, Harbin Veterinary Research Institute, CAAS, China (no. SKLVBP2015013).

Data availability statement

The genomic sequences of the 5'- and 3'-UTR regions of 8 isolates have been deposited in the NCBI GenBank with accession numbers OQ150757-OQ150772.

Statement of conflicts of interest

The authors have declared no conflict of interest.

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