Veterinary Microbiology

Prostaglandin A₁ inhibits the replication of bovine viral diarrhea virus

Lúcio Ayres Caldas a, †, Tânia Rosária Pereira Freitas b, Renata Campos Azevedo c, Wanderley de Souza d

a Instituto Nacional de Metrologia, Qualidade e Tecnologia (INMETRO), Rio de Janeiro, RJ, Brazil
b Ministério da Agricultura, Pecuária e Abastecimento, Laboratório Nacional Agropecuário – LANAGRO – MG, Pedro Leopoldo, MG, Brazil
c Universidade Federal do Rio de Janeiro, Instituto de Microbiologia Professor Paulo de Góes, Rio de Janeiro, RJ, Brazil
d Universidade Federal do Rio de Janeiro, Instituto de Biofísica Carlos Chagas Filho, Rio de Janeiro, RJ, Brazil

A R T I C L E  I N F O

Article history:
Received 22 September 2017
Accepted 28 December 2017
Available online 1 March 2018
Associate Editor: Fernando Spilki

Keywords:
Bovine viral diarrhea virus
BVDV
Prostaglandins
Prostaglandin A₁
Virus-induced vacuolization

A B S T R A C T

Bovine viral diarrhea virus can cause acute disease in livestock, leading to economic losses. We show that Prostaglandin A₁ inhibits bovine viral diarrhea virus replication in Madin-Darby bovine kidney cells (94% inhibition using 5 μg/mL). Light and electron microscopy of infected cells shows that Prostaglandin A₁ also prevents virus-induced vacuolization, but at higher concentrations (10 μg/mL).

© 2018 Sociedade Brasileira de Microbiologia. Published by Elsevier Editora Ltda. This is an open access article under the CC BY-NC-ND license (http://creativecommons.org/licenses/by-nc-nd/4.0/).

Bovine viral diarrhea virus (BVDV) is an insidious, complex and ubiquitous pathogen that can affect cattle of all breeds and all ages, as well as others domestic and wild ruminants. The disease caused by BVDV ranges in severity from asymptomatic to acute infection and fatal mucosal disease, with a mortality index of approximately 5%, which leads to an economic impact of BVDV infections of up to £993 per cow per year.

In pregnant cows, BVDV can reach the fetus and cause abortion, stillbirth or teratogenic defects, depending on the gestational period when infection was established. When the infection is established in the final period of gestation, calves can give birth to immunotolerant and persistently infected calves that are considered reservoirs of the virus. Also, the serum of infected animals may contain active BVDV viral particles, which may result in permanent infection in primary

† Corresponding author at: Universidade Federal do Rio de Janeiro, Instituto de Biofísica Carlos Chagas Filho, Laboratório de Ultraestrutura Celular Hertha Meyer, Rio de Janeiro, RJ, Brazil.
E-mail: lucio@biof.ufrj.br (L.A. Caldas).
https://doi.org/10.1016/j.bjm.2017.12.010
1517-8382/© 2018 Sociedade Brasileira de Microbiologia. Published by Elsevier Editora Ltda. This is an open access article under the CC BY-NC-ND license (http://creativecommons.org/licenses/by-nc-nd/4.0/).
cultures and established cell lines that are routinely cultivated using bovine serum, in the laboratory.

BVDV is the type-species of the Pestivirus genus (from the Flaviviridae family) that also includes the border disease virus (BDV) and the classical swine fever virus (CSFV), responsible for significant economic losses worldwide. The pestivirus capsid has an icosahedral symmetry formed by a single protein (capsid protein C), and is enveloped by a lipoprotein membrane. Its genome consists of a positive sense, single-stranded RNA of about 12,000 nucleotides, containing a large open reading frame coding for a single polyprotein with about 4000 amino acids. This large polyprotein is cleaved (by host and viral proteases) to generate the capsid protein (C), the glycoproteins E1 and E2, a non-structural protease (Npro) and a number of other non-structural proteins (p7, NS2/NS3, NS4A, NS4B, NSSA and NSSB). The E2 protein appears to be of major importance for pestivirus neutralization.

BVDV isolates are divided into the genetic groups BVDV 1 and 2, which are genetically related to a separate species, the BVDV-3 or Hobi group, which was initially identified in fetal bovine serum from Brazil. BVDV isolates are also classified into cytopathogenic (cpBVDV) or non-cytopathogenic (ncpBVDV) depending on their ability to cause cytopathic effect in cell culture. Birk et al. showed that cpBVDV induced extensive vacuolization in Madin-Darby bovine kidney (MDBK) cells; however, this phenomenon appeared morphologically different from that observed in apoptosis and necrosis.

Prostaglandins are natural eicosanoids secreted by a variety of human tissues and capable of causing profound and diverse physiological effects at very low concentrations. All eicosanoids function locally at the site of synthesis, through receptor-mediated G-protein linked signaling pathways. Prostaglandin A$_1$ (PGA$_1$) is capable of blocking the replication of a wide variety of RNA and DNA viruses, including CSFV. Despite the importance of BVDV for animal health, the effect of PGA$_1$ on BVDV replication has not been examined. Therefore, we evaluate the effect of PGA$_1$ on the replication of cpBVDV in MDBK cells, focusing on changes in BVDV-induced vacuolization.

MDBK cells were cultured in vitro in Dulbecco Modified Eagle Medium (DMEM), supplemented with 10% BVDV-free fetal bovine serum (FBS, Gibco), penicillin (500 U/mL), streptomycin (100 μg/mL) and amphotericin B (fungizone, at 2.5 μg/mL).

Monolayers of MDBK cells were infected with cpBVDV-1 (NADL strain) at a multiplicity of infection of 0.1. Viral particles were allowed to interact with host cells for 1 h in the absence of FBS, and then the supernatant was replaced with complete growth medium containing 0.1, 1, 2.5, 5 or 10 μg/mL Prostaglandin A$_1$ (PGA$_1$, Sigma Chemical Co.; stored as a 1 mg/mL solution in 100% ethanol). After treatment of monolayers with PGA$_1$ for 36 h, culture supernatants were tested for the presence of virus by 50% tissue culture infective dose (TCID$_{50}$) titration, and cell monolayers were fixed and stained with the Panoptic Solution Kit® (Laborclin Ltda. Pinhis/PB, Brazil). Panoptic-stained monolayers were observed in a Zeiss-Observer D1 light microscope equipped with a Nomarski differential interferential contrast system.

Cell viability was determined by incubating monolayers in a solution containing 5 μg/mL Neutral red (diluted in PBS) for 3 h at 37 °C, in wells of 96-well plates. Then, the solution was discarded, and cells were incubated with 4% formaldehyde (in PBS) for 1 min, and 100% methanol for 20 min. Samples were analyzed by absorbance at 490 nm, in an ELISA plate reader (SpectraMax M2).

For transmission electron microscopy (TEM) analysis, treated and untreated monolayers in 25 cm$^2$ plastic culture flasks were fixed in 2.5% glutaraldehyde in 0.1 M cacodylate buffer (pH 7.2), post-fixed for 1 h in 1% OsO$_4$/0.8% potassium ferrocyanide in the same buffer, dehydrated in ethanol and ‘flat-embedded’ in Polycarbonate resin (Polysciences1), to preserve the cell architecture. Ultrathin sections (both routine, and en face) were stained with uranyl acetate and lead citrate and observed in a FEI Tecnai T20 transmission electron microscope.

For the visualization of extracellular lamellae by TEM, 0.5 mg/mL ruthenium red was added to the TEM fixation and post-fixation solutions (as well as to the intermediate washes in cacodylate buffer), and cells were processed for TEM as described above. This procedure was also performed during en face sectioning, in order to avoid removal of the cells from the substratum, which could disrupt and disorient their architecture.

<table>
<thead>
<tr>
<th>PGA$_1$ concentration (μg/mL)</th>
<th>TCID$_{50}$/mL</th>
<th>% Inhibition</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>1×10$^6$</td>
<td>0</td>
</tr>
<tr>
<td>0.5</td>
<td>5.6×10$^6$</td>
<td>44</td>
</tr>
<tr>
<td>1.0</td>
<td>5.0×10$^6$</td>
<td>50</td>
</tr>
<tr>
<td>2.5</td>
<td>1.9×10$^6$</td>
<td>81</td>
</tr>
<tr>
<td>5</td>
<td>6.3×10$^5$</td>
<td>94</td>
</tr>
</tbody>
</table>

MDBK cells were infected with BVDV and treated with different concentrations of PGA$_1$. The percentage of BVDV replication inhibition was calculated by measuring the virus yield in the supernatant of infected cells at 36 h after PGA$_1$ treatment. While 1.0 μg/mL PGA$_1$ inhibited 50% of BVDV production, the highest concentration tested in this assay (5 μg/mL PGA$_1$) blocked replication by >90%. Table 1 – Effect of prostaglandin A$_1$ (PGA$_1$) on bovine viral diarrhea virus (BVDV) replication in MDBK cells, expressed as the TCID$_{50}$ and the percentage of replication inhibition.
Although prostaglandins (PGs) are capable of inhibiting the replication of a variety of RNA and DNA viruses, their mode of action is not entirely clear. In some systems, PGs exert potent antiviral activity by inducing the synthesis of proteins from the heat shock protein 70 (HSP70) family, through cycloheximide-sensitive activation of a heat-shock trigger factor.

The reduction of virus-induced vacuolization after PGA₃ treatment (Fig. 2G) is in agreement with the widely described antiviral effect of this drug. This activity, initially attributed to the induction of HSP-70 proteins, can also be triggered by other pathways, such as nuclear factor-κ B regulation and, more recently, by a mechanism targeting the small ribosomal subunit (40S) and the eukaryotic initiation factors eIF3b.

This virus-induced vacuolization seems not to be crucial for viral replication and is probably caused by the interference of BVDV...
Fig. 2 – Transmission electron microscopy (TEM) analysis of bovine viral diarrhea virus (BVDV)-infected cells treated with prostaglandin A₁ (PGA₁). Infected and non-infected MDBK cells were treated with PGA₁ for 36 h, and then processed for routine TEM. Vacuoles (v) were rare or absent in the majority of sections from non-infected cells (A), but were abundant in BVDV-infected cells not treated with PGA₁ (B). Many of these structures exhibited extracellular content, as shown by ruthenium red labeling (arrows) in traditional (C) and en face (D) sections. Treatment with 1 µg/mL (E) or 2.5 µg/mL (F) PGA₁ could not prevent BVDV-induced vacuolization in infected cells, which had large vacuoles (arrows) with multilamellar structures (arrowheads), and often surrounded by smaller vesicles-containing vacuoles (thin arrows). (G) Treatment with 10 g/mL PGA₁, however, dramatically reduced virus-induced vacuolization. n, nucleus. Scale bars: (A) 5 µm; (B, G) 2 µm; (C, E) 500 nm; (D, F) 1 µm.

Furthermore, the targeting of viral proteins by cyclopentenone prostaglandins was already reported by Kalantari et al.¹⁶ The fact that treatment with 5 µg/mL PGA₁ inhibited BVDV production 94%, but did not block virus-induced vacuolization in host cells suggests that the vacuoles are, indeed, not involved in viral replication and that PGA₁ probably reached its saturation point. Our data suggest that the antiviral activity of PGA₁ is not associated with the inhibition of vacuole production, although the viral-induced vacuoles almost disappeared when higher concentrations of PGA₁ were used.
Our work shows that the antiviral activity of PGA₁ during BVDV interaction with MDBK cells does not rely on a significant reduction in virus-induced cytoplasmic vacuolization, but in the viral titers reduction. Since the antiviral effects of PGs on BVDV had not been described, this is the first report of the parameters involved in the protection of MDBK cells from BVDV infection.

**Funding**

This work was supported by Programa Nacional de Apoio ao Desenvolvimento da Metrologia, Qualidade e Tecnologia (Pronametro) and Coordenação de Aperfeiçoamento de Pessoal de Nível Superior (CAPES).

**References**