

# Sequencing and Insights into the Basal Phasi Charoen-like Virus Reveals Absence of Key Virulence Gene

Bredow Benjamin\*

Department of Microbiology, University of California Academy, San Francisco, USA

## Introduction

Advancements in genomics have revolutionized our understanding of viral diversity and evolution. In a significant breakthrough, the nearly complete genome of Phasi Charoen-like virus has been sequenced, shedding light on its genetic makeup and phylogenetic position within the Phlebovirus genus. PCLV represents a basal member of this genus, providing valuable insights into the evolutionary origins and diversity of phleboviruses. This article delves into the exciting findings, highlighting the genomic characterization of PCLV and its significance in the context of phlebovirus evolution. Phleboviruses belong to the family Phenuiviridae and are primarily transmitted by arthropods, such as mosquitoes, ticks, and sandflies.

These viruses can cause a range of diseases in humans and animals, including febrile illnesses, encephalitis, and hemorrhagic fevers. Understanding the genetic diversity and evolutionary relationships within the Phlebovirus genus is crucial for comprehending their pathogenic potential and developing effective control strategies. Researchers undertook the challenging task of sequencing the nearly complete genome of PCLV, a novel member of the Phlebovirus genus. Utilizing state-of-the-art genomic techniques, the researchers deciphered the genetic code of PCLV, enabling detailed analysis of its genomic structure, organization, and evolutionary relationships [1].

## Description

The genome sequencing revealed an intriguing aspect of PCLV—it occupies a phylogenetically basal position within the Phlebovirus genus. This finding suggests that PCLV represents an ancestral lineage that diverged early in the evolution of phleboviruses. This basal positioning provides valuable insights into the origins and evolutionary history of this diverse group of arthropod-borne viruses. Detailed analysis of the PCLV genome allowed researchers to identify unique genomic features and variations that distinguish it from other phleboviruses. These distinctive genetic characteristics provide clues about the molecular mechanisms underlying viral replication, pathogenesis, and host interactions. Understanding these features contributes to our knowledge of phlebovirus biology and may facilitate the development of targeted antiviral strategies [2].

The identification of PCLV as a basal member of the Phlebovirus genus has important evolutionary implications. By unraveling the genetic diversity and relationships within this group, researchers can trace the evolutionary trajectory of phleboviruses and gain insights into their adaptation to different hosts and

environments. Future research efforts may focus on exploring the ecological factors driving the diversification of phleboviruses and their potential impact on public health. The successful sequencing of the nearly complete genome of PCLV represents a significant milestone in the study of phleboviruses. This accomplishment provides a foundation for further research into the genetic diversity, evolution, and pathogenesis of this important group of arthropod-borne viruses. By unraveling the complexities of phlebovirus diversity, we move closer to developing comprehensive strategies for prevention, control, and treatment of diseases associated with these viruses. The sequencing of the nearly complete genome of Phasi Charoen-like virus has unraveled its genetic makeup and revealed its basal position within the Phlebovirus genus. This remarkable achievement contributes to our understanding of phlebovirus evolution, highlighting the diverse nature of this group of arthropod-borne viruses. [3].

The genomic characterization of PCLV opens avenues for further research into viral biology, host interactions, and evolutionary dynamics within the Phlebovirus genus. Ultimately, this knowledge enhances our ability to combat phlebovirus-related diseases and mitigate their impact on human and animal health. Virulence genes play a critical role in the pathogenicity and disease progression of viruses. In an intriguing discovery, Phasi Charoen-like virus a member of the Phlebovirus genus, has been found to lack a key virulence gene that is present in all other known Phleboviruses. This finding offers valuable insights into the genetic diversity and evolution of phleboviruses and their ability to cause disease. Additionally, the study highlights the effectiveness of metagenomic shotgun sequencing as a powerful tool for characterizing viruses in field-caught samples. This article explores the implications of PCLV's gene absence and the significance of metagenomic shotgun sequencing in understanding viral diversity [4].

Virulence genes are critical components that contribute to the ability of viruses to infect, replicate, and cause disease in their host organisms. Phleboviruses, a group of arthropod-borne viruses, possess various virulence factors that influence their pathogenic potential and host interactions. Investigating the presence or absence of specific virulence genes provides crucial insights into the mechanisms underlying virus-host interactions and the evolution of pathogenicity. In the study of PCLV, researchers made a remarkable discovery—the absence of a key virulence gene that is typically present in all other known Phleboviruses. This finding suggests a distinctive genomic profile for PCLV and raises intriguing questions about its pathogenic potential and its interactions with host organisms.

The absence of a key virulence gene in PCLV has significant evolutionary implications for the Phlebovirus genus. It suggests the existence of diverse evolutionary paths and genetic adaptations within this group of viruses. Understanding the genetic diversity and variation in virulence factors is crucial for comprehending the emergence, spread, and adaptation of phleboviruses in different ecological niches. Metagenomic shotgun sequencing has emerged as a powerful tool for characterizing viral diversity in field-caught samples. This approach enables the simultaneous identification and analysis of multiple viruses, including those that may be previously unknown or uncharacterized. By providing a comprehensive snapshot of the viral community present in a sample, metagenomic shotgun sequencing allows researchers to explore the genetic landscape and uncover novel viral species and genetic variations.

Metagenomic shotgun sequencing offers several advantages in the study of viral diversity. It does not require prior knowledge of the specific viruses

\*Address for Correspondence: Bredow Benjamin, Department of Microbiology, University of California Academy, San Francisco, USA, E-mail: bredowbenjamin@gmail.com

Copyright: © 2023 Benjamin B. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Received: 29 May, 2023, Manuscript No. Jgge-23-106413; Editor Assigned: 01 June, 2023, PreQC No. P-106413; Reviewed: 17 June, 2023, QC No. Q-106413; Revised: 22 June, 2023, Manuscript No. R-106413; Published: 29 June, 2023, DOI: 10.37421/2684-4567.2023.7.73

present in a sample, making it particularly useful for uncovering unknown or emerging viruses. Additionally, this approach provides a high-resolution view of viral genomes, allowing for the identification of genomic features, including the absence or presence of critical virulence genes, as observed in the case of PCLV. The discovery of PCLV's lack of a key virulence gene showcases the importance of studying viral diversity and understanding the genetic variations that influence pathogenicity. Such insights are crucial for effective disease surveillance, risk assessment, and the development of targeted control strategies. Metagenomic shotgun sequencing can be applied in various settings, including monitoring zoonotic diseases, investigating outbreaks, and identifying potential emerging pathogens [5].

---

## Conclusion

The absence of a key virulence gene in Phasi Charoen-like virus challenges our understanding of phlebovirus pathogenicity and highlights the genetic diversity within this viral group. This discovery underscores the significance of studying virulence factors and their impact on host interactions. Moreover, the effectiveness of metagenomic shotgun sequencing in characterizing viral diversity emphasizes its potential in uncovering novel viruses and understanding their ecological roles. By expanding our knowledge of viral diversity, we enhance our ability to detect, monitor, and control viral diseases, ultimately advancing public health efforts.

---

## Acknowledgement

None.

---

## Conflict of Interest

None.

---

## References

1. Chandler, James Angus, Panpim Thongsripong, Amy Green and Pattaporn Kittayapong, et al. "Metagenomic shotgun sequencing of a Bunyavirus in wild-caught *Aedes aegypti* from Thailand informs the evolutionary and genomic history of the Phleboviruses." *Virology* 464 (2014): 312-319.
2. Hall, Roy A., Helle Bielefeldt-Ohmann, Breeanna J. McLean and Caitlin A. O'Brien, et al. "Commensal viruses of mosquitoes: Host restriction, transmission, and interaction with arboviral pathogens." *Evolutionary Bioinformatics* 12 (2016): EBO-S40740.
3. Ferreira, Qesya Rodrigues, Fabian Felipe Bueno Lemos, Matheus Nascimento Moura and Jéssica Oliveira de Souza Nascimento, et al. "Role of the microbiome in *Aedes* spp. vector competence: What do we know?." *Virus* 15 (2023): 779.
4. Hermanns, Kyra. "RNA virus diversity in tropical mosquitoes and effects of virus interactions in vectors and hosts." PhD diss 2020.
5. Cottis, Solène, Adrien A. Blisnick, Anna-Bella Failloux and Kenneth D. Vernick. "Determinants of chikungunya and o'nyong-nyong virus specificity for infection of *Aedes* and *Anopheles* mosquito vectors." *Virus* 15 (2023): 589.

**How to cite this article:** Benjamin, Bredow. "Sequencing and Insights into the Basal Phasi Charoen-like Virus Reveals Absence of Key Virulence Gene." *J Genet Genom* 7 (2023): 73.