Outbreaks of canid herpesvirus 1 disease in puppies in southern Brazil

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ABSTRACT.- Cargnelutti J.F., Masuda E.K., Neuls M.G., Weiblen R. & Flores E.F. 2015. Outbreaks of canid herpesvirus 1 disease in puppies in southern Brazil. PesquisaVeterinária Brasileira 35(6):557-561. Setor de Virologia, Departamento de Medicina Veterinária Preventiva, Universidade Federal de Santa Maria, Av. Roraima 1000, Santa Maria, RS 97105-900, Brazil. E-mail: eduardofurtadoflores@gmail.com

Canid herpesvirus 1 (CHV-1) is a widespread pathogen of dogs and produces infertility, abortions and severe systemic disease in young puppies. Clinical data indicate the circulation of CHV-1 among Brazilian dogs yet definitive diagnosis has rarely been accomplished. This article describes the clinicopathological findings of four independent cases/outbreaks of neonatal disease by CHV-1 in Bulldog puppies followed by virus identification and genetic characterization. Three events occurred in a kennel holding dogs of different breeds at reproductive age (March 2013, October 2013 and April 2014). Puppies from three French or English Bulldog litters, aging 9 to 30 days were affected, presenting dyspnea, agonic breathing, pale mucous, abdominal pain and tension, evolving to death within about 24 hours. At necropsy, the puppies presented necrohemorrhagic hepatitis, multifocal and moderate necrohemorrhagic nephritis and fibrinonecrotic interstitial pneumonia. Virus isolation was positive in clinical specimens from one litter and CHV-1 DNA was detected by PCR in tissues from all four cases. Virus-neutralizing assays with samples of the affected kennel revealed 9/12 adult animals with high antibody titers to CHV-1. Nucleotide sequencing of glycoprotein B, C and D genes revealed 99-100% of identity among the viruses and with CHV-1 sequences available in GenBank. Phylogenetic analyses of gC sequences showed a segregation of the samples, even among three isolates from the same kennel. These findings support CHV-1 infection as the cause of disease and death in these dog litters, reinforcing the need for correct etiologic diagnosis, prevention and immunization against CHV-1 in dogs from Southern Brazil.

INDEX-TERMS: Canid herpesvirus 1, CHV-1, diagnosis, dogs, glycoproteins and pathological findings.
Cases of disease clinically compatible with CHV-1 in dogs have been reported in Brazil, and outbreaks of neonatal deaths in French Bulldogs (Kurissio et al. 2012) and in English Bulldogs (Oliveira et al. 2009) have been described. In Brazil, CHV-1 has been isolated from tissues of neonatal deaths in different breeds, including the English Bulldog (Ronsse et al. 2004). The clinical signs of neonatal death in dogs with CHV-1 infection are characterized by respiratory distress, abdominal pain, and jaundice, with a high mortality rate (Ávila et al. 2011). As no vaccine is commercially available in Brazil, the diagnosis of CHV-1 infection is crucial for the implementation of measures to prevent and control outbreaks.

INTRODUCTION

Canid herpesvirus 1 (CHV-1) belongs to the family Herpesviridae, subfamily Alphaherpesvirinae, genus Varicellovirus (ICTV, 2015), and is associated with a variety of clinical manifestations in dogs, including neonatal deaths, respiratory, ocular, and reproductive disorders (Evermann et al. 2011). Like other alphaherpesviruses, CHV-1 establishes latent infections in neural and non-neural tissues after acute infection and can be reactivated in stressful and immunosuppressive situations (Miyoshi et al. 1999).

The outcome and clinical manifestations of CHV-1 infection in dogs are associated with several risk factors, including age, breeding, and kennel cough (Ronssë et al. 2004). Adult dogs infected with CHV-1 usually develop a mild respiratory disease, frequently associated with Parainfluenza virus and Bordetella bronchiseptica infection (Decaro et al. 2008, Evermann et al. 2011). The most important disorders associated with CHV-1 infection, however, generally occur in pregnant bitches and in young puppies. Systemic viremia in pregnant females may result in transplacental CHV-1 transmission and fetal infection, leading to fetal deaths, abortion, mummification, infertility, and the birth of weak puppies (Hashimoto et al. 1983, Anvik 1991, Decaro et al. 2008, Evermann et al. 2011). The age of the puppies at the time of primary infection is determinant to the outcome of the disease and the clinical signs are usually more severe in neonates (Decaro et al. 2008). Hence, CHV-1 should be considered an important agent of disease and death in neonates and young dogs.

Cases of disease clinically compatible with CHV-1 infection have been frequently reported by veterinarians in Brazil, yet a definitive diagnosis is rarely reached. Likewise, serological studies and/or virus identification have not been conducted to date, is spite of this clinical relevance. To date, only few reports were described in Brazil: a pathological description of neonatal death of two litters (Oliveira et al. 2009); a case of virus isolation and molecular identification of CHV-1 in a puppy with neonatal death (Kurissio et al. 2012) and a clinicopathological description of neonatal death in dogs with 10 days-old (Ávila et al. 2011). As no vaccine is commercially available in Brazil, CHV-1 infection of bitches and neonate puppies seem to occur endemically and, as such, have acquired relative importance for dog breeders and domestic dog owners in past years. This report describes four cases/outbreaks of CHV-1 infection in Bulldog puppies and molecular identification of the agent. We hope our report contribute for the understanding of CHV-1 biology and, most importantly, to call attention to and help veterinarians in recognizing such cases.

MATERIALS AND METHODS

Outbreak description

The four cases of CHV-1 infection occurred in Bulldog litters in Porto Alegre, Rio Grande do Sul. Three occurred from March to October 2013 (fall to spring) and the later occurred in April 2014. The main epidemiological and clinical findings are described below:

Case 1 (SV822/13). Two 9 to 15-day-old puppies out of a litter of eight English Bulldogs presented severe dyspnea, pale mucous, abdominal pain and tension, with a fast evolution to death (approx. 24 h). After birth, the bitch developed a severe mastitis that hampered breast-feeding.

Case 2 (SV 823/13). Four 22-day-old litter of English Bulldog puppies developed respiratory agony and pulmonary edema that evolved to death in a few hours.

Case 3 (SV1110/13). A 30-day-old puppy of a litter of French Bulldogs presented nasal discharge, abdominal pustules and neurological signs, characterized by ataxia and incoordination. The clinical course evolved to death in approximately 24 h.

Case 4 (SV242/14). Two 23-day-old puppies of a litter of six English Bulldogs developed abdominal pain and showed pale mucosa, with fast evolution to death (24 h). The bitch was seronegative to CHV-1 when introduced in the kennel, immediately before the pregnancy.

Pathologic analyses

Animals from all reported cases were submitted to necropsy and tissue samples (kidney, lung, and liver) were collected for histopathology and virology diagnosis. For histopathology, samples were fixed in 10% buffered formalin and processed for routine histopathological examination. The slides were stained using hematoxylin and eosin (H & E) standard protocol.

Virus isolation

For virus isolation, tissue fragments (kidney, lung and liver) were grinded, homogenized and resuspended to 10% (weight/volume) in culture medium. After low speed centrifugation, the supernatants were inoculated onto monolayers of primary canine kidney cells, stored at 32°C with 5% of CO₂, and monitored for cytopathic effect (cpe) during three passages of five days each before considering negative for virus.
Virus-neutralization

Thirteen serum samples collected from dogs and bitches of different breeds at reproductive age from the kennel (cases 1, 3 and 4) were submitted to a standard virus-neutralization (VN). VN assays were performed in 96-well plates, testing two-fold dilutions of sera against a fixed dose of virus (100-200 TCID₅₀/well) and MDCK cells as indicators. The CHV-1 used in VN assays was kindly provided by Dr. João Pessoa Araújo Júnior (Unesp, Pirassununga, SP). VN readings were performed after five days of incubation at 32 °C and VN titers were considered as the reciprocal of highest dilution of sera that prevented the production of cpe. Sera from a dog naturally infected with CHV-1 and from a seronegative dog were used as positive and negative controls, respectively.

Molecular analysis

Total DNA was extracted from tissue samples (pool of kidney, liver and lung) using DNAzol Reagent® and submitted to PCR for CHV-1 glycoprotein B gene (gB), using the primers and conditions previously described (Ronsse et al. 2005). DNA extracted from the liver of a dog infected with CHV-1 (SV435/08) (Oliveira et al. 2009) was used as positive control. Additionally, PCR to glycoproteins C and D (gC and gD; Table 1) were performed. The amplicons were purified using the PureLink PCR kit® according to the manufacturer’s instructions and submitted to nucleotide sequencing in an automatic sequencer ABI-PRISM 3100 Genetic Analyzer armed with 50 cm capillaries and POP6 polymer (Applied Biosystems, Carlsbad, CA, USA). The obtained sequences were analyzed by the Staden package (Staden 1996) and aligned using the BioEdit Sequence Alignment Editor Software suite, version 7.0.5.3 (Hall 1999) and compared with other CHV-1 sequences deposited in GenBank. To building the phylogenetic tree, based on sequences of gC gene, it was used the Neighbor-Joining method with 2000 bootstrap replicates implemented by MEGA5.0 (Tamura et al. 2011), based on Tamura 3- parameters model.

Table 1. Primers and PCR conditions used to amplified fragments of glycoproteins C and D of Canid Herpesvirus-1

<table>
<thead>
<tr>
<th>Gene</th>
<th>Primer sequence (5’-3’)</th>
<th>Nucleotide positions</th>
<th>Amplicon size (bp)</th>
<th>PCR conditions</th>
</tr>
</thead>
<tbody>
<tr>
<td>gC</td>
<td>F: CGGCATCTACATCCAAAC</td>
<td>Based on gC - AF361074</td>
<td>65-82</td>
<td>95°C - 5 min, 30x (95°C - 1min, 50°C - 1 min, 72°C - 1min), 72°C - 7 min</td>
</tr>
<tr>
<td></td>
<td>R: GACCGTCAAATCCACAC</td>
<td>832-815</td>
<td></td>
<td></td>
</tr>
<tr>
<td>gD</td>
<td>F: CTTCTACTGACCACTGTG</td>
<td>Based on gD - AF361076</td>
<td>3614-3631</td>
<td>95°C - 5 min, 30x (95°C - 1min, 47.5 °C - 1 min, 72°C - 1min), 72°C - 7 min</td>
</tr>
<tr>
<td></td>
<td>R: TGGGCTCAGGTAATTC</td>
<td>4260-4243</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

RESULTS AND DISCUSSION

At necropsy, all puppies had severe necrohemorrhagic disease, characterized by multifocal to aleatory petechia in liver and kidneys. In the lungs had severe edema and hemorrhage. At histopathologic examination, there were ne-

Fig.1. (A) Pulmonary lobes with severe edema and hemorrhage. (B) Aleatory foci of hemorrhage (petechia) over the renal cortex. (C) Multiple alveoli are necrotic and filled by fibrinous plaques and mild to moderate neutrophilic and histiocytic inflammatory infiltrate. (D) Severe necrosis of hepatocytes with Councilmann-like bodies and moderate hemorrhage. (C, D) Hematoxylin and eosin. Bar = 100.00μm.
In addition to the low body temperature, the lack of zonal temperature fluctuation is frequently reported during autumn and winter, when the environmental temperatures, housing conditions and the body temperature may be subclinical or mild in older dogs due to the ability of some animals to stabilize their body temperatures at approximately 35°C (Huxsoll & Hemelt 1970). The disease is particularly severe and frequently fatal in young puppies in which the optimal virus replication occurs at temperature below 35°C (Jenson & Ederstrom 1955).

The presumptive diagnosis of the four events was based on the clinical signs, high mortality and pathological findings. At necropsy, the macroscopic features were typical of neonatal CHV-1 infection, characterized by a multisystemic necrohemorrhagic disease (Decaro et al. 2008, Oliveira et al. 2009). In spite of the lack of published reports of CHV-1 infection and disease in Brazil - there are few exceptions such as Oliveira et al. (2009), Ávila et al. (2011) and Ku-rissio et al. (2012) - veterinarians frequently report cases compatible with CHV-1-induced disease. Virtually all suspected cases, however, remain without a definitive etiological (virology, serology, molecular) diagnosis. In these particular cases, however, the support of a virology diagnosis using CHV-1 sequences obtained from GenBank and Brazilian samples, such as SV453/08 - identified in a puppy infected with CHV-1 in Southern Brazil in 2008 (Oliveira et al. 2009) (Genbank accession - KJ676501).

Neonatal mortality in puppies is frequently reported during autumn and winter, when the environmental temperatures fluctuate from 0 to +10-15°C. In addition to environmental temperatures, housing conditions and the body temperature exert a strong effect on CHV-1 infection since optimal virus replication occurs at temperature below 35°C (Huxsoll & Hemelt 1970). The disease is particularly severe and frequently fatal in young puppies in which the body temperatures are usually below 35-36°C. The infection may be subclinical or mild in older dogs due to the ability of these animals to stabilize their body temperatures at 38°C (Jenson & Ederstrom 1955).

In addition to the low body temperature, the lack of maternal immunity may worsen neonatal CHV-1 disease
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(Evermann et al. 2011). Maternal antibodies apparently protect puppies from fatal CHV-1 infection even though the animals develop subclinical infection (Decaro et al. 2008, Huxssoll & Hemelt 1970). Thus, in a population of dogs in which the infection is endemic or can be frequently introduced, vaccination of females at heat and/or during pregnancy should be highly recommended to minimize the consequences of CHV-1 infection of puppies.

In the present report, the affected puppies presented severe respiratory signs that evolved to death in a day or so. Interestingly, a puppy from a French Bulldog litter (Case 3) also presented neurological signs, which are rarely observed in puppies naturally infected with CHV-1. For this reason, differential diagnosis for canine distemper virus (CDV), an agent involved in neurologic disease in dogs was performed (Tipold et al. 1992). Again, the pathological and virological findings were conclusive of CHV-1 infection and histopathological analyses dismissed CDV infection.

Sequence analysis of CHV-1 gC sequences (including those from the reported cases and two obtained in GenBank) revealed a high level of nucleotide identity. Nonetheless, SV822/13, SV242/14 and SV1110/13 isolates – all coming from the same kennel in different occasions – clustered in different branches. Two scenarios may be envisioned: i. the most likely hypothesis is that the viruses involved in these events have the same origin. The virus would remain latent in the dogs of the kennel and would be periodically reactivated and transmitted. The selective pressure upon the virus during the transmission-infection cycle would explain the few observed mutations. or ii. these viruses have different origins and have been introduced in the kennel population in different occasions. On the other hand, the closely relationship of gC among CHV-1 samples from different Brazilian and world regions (SV822/13, SV435/08 and isolate Milou) confirms that gC is highly conserved and, therefore, is suitable for CHV-1 diagnosis and phylogenetic studies.

CONCLUSIONS

Veterinary clinicians frequently describe clinical cases compatible with CHV-1-induced disease; but etiologic confirmation has been accomplished in rare occasions.

In this sense, this report contributes for the identification and awareness of a common cause of neonatal deaths in puppies in Southern Brazil, and indicates the need for the adoption of control measures, including vaccination.

Sources and manufacturers.- Life Technologies, CA, USA.

Declaration of conflicting interests.- The authors declared that they had no conflicts of interest with respect to their authorship or the publication of this article.

Acknowledgements.- This study was supported by Conselho Nacional de Desenvolvimento Científico e Tecnológico (CNPq). J.E. Cargnelutti, E.F. Flores and R. Weilen are CNPq research fellows.

REFERENCES
