

Comparison of an ELISA assay for the detection of adhesive/invasive *Neospora caninum* tachyzoites

Comparação de um ensaio de ELISA para a detecção de taquizoítas adesivos/invasivos de *Neospora caninum*

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Abstract

Neospora caninum belongs to the phylum Apicomplexa, the causative agent of neosporosis, which leads to economic impacts on cattle production. A common feature among apicomplexan parasites is the invasive process driven mostly by the parasite. As a first evaluation of candidate molecules that play a possible role by interfering in this invasive process, the *in vitro* invasion assay is a fast and direct way to screen future agonists or antagonists. This work involved the development of a new cell culture ELISA and transient β -galactosidase activity applied to the semi-quantitative detection of *N. caninum* in Vero cell culture. Cell culture ELISA is based on histochemistry and immunology, resulting in a colorimetric reaction. The β -galactosidase activity was obtained by the transient transfection of the lacZ gene under control of RPS13 promoter of *N. caninum*. These methods were used to evaluate the effects of temperature (37°C and 85°C) on the invasion and adhesion of tachyzoites. The three tested methods (real time PCR, β -galactosidase activity and ELISA) showed a similar pattern, indicating that different methods may be complementary.

Keywords: *Neospora caninum*, cell culture ELISA, LacZ gene, transfection, real-time PCR, invasion assay.

Resumo

Neospora caninum, parasita do filo Apicomplexa, é causador da neosporose, doença responsável por perdas econômicas importantes na pecuária. Um fator comum entre os apicomplexas é o processo de invasão majoritariamente dirigido pelo parasita. Dentre as primeiras avaliações de moléculas candidatas, que possivelmente interferem no processo de invasão, o ensaio de invasão *in vitro* é um meio rápido e direto de selecionar futuros agonistas ou antagonistas. Este trabalho desenvolveu um novo ELISA baseado em cultura (Cell-culture ELISA) e um ensaio que mede a atividade transiente de β -galactosidase, aplicados para a detecção semiquantitativa de *N. caninum* em células Vero. Cell-culture ELISA é baseado em histoquímica e imunologia, resultando em uma reação colorimétrica. A atividade da β -galactosidase foi obtida pela transfecção transiente do gene LacZ sob controle do promotor RPS13 de *N. caninum*. Esses métodos avaliaram os efeitos da temperatura (37°C e 85°C) sobre a invasão e adesão. Os três métodos testados (*real time* PCR, atividade de β -galactosidase e ELISA) mostraram um padrão similar, indicando que diferentes métodos podem ser complementares. Adicionalmente, esse ELISA é adequado para aplicação em laboratórios carentes de uma complexa estrutura para métodos de detecção moleculares.

Palavras-chave: *Neospora caninum*, cell-culture ELISA, gene LacZ, transfecção, PCR em tempo real, ensaio de invasão.

Introduction

Neospora caninum is an apicomplexan protozoan first isolated in 1988 by Dubey and coworkers when they investigated the origin of a severe neuromuscular disorder in dogs (DUBEY et al., 1988). Infected cows represent economic losses due to fetal

abortion, stillbirth, clinical and subclinical disease, impaired milk production, neonatal deaths, reduced fertility and reduced value of infected cows (CORBELLINI et al., 2006). The losses resulting from neosporosis have recently been estimated to exceed US\$1.3 billion, affecting predominantly dairy cattle (REICHEL et al., 2013). Like any member of the phylum Apicomplexa, *N. caninum* is an obligatory intracellular parasite that invades the host cells by a conserved mechanism. Invasion of the host cell by the parasite occurs mainly through an active process, unlike the entry pathway of invasive bacteria (CESBRON-

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DELAUW et al., 2008). The infectious stage of Apicomplexa invades through a multifactorial process driven basically by the protein release of organelles called micronemes, rhoptries, and dense granules (MORAHAN et al., 2009). The direct or indirect detection of *N. caninum*, *in vivo* or *in vitro*, can be accomplished by several methods that include PCR and/or real-time PCR (BELL; RANFORD-CARTWRIGHT, 2002; DARWICH et al., 2012; HUGHES et al., 2006; MORENO et al., 2012), β -galactosidase (HOWE et al., 1997) and ELISA (JENKINS et al., 2002). Real-time PCR in *N. caninum* research using primers for the N5 genomic region of *N. caninum* (GenBank accession no. X84238) is widely applied as a quantitative method due to its fidelity and sensitivity (COLLANTES-FERNANDEZ et al., 2002; GHALMI et al., 2008; NAGULESWARAN et al., 2003). *N. caninum* expressing β -galactosidase is a useful tool to detect tachyzoite integrity in secretion assays (LOVETT et al., 2000; PEREIRA et al., 2011), especially when stored as a stable cytoplasmic enzyme (HOWE et al., 1997). The reproducible and stable reaction with CPRG is well established and widely applied in several microorganisms and cell lineages (D'ANGELO et al., 2009; ESPERANDIM et al., 2010; VAVROVA et al., 2010). The enzyme-linked immunoassay (ELISA) has proven to be a useful technique for antibody detection against a variety of antigens, since it enables rapid determination and antibody titration, making it very suitable for serological surveys (BJORKMAN; LUNDEN, 1998). There are several ELISA methodologies for the detection of *N. caninum* antibodies, based on either whole or fixed *N. caninum* tachyzoites, aqueous or detergent-soluble tachyzoite extracts, tachyzoite antigens incorporated into iscoms (immunostimulating complexes) or recombinant tachyzoite antigens (JENKINS et al., 2002).

All these immunoenzymatic techniques have been used for diagnosis, but none of them for measuring the invasion of *N. caninum*. This work introduces a modified ELISA, based on a Vero cell culture, as a semi-quantitative assay to detect adhesive and/or invasive tachyzoites after an invasion assay. Unlike the existing ELISAs, this assay is based on histochemistry and immunology, allowing for the relative mensuration of invasion by means of a colorimetric reaction. This new method offers a direct and low cost route for detecting *N. caninum* after invasion assays, without steps such as protein or DNA extraction and purification. In addition to real time PCR, for comparison, *N. caninum* expressing β -galactosidase was obtained after transient transfection with a LacZ gene vector controlled by the RPS13 promoter of *N. caninum*. This paper reports the use of *N. caninum* promoters for heterologous expression, since most other studies have employed *T. gondii* promoters. The adhesion and/or invasion pattern of *N. caninum*, as well as the different effects of incubation temperature and time, were analyzed by different methods. In addition, the ELISA and β -galactosidase assays proved to be suitable alternatives to real-time PCR as screening methods for evaluating the effects of drugs on the *N. caninum* invasion process.

Materials and Methods

Tissue culture and parasite purification

N. caninum Nc-1 isolate was cultivated in Vero cell cultures maintained in RPMI-1640 medium (Sigma) supplemented with

5% fetal calf serum, 2.05 mM glutamine, 50 U of penicillin/streptomycin at 37°C, and 5% CO₂ in T-25 cm² and 75 cm² tissue culture flasks (for *N. caninum* culture) or on 24 and 96-well culture plates (for invasion assay). The purification of tachyzoites was performed by exclusion chromatography in Sephadex G-25 (PD-10 columns, GE). The recovered parasites were counted in a Neubauer chamber after adequate dilution; the average recovery from a 25 cm² flask culture was 2 × 10⁷ tachyzoites.

Evaluation of N. caninum viability by flow cytometry

Freshly purified *N. caninum* tachyzoites (1 × 10⁶ in 250 μ l PBS) incubated at 37°C for 1 h or 85°C for 20 min were evaluated by flow cytometry (FACSCanto, BD Biosciences) with an argon laser set to 488 nm, and evaluated using FACSDiva BD 6.1.3 software. The gate corresponding to *N. caninum* characteristics was drawn by forward (size, FSC) and side scattering (complexity, SSC). Tachyzoite viability was evaluated by the incorporation of propidium iodide (20 ng/ μ l, Santa Cruz Biotechnology), with excitation at 535 nm and detection at 617 nm. The percentage of live and dead (negative and positive to propidium iodide, respectively) tachyzoites was determined.

Invasion assays

All the invasion assays were performed according to the following protocol. Briefly, three concentrations of *N. caninum* tachyzoites (3.5 × 10⁶, 1.75 × 10⁶ and 3.5 × 10⁵ for ELISA and β -galactosidase, or 3.5 × 10⁵, 1.75 × 10⁵ and 3.5 × 10⁴ for real-time PCR) were diluted in 650 μ l RPMI. Tachyzoites were treated differently: i) applied immediately after purification, ii) 37°C for 1 h, or iii) 85°C for 20 min. These suspensions (650 μ l) were divided into three parts (200 μ l each) and applied to Vero cells cultured in 96-well (ELISA) or 24-well plates (β -galactosidase or real-time PCR). The tachyzoites were allowed to invade for 2 h at 37°C, after which each well was washed three times with PBS and subjected to the three specific detection methods: ELISA, β -galactosidase and real-time PCR.

Detection of N. caninum by ELISA

The 96-well plates (TPP) were fixed with 3% formamide in PBS for 15 min, permeabilized with PBS/0.2% Triton X-100 for 30 min, washed once with PBS, and incubated with blocking buffer solution (2% BSA, 50 mM glycine in PBS) for 16 h at 4°C. After washing with PBS, antiserum against *N. caninum* (anti-Nc tachyzoite protein extract, 1:1000 (PEREIRA et al., 2011)) was incubated for 1 h, followed by three washes in PBS. The secondary antibody goat anti-rabbit-peroxidase conjugate (1:10000, ZyMed-Invitrogen) was incubated for 1 h and then washed three times. The wells were incubated for 15 min with a TMB solution (ZyMed-Invitrogen). The blank was composed of the same antiserum but incubated in Vero cells, since the anti-Nc tachyzoite protein extract was raised against protein extract of *N. caninum* that had been cultivated in Vero cells, thus eliminating the response against these cells after the subtraction of Vero absorbance. The reaction was stopped with a 2 M HCl

solution and measured at 450 nm (ELISA reader, SpectraMax Plus, Molecular Devices). These assays were performed in triplicate.

Construction of β -galactosidase vector

N. caninum expressing β -galactosidase was obtained via the transient insertion of the NcRPS13 subIV23LacZ plasmid. This vector was obtained after replacing the *Toxoplasma gondii* promoter region of RPS13 ((TgRPS13 with 1353bp from TgRPS13subIV23LacZ (van POPPEL et al., 2006)) with the homologous region from *N. caninum* (NcRPS13) (Figure 1). Primers with *Hind*III and *Nru*I restriction sites were designed to amplify a region of 663bp upstream of the *N. caninum* RPS13 ORF. This fragment was linked to TgRPS13subIV23LacZ, which was pretreated with *Hind*III and *Nru*I, generating NcRPS13subIV23LacZ. The β -galactosidase activity was evaluated (data not shown). The tetO region is a site with affinity for the TetR protein, which, once bound to the tetO site, blocks expression and is released from the site by tetracycline. This system is based on the *E. coli* tetracycline operon, which has been adapted for *T. gondii* and *Plasmodium falciparum* (DAS et al., 2011; MEISSNER et al., 2001, 2002; van POPPEL et al., 2006). However, only the β -galactosidase expression was employed in this study.

Transfection of *N. caninum* for β -galactosidase expression

Freshly purified tachyzoites (5×10^7) were resuspended with 25 μ g of NcRPS13 subIV23LacZ in cytomix buffer (120 mM KCl, 0.15 mM CaCl₂, 10 mM K₂HPO₄/KH₂PO₄, 25 mM HEPES, 2mM EDTA, and 5 mM MgCl₂, pH 7.6) supplemented with 5 mM glutathione and stored on ice. Cells were transferred to a 0.4 cm gap cuvette (in ice) and electroporated at 1.8 kV, 25 μ Fd and 100 Ω in a BioRad GenePulser Xcell system. The electroporated cells were incubated in Vero cells cultured (75 cm² flask, TPP) in incomplete RPMI (without fetal bovine serum) until total lysis of host cells occurred. The tachyzoites were purified by exclusion chromatography in Sephadex G-25 (PD-10 columns, GE) under the same conditions as the wild tachyzoites (see 2.1).

Detection of *N. caninum* by β -galactosidase

Adhesive/invasive *N. caninum* were incubated in lysis buffer (4-(2-hydroxyethyl)-1-piperazine ethanesulfonic acid, 100 mM

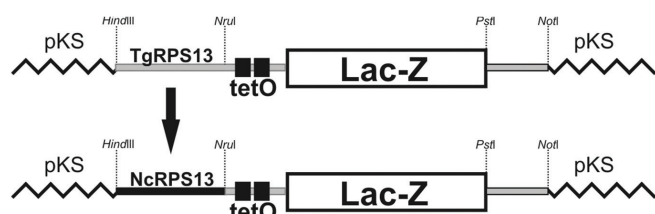


Figure 1. NcRPS13subIV23LacZ vector. The promoter region of the RPS13 gene from *N. caninum* (NcRPS13) was ligated (flanked by *Hind*III and *Nru*I) to control the expression of LacZ, replacing the original promoter TgRPS13 (*T. gondii*) of vector TgRPS13subIV23LacZ from van Poppel et al. (2006).

HEPES, 1mM MgSO₄, 1% Triton X-100, 5 mM Dithiothreitol, DTT, pH 8.0) at 50°C for 1h. The lysate was centrifuged at 10,000g at 4°C for 10 min and 20 μ l of supernatant was solubilized in 190 μ l of buffer lysis with 40 μ l of 6 mM CPRG (chlorophenol red- β -D-galactopyranoside, Roche). The reaction was measured at 570 nm after 18 h at 37°C (ELISA reader, SpectraMax Plus, Molecular Devices). The invasion assays were performed in triplicate and the lacZ activity in each well was evaluated in duplicate (final n = 6).

Detection of *N. caninum* by real-time PCR

N. caninum DNA was extracted and purified from each well according to the manufacturer's instructions (Wizard SV Genomic DNA Purification System, Promega). Parasites were quantified by real-time PCR, using Nc 5 primers with the forward primer linked to FAM fluorophore Lux (Invitrogen) (PEREIRA et al., 2011). The experiment was carried out in a Mastercycler ep RealPlex (Eppendorf) and the results analyzed using RealPlex software (Eppendorf). As external standards, DNA corresponding to 1 to 10⁵ *N. caninum* tachyzoites was used for the standard curve and the samples were calculated by interpolation from this standard curve. The invasion assay was performed in triplicate and the real-time reaction in duplicate for each well (n = 6).

Statistical analyses

The results of invasion assays were expressed as mean \pm standard deviation and analysis of variance (ANOVA) was employed for the dilutions and for 37°C/fresh and 85/37°C relations using the Tukey test, considering statistical significance at a level below 5%. The overall analyses of dilutions were performed by linear regression and the r² values were calculated. The analyses were performed using the Graph Pad PRISM 4.0 (Graph Pad, USA) statistical program.

Results

Evaluation of live/dead *N. caninum* by flow cytometry

The percentage of live parasites was determined to corroborate the invasion assays, confirming that the decrease in invasion capacity after incubation was due to the infeasibility of tachyzoites. This assay was also useful for detecting dead parasites that maintained their adhesive function. The gate representing size (forward scatter, axis x) and granularity (side scatter, axis y) of the *N. caninum* tachyzoites treated at different temperatures (Figures 2A, B, C-1) was delimited and the groups were separated according to their fluorescence emission for live and dead tachyzoites after propidium iodide treatment. The rate of live parasites was 81.9% for freshly purified organisms, and 76.5% for 37°C and 1.5% for 85°C incubated parasites (Figures 2A, B, C-2).

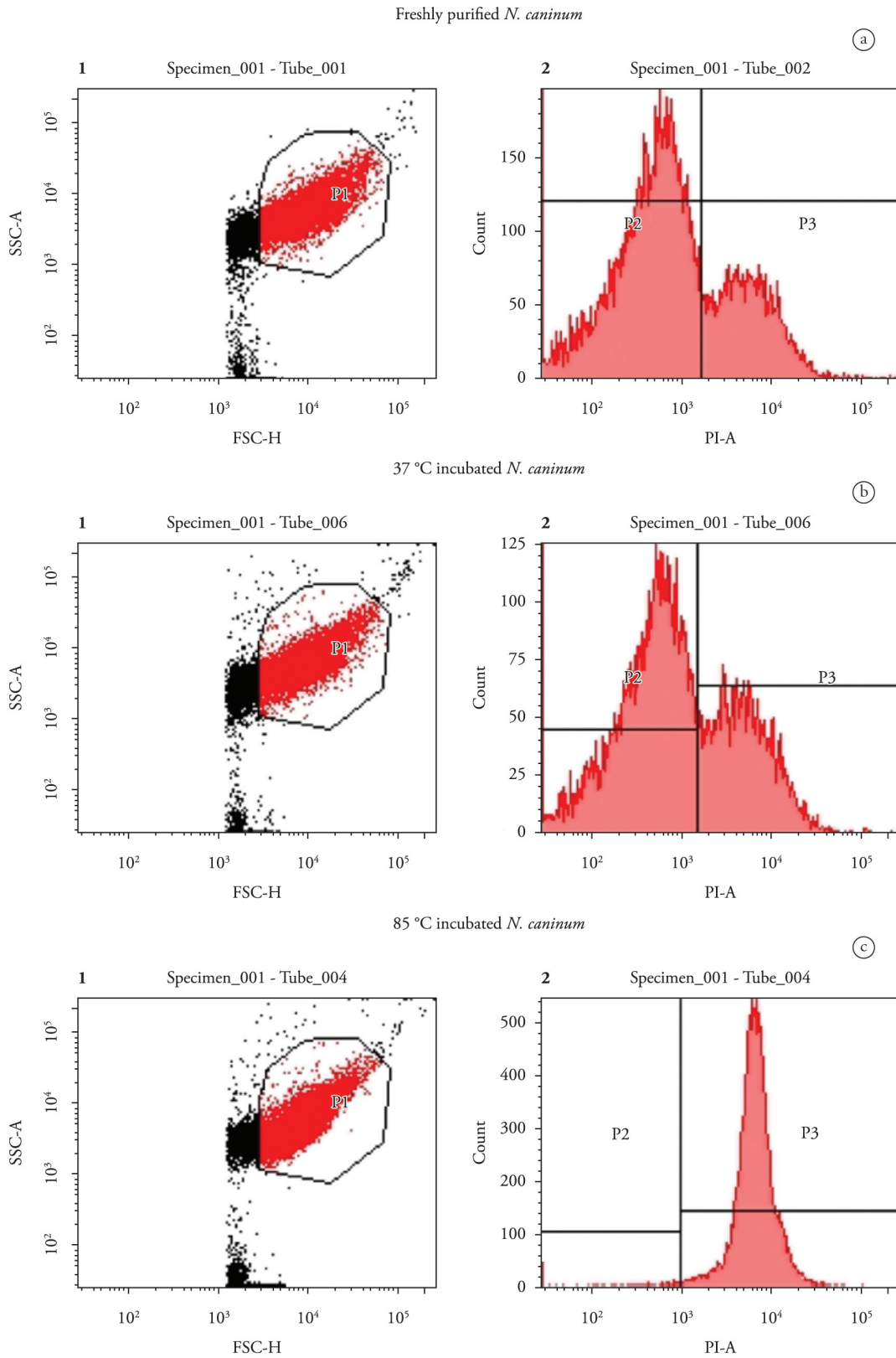


Figure 2. Evaluation of tachyzoite mortality with flow cytometry after different temperature treatments. Tachyzoites (1×10^6) incubated at 37°C, 85°C or freshly purified were mixed with propidium iodide (PI, 20 ng/ μ l) and immediately analysed by flow cytometry. Cytograms 1A, 1B and 1C plot forward scatter (FSC), representing size, against side scatter (SSC) representing granularity. Cytograms 2A, 2B and 2C plot count against propidium iodide emission (PI-A). A: Freshly purified tachyzoites with no temperature treatment; B: Tachyzoites incubated at 37°C for 1 hour; C: Tachyzoites incubated at 85°C for 20 minutes. The squares indicate the percentage of tachyzoites detected negatively (P2) and positively (P3) by propidium iodide.

Semi-quantification of incubated tachyzoites by ELISA, β -galactosidase and real-time PCR

N. caninum tachyzoites detected by ELISA, β -galactosidase and real-time PCR exhibited a similar tendency. In the ELISA assay, the absorbance of tachyzoites incubated at 85°C and 37°C showed lower values than those of freshly purified ones. The three concentrations of tachyzoites exhibited a linear pattern with r^2 of 0.94, 0.98 and 0.93, respectively, for incubations at 85°C and 37°C and freshly purified tachyzoites (Figure 3A). The lacZ activity indicated the same decreasing pattern of invasive/adhesive tachyzoites when incubated at 37°C (the r^2 of 37°C-incubated and freshly purified tachyzoites were 0.95 and 0.96, respectively, Figure 3B). After incubation at 85°C, the detection of LacZ activity was not feasible because the enzyme β -galactosidase was denatured. Due to the high sensitivity of the real-time PCR assay, the number of tachyzoites employed in the invasion assay was 10-fold lower than for the ELISA and β -galactosidase assays, with 1×10^5 , 5×10^4 and 1×10^4 (Figure 3C). The 85°C-incubated tachyzoites indicated an r^2 of 0.87 with the three concentrations of tachyzoites, the incubation at 37°C yielded an r^2 of 0.95 and the freshly purified tachyzoites an r^2 of 0.96.

Effects of temperature on the efficiency of tachyzoite invasion

Two ratios were calculated to analyze the effects of different temperatures on the tachyzoites. First, a correlation between tachyzoites incubated at 37°C and the freshly purified cells was calculated (ratio of 37°C/fresh*100), which represents the tachyzoites able to adhere to and/or invade host cells after a 37°C incubation. Secondly, the correlation between tachyzoites incubated at 85°C and 37°C was calculated (ratio of 85°C/37°C*100), which represents the tachyzoites able to adhere to host cells even after death.

The ELISA assay applied as an alternative method to measure invasion indicated a high correlation between tachyzoites incubated at 37°C and those that were freshly purified: 77.2% (1×10^6 tachyzoites/well), 66.2% (5×10^5) and 50.9% (1×10^5) (Fig. 4A). On the other hand, 32.2% (1×10^6), 21.8% (5×10^5) and 24.8% (1×10^5) of dead tachyzoites were still able to adhere (Figure 4A). The real-time PCR indicated adhesion after death at 85°C, with the ratio of 85°C/37°C decreasing to 39.9% (1×10^5), 44.5% (5×10^4) and 27.0% (1×10^4) (Figure 4C), while that of 37°C/freshly purified tachyzoite was 92.7% (1×10^5), 71.1% (5×10^4) and 59.5% (1×10^3) (Figure 4C). The β -galactosidase method maintained the correlation between 37°C/freshly purified with 85.03% for 1×10^6 , 79.44% for 5×10^5 and 78.97% for 1×10^5 (Figure 4B), which was similar to that of ELISA and real-time PCR. Overall, there were no significant differences among the dilutions, except for tachyzoites at 1×10^5 and 1×10^6 with the correlation 37°C/freshly purified tachyzoites (Figure 4A).

Discussion

N. caninum direct or indirect detection in cell cultures and tissues is well established by traditional methods, such as IFAT

(YAKHCHALI et al., 2010), and highly specific methods such as real-time PCR (DEBACHE et al., 2010; GHALMI et al., 2008; NAGULESWARAN et al., 2003; PEREIRA et al., 2011). The three methods tested in this study for the detection of *N. caninum* invasion and/or adherence in cell culture showed a similar pattern, indicating that different methods may be complementary, since

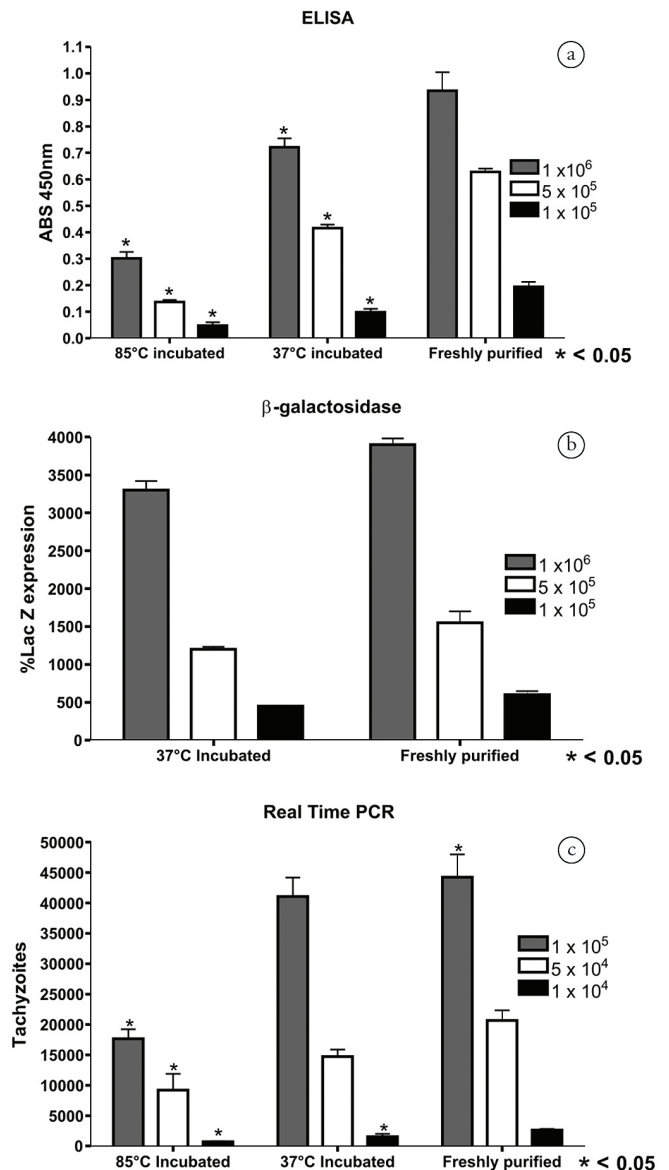


Figure 3. Detection of *N. caninum* incubated at 37°C, 85°C or freshly purified by ELISA, β -galactosidase and real-time PCR. The tachyzoites were diluted 1×10^6 , 5×10^5 and 1×10^5 /well for ELISA and β -galactosidase and 1×10^5 , 5×10^4 and 1×10^4 for real-time PCR and divided in three groups before invasion. After purification, one part was incubated at 37°C for 1hr, the second part incubated at 85°C for 20 min and the third part was immediately allowed to invade Vero cells (freshly purified). A: ELISA; B: β -galactosidase; C: Real-time PCR. The detection was performed with antisera (ELISA), expression of LacZ gene in *N. caninum* transiently transfected with a LacZ plasmid (β -galactosidase), or with *N. caninum* DNA measurement (real-time PCR). * represents significant difference < 0.05 in relation to the freshly purified group.

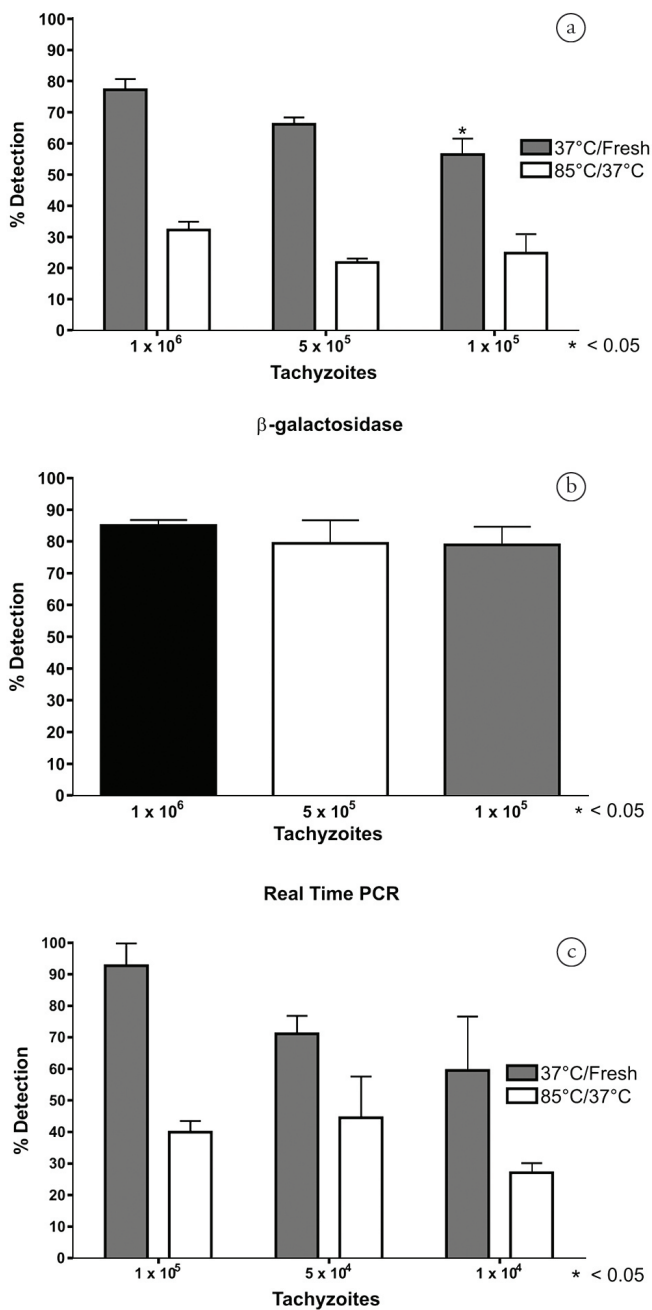


Figure 4. Decrease in adhesion/invasion of *N. caninum* after incubation at 37°C or 85°C by ELISA, β -galactosidase and real-time PCR. The correlations between 37°C/freshly purified and 85°C/37°C incubated tachyzoite were calculated after detection by ELISA (A, anti-tachyzoite protein extract), and real-time PCR (C, primers Nc-5). For the β -galactosidase activity (% lacZ expression), only the 37°C/fresh correlation was obtained because the enzyme was inactivated at 85°C. Tachyzoites were incubated in three dilutions: 1×10^6 , 5×10^5 and 1×10^5 /well for ELISA and β -galactosidase and 1×10^5 , 5×10^4 and 1×10^4 for real-time PCR. A: ELISA; B: β -galactosidase; C: Real-time PCR. *represents a significant difference (<math>p < 0.05</math>).

ELISA, β -galactosidase and real-time PCR are based on different principles, i.e., antibodies, enzymatic reporter and genomic DNA, respectively. The number of tachyzoites per well applied in β -galactosidase and ELISA ranges from 10^5 - 10^6 , limiting evaluations to lower parasite loads; hence, real-time PCR may be the best choice. Thus, the first two methods are suitable for an initial evaluation of parasite invasion and real-time PCR participates as an unequivocal quantitative assay. The incubation of *N. caninum* or *T. gondii* at 37°C is a current way to determine the *in vitro* effects of drugs or sera on the invasion phenomenon (NAGULESWARAN et al., 2003; PEREIRA et al., 2011; SAWMYNADEN et al., 2008). Our results indicate that all these methods show losses of invasive/adhesive tachyzoites incubated at 37°C; therefore, this is an important effect to take into account when designing assays that require long incubation times.

On the other hand, *N. caninum* maintains a basal level of adhesion even after incubation at 85°C, which is a temperature that leads to the death of 98.5% of the tachyzoites, according to propidium iodide positivity. Such a high temperature probably denatures many of the tachyzoite's surface proteins; however, a basal adhesion capacity is detected and may indicate mechanisms of parasite/host affinity independent of the parasite's viability. Propidium iodide is only internalized when the membrane is disrupted; conversely, membrane integrity is necessary to enable the initial tachyzoite invasion process (CARRUTHERS; BOOTHROYD, 2007).

A semi-quantitative method for the detection of *N. caninum* based on ELISA after invasion assays in cell culture integrates characteristics of histochemistry (fixation, permeabilization) and immunology (ELISA, indirect detection of antibodies by colorimetric peroxidase reaction), offering an additional fast, low cost method to determine the influence of substances on the *in vitro* invasion process. Albeit highly specific and sensitive, real-time PCR requires high quality inputs, such as primers, probes, thermal cyclers with suitable detectors, software and an optimal standardization (BELL; RANFORD-CARTWRIGHT, 2002). On the other hand, it allows for quantitation and requires lower cell concentrations in the assays. The high specificity and sensitivity of real-time PCR led to higher ratios (37°C/fresh and 85°C/37°C), probably due to the higher tachyzoite detection capacity of this technique compared to that of ELISA and β -galactosidase. The β -galactosidase method is less laborious to standardize; however, it requires tachyzoites expressing β -galactosidase (HOWE et al., 1997, the expression of LacZ gene was controlled by *T. gondii* promoters), which is absent in wild-type Nc-1 tachyzoites. Although extremely well developed in *T. gondii*, the expression of heterologous genes in *N. caninum* is not an ordinary technique.

This newly described ELISA assay requires an antiserum against *N. caninum* tachyzoite protein extract. Its specificity/sensitivity led to a detection cut-off of 1×10^5 tachyzoites per well, which was verified by the differences observed between concentrations of 1×10^5 and 1×10^6 tachyzoites incubated at 37°C (Figure 4A), with a significant difference probably due to the small number of tachyzoites required by this method. The incubations indicated that the number of parasites is a limiting factor for optimal detection by ELISA, although a similar condition was found for β -galactosidase and only real-time PCR presented higher sensitivity.

The polyclonal sera against the tachyzoites extract obtained from urea 8M buffer raises a wide range of antibodies, which offers a robust absorbance in *N. caninum* detection. A significant number of antibodies recognize only *N. caninum* proteins, and for 1×10^6 tachyzoites, ELISA is as adequate as beta-gal or real time assays, which validate the method. The difference between *N. caninum* and Vero cell absorbance is 0.720 in parasites incubated at 37°C for 1 hour, a suitable range for invasion evaluations (Figure 3A). The ELISA method described here appears to be a low cost route to evaluate the invasion process of *N. caninum*, a simple and fast method for laboratories whose routine lacks a structure for the application of molecular tools. In addition, β -galactosidase activity showed the applicability of *N. caninum* promoters for heterologous expression. The ELISA and the β -galactosidase assays exhibited a pattern similar to that of the highly specific and sensitive real-time PCR, indicating the effects of incubation and temperature on the viability and invasion process of *N. caninum*. Overall, this ELISA method is expected to support the design of more adequate assays for screening agonists and antagonists that are active in the invasion process.

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