

# Identification of a Genotype IX Newcastle Disease Virus in a Guangxi White Duck

Zhixun Xie, Liji Xie, Zongli Xu, Jiabo Liu, Yaoshan Pang, Xianwen Deng, Zhiqin Xie, Qing Fan, Sisi Luo

Guangxi Key Laboratory of Animal Vaccines and Diagnostics, Guangxi Veterinary Research Institute, Nanning, Guangxi Province, China

**We report the complete genomic sequence of a novel Newcastle disease virus (NDV) strain, duck/China/Guangxi19/2011, isolated from a white duck in Guangxi Province, southern China. Phylogenetic analysis based on a fusion gene comparison with different NDV strains revealed that duck/China/Guangxi19/2011 is phylogenetically close to genotype IX NDV, and the deduced amino acid sequence of the fusion protein cleavage site was 112R-R-Q-R-R-F117. The whole nucleotide sequence had the highest homology (99.7%) to the sequence of strain F48E8 (GenBank accession number FJ436302). This study will help us understand the epidemiology and molecular characteristics of genotype IX Newcastle disease virus in ducks.**

Received 16 September 2013 Accepted 18 September 2013 Published 10 October 2013

Citation Xie Z, Xie L, Xu Z, Liu J, Pang Y, Deng X, Xie Z, Fan Q, Luo S. 2013. Identification of a genotype IX Newcastle disease virus in a Guangxi white duck. *Genome Announc.* 1(5):e00836-13. doi:10.1128/genomeA.00836-13.

Copyright © 2013 Xie et al. This is an open-access article distributed under the terms of the [Creative Commons Attribution 3.0 Unported license](https://creativecommons.org/licenses/by/3.0/).

Address correspondence to Zhixun Xie, xiezixun@126.com.

Newcastle disease virus (NDV) is a single-stranded, negative-strand RNA virus. The full genome of NDV has three sequence types, with lengths of 15,198 nucleotides, 15,192 nucleotides, and 15,186 nucleotides. NDVs of 15,198 nucleotides belong to class I (containing 9 genotypes), and NDVs of 15,192 nucleotides and 15,186 nucleotides belong to class II (containing 11 genotypes) (1–3). The NDV genome sequence contains six open reading frames, which encode 6 kinds of proteins (3'-NP-P-M-F-HN-L-5'), nucleocapsid proteins, phosphoproteins, matrix proteins, fusion proteins, hemagglutinin-neuraminidase proteins, and large polymerases (4–6).

In December 2011, NDV was isolated from a white duck in Guangxi Province, southern China. The isolate was named duck/China/Guangxi19/2011. Nucleotide sequences of duck/China/Guangxi19/2011 were amplified by PCR. The amplified products were purified and cloned into the pMD18-T vector (TaKaRa) and then sequenced (TaKaRa, Dalian, China). Sequences were assembled and manually edited to generate the final genome sequence.

Sequence analysis showed that the full genome sequence of duck/China/Guangxi19/2011 is 15,192 nucleotides and has the highest homology (99.7%) to the sequence of strain F48E8 (GenBank accession number FJ436302, class II, genotype IX). The amino acid sequence identities of the NP, P, M, F, HN, and L proteins between duck/China/Guangxi19/2011 and F48E8 are 99.6%, 99.0%, 98.6%, 99.3%, 99.1%, and 99.6%, respectively. The amino acid sequence identities of the NP, P, M, F, HN, and L proteins between duck/China/Guangxi/2011 and strain LaSota (GenBank accession number AF077761, class II, genotype II) are 91.5%, 86.5%, 90.2%, 91.9%, 90.6%, and 93.4%, respectively. The amino acid sequence identities of the NP, P, M, F, HN, and L proteins between duck/China/Guangxi/2011 and the Newcastle disease virus isolate SDWF02 (GenBank accession number HM188399, class II, genotype VII) are 94.5%, 80.8%, 89.2%, 90.3%, 88.3%, and 93.6%, respectively.

The sequence at the fusion protein cleavage site is a major

determinant of NDV pathogenicity (7–9). The F gene of duck/China/Guangxi19/2011 has the highest sequence homology (99.8%) to strain F48E8, and its virulence fusion protein cleavage site sequence (112R-R-Q-R-R-F117) (10) is in accord with the detected biological characteristics (mean death time, 51.4 h; intracerebral pathogenicity index, 1.80; intravenous pathogenicity index, 2.82).

The first to 21st amino acid sites of the F protein are the signal peptide areas of the N terminus and comprise one of the main variant areas of the F protein (11). There are four amino acid mutations in duck/China/Guangxi19/2011 compared with strain F48E8. The sites of amino acid mutations are the third amino acid, for which proline (hydrophobic) in F48E8 is mutated to serine (hydrophilic) in duck/China/Guangxi19/2011; the fourth amino acid, for which lysine (alkaline) in F48E8 is mutated to arginine (basic) in duck/China/Guangxi19/2011; the 380th amino acid, for which threonine (hydrophilic) is mutated to alanine (hydrophobic) in duck/China/Guangxi19/2011; and the 553rd amino acid, for which methionine (hydrophobic) is mutated to isoleucine (hydrophobic) in duck/China/Guangxi19/2011. Further study is needed to determine whether these variations affect viral fusion.

This report of the phylogenetic analysis of the whole-genome sequence of genotype IX NDV isolated from a white duck will further understanding of the epidemiology and molecular characteristics of NDV in duck.

**Nucleotide sequence accession number.** The GenBank accession number for duck/China/Guangxi19/2011 is [KC920893](https://www.ncbi.nlm.nih.gov/nuccore/KC920893).

## ACKNOWLEDGMENTS

This work was supported by the Guangxi Science and Technology Bureau (10100014-5 and 1222003-2-4) and by the Guangxi Government Senior Scientist Foundation (2011B020).

## REFERENCES

1. Knipe DM, Lamb R, Parks G. 2007. Paramyxoviridae: the viruses and their replication, p 1449–1496. In Knipe DM, Howley PM (ed), *Fields virology*, 5th ed. Lippincott Williams & Wilkins, Philadelphia, PA.

2. Huang Y, Wan HQ, Liu HQ, Wu YT, Liu XF. 2004. Genomic sequence of an isolate of Newcastle disease virus isolated from an outbreak in geese: a novel six nucleotide insertion in the non-coding region of the nucleoprotein gene. *Arch. Virol.* **149**:1445–1457.
3. Czegledi A, Ujvari D, Samogyi E, Wehmanna E, Werner O, Lomniczi B. 2006. Third genome size category of avian paramyxovirus serotype 1 (Newcastle disease virus) and evolutionary implications. *Virus Res.* **120**: 36–48.
4. Maminaiina OF, Gil P, Briand FX, Albina E, Keita D, Andriamanivo HR, Chevalier V, Lancelot R, Martinez D, Rakotondravao R, Rajaonari-son JJ, Koko M, Andriantsimahavandy AA, Jestin V, Servan de Almeida R. 2010. Newcastle disease virus in Madagascar: identification of an original genotype possibly deriving from a died out ancestor of genotype IV. *PLoS One* **5**:e13987. doi:[10.1371/journal.pone.0013987](https://doi.org/10.1371/journal.pone.0013987).
5. Rima BK, Wishaupt RG, Welsh MJ, Earle JA. 1995. The evolution of morbilliviruses: a comparison of nucleocapsid gene sequences including a porpoise morbillivirus. *Vet. Microbiol.* **44**:127–134. doi:[10.1016/0378-1135\(95\)00005-U](https://doi.org/10.1016/0378-1135(95)00005-U).
6. García-Sastre A, Cabezas JA, Villar E. 1989. Proteins of Newcastle disease virus envelope: interaction between the outer hemagglutinin-neuraminidase glycoprotein and the inner non-glycosylated matrix protein. *Biochim. Biophys. Acta* **999**:171–175.
7. Panda A, Huang Z, Elankumaran S, Rockemann DD, Samal SK. 2004. Role of fusion protein cleavage site in the virulence of Newcastle disease virus. *Microb. Pathog.* **36**:1–10.
8. Xie Z, Xie L, Chen A, Liu J, Pang Y, Deng X, Xie Z, Fan Q. 2013. Complete genome sequence analysis of a Newcastle disease virus isolated from a wild egret. *J. Virol.* **86**:13854.
9. de Leeuw OS, Koch G, Hartog L, Ravenshorst N, Peeters BP. 2005. Virulence of Newcastle disease virus is determined by the cleavage site of the fusion protein and by both the stem region and globular head of the haemagglutinin-neuraminidase protein. *J. Gen. Virol.* **86**:1759–1769.
10. Manin TB, Shcherbakova LO, Bochkov IA, El'nikov VV, Pchelkina IP, Starov SK, Drygin VV. 2002. Characteristics of field isolates of Newcastle disease virus isolated in the course of outbreaks in the poultry plant in the Leningrad region in 2000. *Vopr. Virusol.* **47**:41–43. (In Russian.)
11. Collins MS, Strong I, Alexander DJ. 1994. Evaluation of the molecular basis of pathogenicity of the variant Newcastle disease viruses termed “pigeon PM V-1 viruses.” *Arch. Virol.* **134**:403–411.