Identification of canine papillomavirus type 1 (CPV1) DNA in dogs with cutaneous papillomatosis

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ABSTRACT.- De Alcântara B.K., Alfieri A.A., Rodrigues W.B., Otonel R.A.A., Lunardi M., Headley S.A. & Alfieri A.F. 2014. Identification of canine papillomavirus type 1 (CPV1) DNA in dogs with cutaneous papillomatosis. Pesquisa Veterinária Brasileira 34(12):1223-1226. Laboratório de Virologia Animal, Departamento de Medicina Veterinária Preventiva, Universidade Estadual de Londrina, Rodovia Celso Garcia Cid, Campus Universitário, Cx. Postal 10011, Londrina. PR 86057-970, Brazil. E-mail: alfieri@uel.br

Canine oral papillomavirus (COPV), also known as Canine Papillomavirus type 1 (CPV1), induces papillomas at the mucous membranes of the oral cavity and at the haired skin of dogs. The classification of Papillomavirus (PV) types is based on the L1 capsid protein and nucleotide sequence; so far, 14 CPV types have been described in several countries, but the molecular characterization of CPV in Brazil is lacking. This study investigated the presence of the PV in seven papillomas from four mixed breed dogs from Londrina/PR, Southern Brazil, by partial sequencing of the L1 gene. Seven exophytic cutaneous lesions were surgically removed and processed for histopathological and molecular characterization. Histopathology confirmed the lesions as viral papillomas due to typical histological features. Polymerase Chain Reaction (PCR) assay using the FAP59 and FAP64 primers targeted the L1 gene followed by sequence analysis of the amplicons identified CPV1 in all evaluated papilloma samples. This study represents the first description of CPV1 DNA associated with canine papillomatosis in Brazil.

INDEX TERMS: Papillomatosis, canine, CPV1, papilloma, warts, L1 gene.

INTRODUCTION

The Papillomavirus (PV) is a group of DNA viruses that induces benign and malignant proliferative lesions at the epidermis and mucous membranes of their natural hosts, and probably occur in most mammals and birds (De Villiers et al. 2004).
The classification of the papillomavirus types is based on the L1 nucleotide sequence. The L1 gene is highly conserved, and a new putative PV type is considered when the L1 nucleotide sequence is at least 10% different from other PV types (De Villers et al. 2004). Based on this classification, more than 200 types of PVs have been characterized, most being isolated from humans (Bernard et al. 2010); currently 14 types of canine papillomavirus (CPVs) have been identified (Yuan et al. 2012a, Lange et al. 2013).

Moreover, CPVs can be allocated into three distinct Papillomavirus genera: *Lambdapapillomavirus* (CPVs 1 and 6); *Taupapillomavirus* (CPVs 2, 7, and 13); and *Chipapillomavirus* (CPVs 3, 4, 5, 8, 9, 10, 11, and 14), while CPV 12 is not currently classified into a definite genus (Bernard et al. 2010, Lange & Favrot 2011, Lange et al. 2012).

CPV1, originally denominated canine oral papillomavirus (COPV) (Bernard et al. 2010), induces papillomas at the mucous membranes of the oral cavity and at the haired skin of dogs (Gross et al. 2005); papillomas may progress to squamous cell carcinoma (Teifke et al. 1998). Additionally, CPV1 has been associated with inverted papillomas (Lange et al. 2009), and conjunctival epithelial hyperplastic lesions (Brandes et al. 2009) in dogs.

CPV2 has already been isolated from exophytic and endophytic papillomas as well as invasive squamous cell carcinoma, CPV6 was isolated from endophytic papilloma, and CPV7 from an exophytic lesion (Lange & Favrot 2011). CPVs 3, 4, 5, 8, 9, 10, 11, and 14 are associated with the development of pigmented plaques (Lange et al. 2013). Additionally, CPV13 was identified in oral papillomas (Lange et al. 2012). Moreover, the histopathological characterization of canine viral papilloma is obtained by the finding of typical viral-induced alterations in the cytoplasm and/or nucleus of affected cells (Gross et al. 2005, Lange & Favrot 2011). The role of CPV12 as a pathogen of papillomatosis has not been elucidated (Yuan et al. 2012b unpublished data).

Isegawa et al. (1994) revealed that the CPV1 L1 gene was more closely related with human papillomavirus type 1 (HPV1A) and animal PVs associated with cutaneous lesions in the rabbit, European elk, deer, and cattle. Consequently, CPV1 has been used as a model for animal and human PVs infection and vaccine trials (Campo 2002).

Although oral papillomas were described in dogs from Brazil (Souza et al. 2006, Bianchi et al. 2012), molecular studies that characterized the PVs types in canine papillomas from Brazil were not located. This study investigated the occurrence of the CPV DNA in papillomas of dogs from Brazil by using the partial L1 gene amplification and sequence analysis.

**MATERIALS AND METHODS**

**Dogs, cutaneous lesions, and sampling.** Cutaneous lesions, clinically classified as papillomas, were surgically removed and collected from four mixed breed dogs. The lesions were surgically removed at the veterinary hospital, according with the protocols of animal welfare. All tissue samples were divided into two sections; one fixed in 10% neutral buffered formalin solution and routinely processed for histopathological evaluation, and the other maintained at -20°C until used for molecular diagnosis.

**DNA extraction, PCR, and sequencing.** DNA was extracted from all samples by using the Qiagen DNeasy tissue kit (Qiagen Sample and Assay Technologies, Hilden, Germany). A negative control without tissue was included in all procedure to monitor possible contamination.

The PCR assay was done by using primers FAP59 and FAP64 designed to amplify a 480 base pair (bp) fragment of the PV L1 gene (Forslund et al. 1999), with modifications (Claus et al. 2007). All PCR products were analyzed by electrophoresis in 2% agarose gel, stained with ethidium bromide, and visualized under UV light.

The PCR products were purified by using the Illustra GFX PCR DNA and Gel Band Purification kit (GE Healthcare, Little Chalfont, UK). Direct sequencing was then performed using the DYEnamic ET dye terminator cycle sequencing kit (GE Healthcare, Little Chalfont, UK) with forward and reverse primers, in the 3500 Genetic Analyzer (Applied Biosystems, Carlsbad, USA). The obtained sequences were examined with the PHRED software for quality analysis by chromatogram readings. The sequences were accepted if the base quality was equal to or higher than 20. Consensus sequences were obtained by using the CAP3 software, and the sequence identity was compared with similar sequences deposited in the GenBank by using the BLAST software. The guidelines of the Papillomavirus Nomenclature Committee 1995 (14th International Papillomavirus Conference, Quebec City, Quebec, Canada) were used to identify PV types (De Villiers et al. 2004).

A multiple alignment was performed by using CLUSTAL W (version 1.4) within the MEGA 5.1 software; a sequence identity matrix was generated by using BioEdit software version 7.0.8.0. Phylogenetic trees were reconstructed by the maximum-likelihood method and distances were estimated by the Kimura two parameter (Kimura 1980), using MEGA 5.1 software. Statistical analyses of phylogenetic trees were determined by 1,000 bootstrap replicates.

**RESULTS**

Pathological findings. Warts were observed at the lips and nose of two dogs, at the mucous membranes of the oral cavity of one, and at the lips and the left and right hind limbs of the other. In one dog, the warts resulted in deformation of the nasal cavity (Fig.1). Grossly, all collected tissues were exophytic cutaneous warts. The histopathological findings of all submitted tissue sections were similar, being charac-
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There was hyperkeratotic hyperplasia with the forming of finger-like projections of the hyperplastic squamous epithelium, the occurrence of giant keratohyalin granules within the stratum granulosum, and discrete koilocytosis of the stratum spinosum and granulosum (Fig. 2); distinct basophilic intranuclear inclusion bodies were not observed.

**Molecular characterization of canine papillomavirus.** Amplicons of the expected length (approximately 480 bp), were amplified from all samples by the PCR assay; negative controls yielded no amplified products. Six of these sequences were similar with 100% identity among them; however, one nucleotide sequence from an oral papilloma had 99.7% identity with the other sequences from this study. The nucleotide sequences obtained during this study demonstrated 99.3% to 100% similarity with other sequences of CPV1 available at GenBank.

The strains identified in this study and used during the sequence alignment were designated as CPV1 BR/UEL1 and CPV1 BR/UEL2 (GenBank accession numbers: KF199909 and KF199910). Analysis of the phylogenetic tree demonstrated that the Brazilian strains of CPV1 clustered with known strains of the *Lambdapapillomavirus*, particularly with those of CPV1 (Fig. 3).

The GenBank accession numbers of the strains used are: BPV1 (X02346), CPV1 (L22695), CPV2 (AY722648), CPV3 (NC_008297), CPV4 (NC_010226), CPV5 (FJ492743), CPV6 (FJ592744), CPV7 (FJ492743), CPV8 (HM796884), CPV9 (NC_016074), CPV10 (NC_016075), CPV11 (NC_016076), CPV12 (JQ754321), CPV13 (JX141478), CPV14 (JQ701802), FdPV1 (NC_004765), and FdPV2 (EU796884).

**DISCUSSION**

The gross and histopathological features of the cutaneous lesions are consistent with those described in canine oral papillomas (Gross et al. 2005, Ginn et al. 2007), since the cytoplasmic alterations observed during this study are typical of viral-induced cytopathic effects and are diagnostic for this lesion (Gross et al. 2005, Lange & Favrot 2011).

Similar histopathological findings were recently described in dogs with oral papillomavirus from Korea identified by molecular biology (Yhee et al. 2010) and in dogs from Brazil characterized by immunohistochemistry for *Papillomavirus* (Bianchi et al. 2012), however the last investigation did not specify the PV type identified in dogs from Brazil.

Confirmation of the participation of CPV1 within these lesions was obtained due to the successful amplification of viral DNA with subsequent sequencing of the partial L1 gene of CPV1; similar results have been described (Yhee et al. 2010). However, it must be highlighted that the degenerate primers (FAP 59 and FAP 64), originally designed to detect a broad range of cutaneous HPVs (Forslund et al. 1999), originally designed to detect a broad range of cutaneous HPVs (Forslund et al. 1999), successfully amplified the partial fragment of the CPV1 L1 gene, suggesting that direct sequencing is an important tool for the molecular characterization of PV types of infectious agents.

Bianchi et al. (2012) using monoclonal *Papillomaviridae* antibodies, by immunohistochemistry, demonstrated that *Papillomavirus* was present in 83.3% (20/24) of papillomas from dogs in Rio Grande do Sul. However, to the best of the authors’ knowledge, the present study represents the first molecular characterization of CPV1 associated with papillomatosis in dogs from Brazil.

The importance of this study was to characterize the PVs types present in dogs of Brazil, since immunity is type-specific, and CPV1 L1 virus like-particles vaccines can protect dogs against the development of mucosal papillomas (Suzich et al. 1995, Stanley et al. 2001), while the immu-
nity can be passively transferred to offsprings (Nicholls & Stanley 2000). Additionally, considering the great diversity of canine PVs, the efficiency of degenerate primers, as demonstrated during this investigation, facilitate the identification of putative new PV types (Zaugg et al. 2005, Lange et al. 2013). Hence, the knowledge of the CPV types circulating in local canine populations is essential for the establishment of control measures.

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