

Isolation and molecular characterization of virulent Newcastle disease viruses in Mali in 2007 and 2008

Patricia Gil, Renata Servan de Almeida, Saliha Hammoumi, Sophie Molia, Véronique Chevalier, H. A. Traore, K. Samaké, Emmanuel Albina

▶ To cite this version:

Patricia Gil, Renata Servan de Almeida, Saliha Hammoumi, Sophie Molia, Véronique Chevalier, et al.. Isolation and molecular characterization of virulent Newcastle disease viruses in Mali in 2007 and 2008. EPIZONE 3rd annual meeting, May 2009, Antalya, Turkey. 1 p., 2009. hal-02819510

HAL Id: hal-02819510 https://hal.inrae.fr/hal-02819510v1

Submitted on 6 Jun 2020

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

Isolation and molecular characterization of virulent Newcastle disease viruses in Mali in 2007 and 2008

P.Gil¹, R.Servan¹, S.Hammoumi¹, S.Molia², V.Chevalier², H.A.Traoré³, K.Samake³, E.Albina¹

1, CIRAD, UMR «Contrôle des maladies», F-34398 Montpellier, France

2, CIRAD, UPR «Agirs», F-34398 Montpellier, France



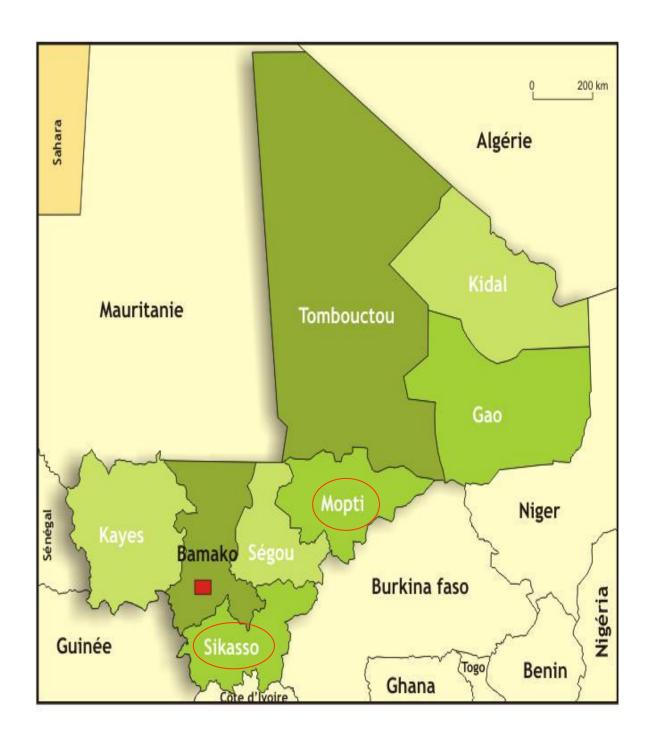


INTRODUCTION

ewcastle disease virus (NDV) is the causal agent of a fatal respiratory and neurological disease that can result in 100% morbidity and mortality in chicken flocks. ND was in the former list A diseases of the World Organisation for Animal Health (OIE) and is still one of the most serious animal diseases in developing countries.

Although ND is endemic in Africa, little is known about the NDV strains circulating in this continent. The objective of this study was to detect, isolate and characterise the NDV strains circulating in Mali.

In this study, cloacal and tracheal swabs were collected on healthy chickens in two different regions in Mali in a framework of a surveillance programme implemented in 2007 and 2008. These samples were analysed by RT-PCR. Positive samples were inoculated into eggs and isolated viruses were sequenced.



MATERIAL AND METHODS

NA extraction was performed using the Nucleospin Virus Kit (Macherey Nagel) on an automatic workstation (Beckman FXP), allowing the extraction of 90 samples in 50 minutes.

Virus RNA detection was done by a one-step RT-PCR targeting the F gene of NDV. All 1076 samples from domestic birds and all 1041 samples from wild birds were analysed and the positive ones were further inoculated into the allantoic cavities of 9 day-old to 11 day-old embryonated eggs. The complete procedure is represented in fig 1.

Nucleotide sequencing was achieved on 1659 pb of the F gene and on the full sequence of the HN gene. Phylogenetic analyses were carried out using the neighbor-joining method.

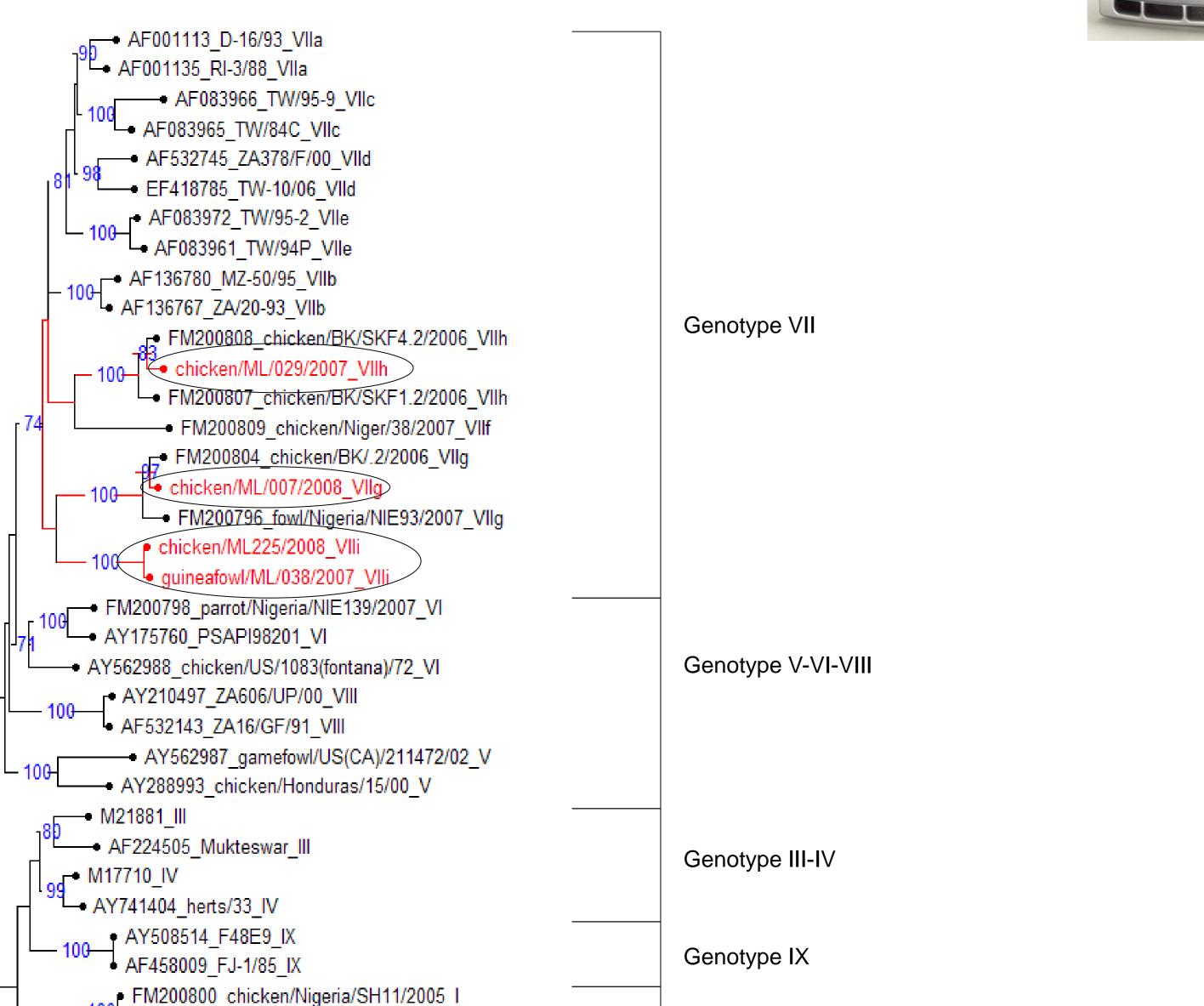


Fig2: Phylogenetic analysis of partial *F* sequences based on nucleotides 31-377. The tree was constructed using neighbor-joining method with 1000 bootstrap replicates. Only bootstrap values higher than 70 are shown. Malian NDV sequences are in red.

Genotype I-II

AY175734_HNZDK98194_classl

AY135759_HIECK87191_classl

90 samples dispensed Sample in a 96 deep well plate 4 ten-fold dilutions of NDV laSota Discard swabs **RNA** extraction from 100 µl in 50 min Q RT-PCR : F gene **Elution** in a final - Primers F+4839, F2AS (Wise et al, 2004-Pham et al 2005) volume of 50 μL - Sybrgreen QRT-PCR one-step Master mix kit (Stratagene) Collection Positive samples swabs Pathotyping of the cleavage site and Inoculation on embryonated eggs Fig1: Procedure for analysis at CIRAD laboratory

RESULTS AND DISCUSSION

DV prevalence was 1.7% for domestic birds and 2.3% for wild birds. Four viruses were isolated from domestic birds after two passages into embryonated eggs. Unfortunately no isolates from wild birds could be obtained. The sequence of the cleavage site shows the presence of basic amino acids and thus characterise these isolates as virulent strains.

Two motifs were identified for the cleavage sites were identified with at least three basic amino acids (112-RRRKR*FV-118 and 112-RRQKR*FI-118). The presence of V₁₁₈ associated with the cleavage site RRRKR*FV in two of these isolates was only reported recently, in the neighbour country Burkina Faso (1).

Phylogenetic analysis based on a fragment of 356 nucleotides corresponding to position 48 to 422 of the F gene showed that these isolates are branched with the genotype VII (fig 2). This genotype represents the currently circulating genotype in Europe, Asia and Africa (2-3). Phylogenetic analysis performed on the full sequence of the F and HN genes confirmed that the Malian NDV isolates belong to the genotype VII. Two of the isolates clustered in the two subgenotypes (VIIg and VIIh) recently described by Snoeck et al in Burkina Faso, Niger and Nigeria (1). This suggest a circulation of the strains through animal trade between these countries. The two other isolates may define a new subgenotype (VIII, fig 2).

These results show that the two survey regions were good sites for NDV ecology since they are major trade crossroads for domestic birds of different neighboring countries.

CONCLUSION

AF532741 ZA341/P/99_I

FM200801_chicken/Nigeria/N2/2006_ll

his study reports on original NDV isolates detected in Mali during a surveillance campaign in domestic and wild birds over 2007 and 2008. Four strains, genetically characterise as virulent may define new subgenotypes. These virulent strains were surprisingly isolated from unvaccinated and apparently health poultry. Question about the real virulence of these strains warrants further experimental demonstration.

References: 1- Snoeck CJ, Ducatez MF, Owoade AA, Faleke OO et al , Newcastle disease virus in West Africa: new virulent strains identified in non-commercial farms2009. Archives of Virology (2009)-154

2- Lomniczi B, Wehmann E et al: Newcastle disease outbreaks in recent years in Western Europe caused by an old (VI) and a novel genotype (VII). Archives of Virology (1998) -146

3- Herczeg J, Wehmann E et al: Two novel genotype groups (VIIb and VIII) responsible for recent Newcastle disease outbreks in Southern Africa, one of which reached Southern Europe. Archives of Virology (1999) 144



de coopération internationale en recherche agronomique pour le développement



Patricia.gil@cirad.fr