Sheep polyclonal antibody to map *Haemonchus contortus* mimotopes using phage display library

Anticorpo policlonal de ovinos para mapear mimetopos de *Haemonchus contortus* usando a biblioteca de Phage display

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Abstract

The aim of this study was to evaluate phage display technology for mapping *Haemonchus contortus* mimotopes. We screened the PhD-7 Phage Display Peptide Library Kit with a sheep polyclonal antibody against *H. contortus*. After four rounds of selection, 50 phage peptide clones were selected by biopanning and sequenced. Two clones displaying peptide mimotopes of *H. contortus* proteins were chosen for sheep immunization: clone 6 - mimotope of glyceraldehyde-3-phosphate dehydrogenase (GAPDH) and clone 17 - mimotope of a disorganized muscle family member (Dim 1). Twelve sheep were allocated into 3 groups of 4 animals as follow: G1: control group; G2/GAPDH: immunized with clone 6; and G3/Dim1: immunized with clone 17. Four immunizations were performed at intervals of seven days (0, 7, 14, and 21 days). On day 28 post initial vaccination, all groups were orally challenged with 2500 *H. contortus* infective larvae. The mimotope peptides selected by phage display were recognized by IgG from sheep naturally infected with *H. contortus*. The immunization protocol showed an increase in IgG anti-M13 phage titers, but no effect was observed in IgG-specific for the anti-mimotope peptides. This is the first report of successful use of a phage display library for the identification of mimotopes of *H. contortus* proteins.

Keywords: Mimotopes, biopanning, ELISA, ruminants, immunogenic.

Resumo

O objetivo deste estudo foi avaliar a tecnologia de phage display no mapeamento de mimetopos de *Haemonchus contortus*. Anticorpo policlonal de ovinos anti-*H. contortus* foi usado para seleção a partir da biblioteca PhD-7 Phage Display Peptide Library Kit (New England BioLabs). Após quatro rodadas, 50 clones de fagos expressando peptídeos foram selecionados e sequenciados. Dois clones que exibiram mimetopos de *H. contortus* foram escolhidos para imunização de ovinos: clone 6 – mimetopo de gliceraldeído-3-fosfato desidrogenase (GAPDH) e clone 17 – mimetopo da família do músculo desorganizado (Dim 1). Doze ovinos foram alocados em 3 grupos de 4 animais, da seguinte forma: G1: grupo controle, G2/GAPDH: imunizado com o clone 6 e G3/Dim1: imunizado com o clone 17. Quatro imunizações foram realizadas (0, 7, 14 e 21 dias). No dia 28 após a primeira imunização, todos os grupos foram desafiados oralmente com 2500 larvas infectantes de *H. contortus*. Os peptídeos mimetopos selecionados foram reconhecidos por IgG de ovinos naturalmente infectados por *H. contortus*. O ensaio de imunização revelou um aumento dos títulos de IgG anti-fago M13, mas não ocorreu aumento de IgG anti-peptídeos mimetopos. Este é o primeiro relato de uso bem sucedido da biblioteca de Phage display para a identificação de mimetopos de *H. contortus*.

Palavras-chave: Mimetopos, biopanning, ELISA, ruminantes, imunogênicos.

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Introduction

*Haemonchus contortus* is a hematophagous gastrointestinal parasite of ruminants that is extremely important on a global scale. It is one of the most prevalent parasites, causing a lot of damage to animals, thereby having a substantial impact on farm production. The parasite has been reported to outlive the use of different anthelmintic products (FORTESS & MOLENTO, 2013).

Phage display is a low-cost and fast method for mapping the epitope of an antigen that is involved in a specific protein interaction with the antibody. The identification of epitopes is essential in diagnostics, immunotherapy, drug discovery and vaccine development to combat diseases (PANDE et al., 2010). This application, enables selection of mimotopes, peptides mimicking natural epitopes of a pathogen, even without prior knowledge of the natural ligand area (ELLIS et al., 2012).

Concerning parasite control, phage display library technology is still a relatively novel research area (ELLIS et al., 2012) and it has not been applied previously in the screening of *H. contortus* mimotopes.

Several reports have demonstrated the use of phage display libraries for targeting and diagnosis of parasites (HELL et al., 2009; GAZARIAN et al., 2012; CHEONG et al., 2016; RHAIEEM & HOUMEL, 2016), and promising studies on protective mimotopes have been reported on *Plasmodium falciparum* (ADDA et al., 1999), *Schistosoma japonicum* (TANG et al., 2004), *Trichinella spiralis* (GU et al., 2008), *Fasciola hepatica* (VILLA-MANCERA et al., 2008), *Rhipicephalus (Boophilus) microplus* (PRUDENCIO et al., 2010), and *Taenia solium* (ASSANA et al., 2010). The objective of this study was to evaluate for the first time, the use of phage display technology for mapping *H. contortus* mimotopes.

Materials and Methods

*Haemonchus contortus polyclonal antibodies*

Forty-eight adult sheep from a herd naturally infected with *H. contortus* and exposed to repeated infections, were used to obtain serum. We selected serum only from the animals with high IgG titer and zero parasite fecal egg count (FEC).

Blood samples (10 ml) were collected from the jugular vein of all animals. The serum was separated by centrifugation at 3000 x g for 10 min and stored at -20°C until its use for the ELISA analysis of the selected clones.

**Extraction of Haemonchus contortus proteins**

Fifty worms were crushed in liquid nitrogen using a mortar and pestle. Protein extraction was performed using chilled 0.1% [v/v] PBS and Tween 20, with shaking for 1 h at 4°C. The samples were centrifuged at 10,000 x g for 20 min at 4°C. The supernatant was filtered and aliquots were kept at -20°C until use. The protein concentration was determined by Qubit protein assay (Invitrogen, USA).

Enzyme-linked immunosorbent assay (ELISA)

The 96-well ELISA plates (PolySorp, Nunc) were coated with 5 µg ml⁻¹ (100 µl well⁻¹) of *H. contortus* total proteins (DIAZ et al., 2015) (5 µg ml⁻¹ of chemically-synthesized peptides or 10¹¹ plaque-forming units or 5 µg ml⁻¹ according to the assay) diluted in coating buffer (0.1 mM sodium carbonate-bicarbonate, pH 9.6) and incubated at 4°C overnight. The plates were washed four times with 0.05% [v/v] Tween 20 solution (Merck, USA). The blocking of the reaction was performed with 3% [v/v] PBS for 1 h at 20°C. The plates were washed four times with 0.05% [v/v] Tween 20 solution. Sheep sera were used at 1:100, 1:300, 1:600, 1:1200, 1:2400, 1:4800, 1:9600 and 1:19200, depending on the assay, in PBS 1 x 0.2% [v/v] Tween 20 and incubated for 1 h at room temperature. After incubation, the plates were washed four times with Tween 20 solution 0.05% [v/v] and the anti-sheep IgG HRP 1:10.000 (Sigma A3415) was added. The plates were washed four times with 0.05% [v/v] Tween 20 solution and ABTS [2,2′-azino-bis(3-ethylbenzothiazoline-6-sulphonic acid)] buffer was added. The plates were then incubated in the dark at 20°C for 30 min. The optical density was determined using a microplate reader (BioTek, USA) at 405 nm. The cut-off optical density (OD) was determined using the mean OD + 3xSD (standard deviation) of the negative control (serum of sheep not infected with *H. contortus*).

Titration, purification and amplification of the phage

The PhD-7 Phage Display Peptide Library Kit (New England BioLabs, USA) was used according to methodology previously described by Wu et al. (2006). The technology consists of a randomized linear 7-mer peptides fused to a minor coat protein (pIII) of M13 phage.

Peptide selection from Phage-Display Library

A 96-well plate (PolySorp, Nunc) was coated with IgG protein-G purified from sheep polyclonal sera that recognized *H. contortus* (1:100), in 100 µl of binding buffer (0.1M NaHCO3, pH 9.6), overnight at 4°C. The panning process was performed according to Wu et al. (2006).

Four rounds of selection were performed, after which an individual plaque was picked up at random and subjected to analysis by ELISA and DNA sequencing, following amplification in *Escherichia coli* ER2537.

Analysis of the selected clones

After the four selection rounds, each clone obtained had its DNA purified and sequenced using a capillary electrophoresis apparatus (ABI3130) using BigDye® terminator v 3.1 and POP 7 polymer. The sequence analyses were carried out using the software Vector NTI (version 6.0) and Molecular Evolutionary Genetics Analysis (version 6.06). We evaluated the identity among the clones and also with *H. contortus* protein sequences available in GenBank.
Evaluation of peptides as mimotopes

Serum samples of 25 sheep naturally infected with *H. contortus* were subjected to ELISA testing (section 2.3) to evaluate the peptides’ ability as mimotopes. The plates were coated with two chemically-synthesized peptides, which had the same sequences and conformation (7 amino acids - aa) displayed on two clones selected by the biopanning, and used in the vaccination assay.

Immumization protocol

To evaluate the potential of the selected mimotopes as experimental vaccine candidates, two clones displaying the peptides were selected to perform the *in vivo* study. Clone 6 - similar to the glyceraldehyde-3-phosphate dehydrogenase (GAPDH) and clone 17 - similar to a disorganized muscle family member (Dim 1) were used. Twelve sheep, 8-10 months old, raised indoors under nematode-free conditions, were allocated into 3 groups of 4 animals as follows: G1: control group - infected but not immunized; G2: GAPDH - immunized with clone 6; and G3: Dim 1 - immunized with clone 17. Vaccinated animals were immunized with 5x10^{11} pfu with the adjuvant Montanide ISA 25 VG according to the manufacturer’s recommendations. G1 was unvaccinated but received 1 ml of the same adjuvant. Four immunizations were performed at seven-day intervals (0, 7, 14 and 21 days). On day 28 post-vaccination, all groups were orally challenged with 2500 infective *H. contortus* larvae (L3). All sheep were humanely slaughtered on day 63.

Serological analysis

Sheep serum samples, corresponding to days 0; 14; 21; 28; 44; 51; and 63 were used for testing antibody response (ELISA section 2.3). All samples were examined against five antigens: 1: M13 phage, 2: Clone 6, 3: Clone 17, 4: chemically-synthesized GAPDH (7 aa), and 5: chemically-synthesized Dim 1 (7 aa).

Haematological analysis

Blood samples were collected on day 0; 14; 28; 44 and 50 following the first vaccination, by jugular venipuncture into vacuum glass tubes containing EDTA as anti-coagulant. The samples were used for leukogram and haemoglobin determination. Differential cell counts were also carried out.

Fecal egg count (FEC)

Fecal samples were collected from each sheep at day 0; 28; 44; 47; 51 and 55 of the experiment. FEC was determined by a modified Gordon & Whitlock (1939) method, with a sensitivity of 25 eggs per gram of feces. Previously the experiment start (day 0), we analyzed fecal samples of all animals for 4 consecutive days (day -4; day -3; day -2 and day -1) to confirm the absence of infection by gastrointestinal parasites.

Abomasum worm counts

The number of worms in the abomasum from all slaughtered animals was determined on day 63. The abomasal content was collected and the mucosa was scraped and washed with warm 0.9% sodium chloride to detach the worms. All *H. contortus* were picked up, collected, counted and sorted according to their sex.

All protocols were approved by the Animal Care and Ethics Committee of Camaguey University, Cuba (protocol No. 2013001).

Statistical analyses

The data were analyzed using the Shapiro-Wilk normality test. The Kruskal-Wallis test was used to evaluate differences between groups (P<0.05).

Results

Phage display library biopanning

To map the epitopes of *H. contortus*, a random heptapeptide phage display library (PhD-7) comprised of 1.0x10^{12} independent phage clones was screened with the coated sheep polyclonal antibodies (*H. contortus*-IgG). After four rounds of biopanning, the selected phage bound to *H. contortus*-IgG were well enriched (2100X), as indicated by the increased recovery. The number of phage peptides bound to *H. contortus*-IgG increased from 2.2x10^{4} pfu in the first round to 4.6x10^{7} pfu in the fourth round.

After four rounds of selection, 50 peptide phage clones were selected by biopanning and sequenced by capillary electrophoresis. The analysis of the sequences (Figure 1) revealed degrees of identity among some clones and also to proteins of *H. contortus*, namely, Glyceraldehyde-3-phosphate dehydrogenase (GAPDH) and a Disorganized Muscle Family Member (Dim 1).

The identity among clones 9; 10 and 32 was 100%; and among the clones 21; 25; 26; 37; 39 and 40 was 57.14%. The identity between GAPDH and clone 6 was 71.42%; the identity among Dim1, clone 11 and clone 17 was 57.14.

![Figure 1](image-url)

**Figure 1.** Phage display technology was employed to select mimotopes of *Haemonchus contortus* peptides. Analyses of the phage cloned sequences were performed with MEGA software Version 6.06. Consensus sequence (*) was observed among amino acids of some clones (a, b) and also to two *H. contortus* proteins, Glyceraldehyde-3-phosphate dehydrogenase (c) and a Disorganized muscle family member – Dim 1 (d).
The evaluation of the chemically-synthesized peptides as mimotopes is demonstrated in Figure 2 (a and b). The serum samples analyzed against the peptide GAPDH (Figure 2a) revealed 24 samples with IgG higher than the cut-off and just one sample below the cut-off. When used against Dim 1 (Figure 2b), 12 samples demonstrated IgG means above and 12 samples below the cut-off value.

**Immunization trial**

Circulating antibody responses were observed in both vaccinated groups from day 14 onwards (Figure 3b and 3c). The same animals also revealed IgG against the M13 phage (Figure 3a). The IgG titers of the vaccinated groups were similar and followed the same pattern, rising after the first immunization (Day 0) and reaching a peak on day 28. The animals did not show an IgG response specifically against the GAPDH (Figure 3d) or Dim 1 (Figure 3e) chemically-synthesized peptides.

The infection with *H. contortus* was confirmed on day 44. We did not detect any significant FEC reduction within the experimental groups; nor for the worm counts (P>0.05) after vaccination (data not shown). The results from leukogram and haemoglobin analyses are shown in Table 1, with a lack of significant variations during the experimental period.

**Table 1.** Hemoglobin, total leukocyte number and percentage of lymphocyte, neutrophil and eosinophil cells in blood are shown as means ± SD. Twelve 8-10 month-old Suffolk sheep were allocated into 3 groups of 4 animals as follows: Control: infected but not immunized; GAPDH: immunized with clone 6; and Dim 1: immunized with clone 17. Four immunizations were performed (0, 7, 14 and 21 days) with 5x10¹⁷ phage particles displaying mimotopes of *H. contortus* peptides - On day 28 all groups were orally challenged with 2500 infective *H. contortus* infective larvae. All groups demonstrated similar results (P>0.05) by the Kruskall-Wallis test.

<table>
<thead>
<tr>
<th>Day of experiment</th>
<th>GAPDH</th>
<th>Dim 1</th>
<th>Control</th>
</tr>
</thead>
<tbody>
<tr>
<td>Day 0</td>
<td>10.64 ± 2.90</td>
<td>11.90 ± 2.92</td>
<td>11.03 ± 4.05</td>
</tr>
<tr>
<td>Day 14</td>
<td>14.50 ± 2.23</td>
<td>11.66 ± 3.82</td>
<td>13.37 ± 3.03</td>
</tr>
<tr>
<td>Day 28</td>
<td>11.75 ± 2.42</td>
<td>13.52 ± 3.97</td>
<td>13.20 ± 2.02</td>
</tr>
<tr>
<td>Day 44</td>
<td>12.56 ± 2.68</td>
<td>14.73 ± 1.68</td>
<td>14.28 ± 3.05</td>
</tr>
<tr>
<td>Day 50</td>
<td>13.89 ± 1.96</td>
<td>14.41 ± 2.06</td>
<td>13.10 ± 0.81</td>
</tr>
<tr>
<td>Total leukocytes (x 10⁶/µL)</td>
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<td></td>
<td></td>
</tr>
<tr>
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<td>11.85 ± 2.06</td>
<td>12.00 ± 1.48</td>
<td>12.12 ± 1.42</td>
</tr>
<tr>
<td>Day 14</td>
<td>12.20 ± 2.14</td>
<td>10.97 ± 0.28</td>
<td>12.12 ± 1.38</td>
</tr>
<tr>
<td>Day 28</td>
<td>11.07 ± 1.96</td>
<td>10.70 ± 2.09</td>
<td>12.50 ± 1.58</td>
</tr>
<tr>
<td>Day 44</td>
<td>11.37 ± 1.15</td>
<td>12.47 ± 0.75</td>
<td>12.27 ± 1.61</td>
</tr>
<tr>
<td>Day 50</td>
<td>12.05 ± 2.04</td>
<td>10.92 ± 1.90</td>
<td>12.57 ± 1.19</td>
</tr>
<tr>
<td>Lymphocytes (%)</td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>Day 0</td>
<td>58.75 ± 3.86</td>
<td>59.50 ± 4.72</td>
<td>57.50 ± 2.38</td>
</tr>
<tr>
<td>Day 14</td>
<td>68.00 ± 8.16</td>
<td>64.50 ± 6.35</td>
<td>66.00 ± 6.22</td>
</tr>
<tr>
<td>Day 28</td>
<td>61.00 ± 4.08</td>
<td>65.00 ± 0.82</td>
<td>61.50 ± 1.91</td>
</tr>
<tr>
<td>Day 44</td>
<td>59.25 ± 3.30</td>
<td>53.75 ± 0.50</td>
<td>56.75 ± 6.95</td>
</tr>
<tr>
<td>Day 50</td>
<td>64.25 ± 3.77</td>
<td>62.00 ± 6.22</td>
<td>60.50 ± 2.65</td>
</tr>
<tr>
<td>Neutrophil (%)</td>
<td></td>
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<td></td>
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<tr>
<td>Day 0</td>
<td>36.75 ± 3.30</td>
<td>36.75 ± 3.77</td>
<td>39.00 ± 2.44</td>
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<tr>
<td>Day 14</td>
<td>26.75 ± 7.80</td>
<td>31.00 ± 5.35</td>
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</tr>
<tr>
<td>Day 28</td>
<td>34.25 ± 3.59</td>
<td>31.00 ± 1.56</td>
<td>32.75 ± 1.72</td>
</tr>
<tr>
<td>Day 44</td>
<td>36.00 ± 3.37</td>
<td>41.25 ± 1.73</td>
<td>37.00 ± 5.71</td>
</tr>
<tr>
<td>Day 50</td>
<td>30.75 ± 2.75</td>
<td>32.75 ± 5.44</td>
<td>34.50 ± 2.38</td>
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<tr>
<td>Eosinophil (%)</td>
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<td></td>
</tr>
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<td>1.50 ± 0.58</td>
</tr>
<tr>
<td>Day 14</td>
<td>2.25 ± 0.50</td>
<td>2.25 ± 0.50</td>
<td>1.75 ± 0.96</td>
</tr>
<tr>
<td>Day 28</td>
<td>1.75 ± 0.50</td>
<td>1.75 ± 0.96</td>
<td>2.75 ± 1.26</td>
</tr>
<tr>
<td>Day 44</td>
<td>1.75 ± 0.50</td>
<td>1.50 ± 0.58</td>
<td>2.25 ± 0.50</td>
</tr>
<tr>
<td>Day 50</td>
<td>2.50 ± 0.58</td>
<td>1.75 ± 0.96</td>
<td>1.75 ± 0.50</td>
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<tr>
<td>Monocytes (%)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Day 0</td>
<td>1.50 ± 1.00</td>
<td>1.75 ± 0.50</td>
<td>1.50 ± 0.58</td>
</tr>
<tr>
<td>Day 14</td>
<td>1.50 ± 0.58</td>
<td>1.25 ± 0.50</td>
<td>1.75 ± 0.50</td>
</tr>
<tr>
<td>Day 28</td>
<td>2.00 ± 0.82</td>
<td>1.50 ± 0.58</td>
<td>2.00 ± 0.82</td>
</tr>
<tr>
<td>Day 44</td>
<td>2.00 ± 0.82</td>
<td>1.75 ± 0.50</td>
<td>1.75 ± 0.96</td>
</tr>
<tr>
<td>Day 50</td>
<td>1.75 ± 0.50</td>
<td>1.50 ± 0.58</td>
<td>1.25 ± 0.50</td>
</tr>
<tr>
<td>Basophils (%)</td>
<td></td>
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<td></td>
</tr>
<tr>
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<td>0.25 ± 0.50</td>
<td>0.50 ± 0.58</td>
<td>0.50 ± 0.58</td>
</tr>
<tr>
<td>Day 14</td>
<td>1.25 ± 0.96</td>
<td>0.75 ± 0.96</td>
<td>0.50 ± 0.58</td>
</tr>
<tr>
<td>Day 28</td>
<td>0.75 ± 0.50</td>
<td>0.50 ± 0.58</td>
<td>0.00 ± 0.00</td>
</tr>
<tr>
<td>Day 44</td>
<td>0.50 ± 0.58</td>
<td>0.75 ± 0.96</td>
<td>1.50 ± 0.58</td>
</tr>
<tr>
<td>Day 50</td>
<td>0.25 ± 0.50</td>
<td>1.50 ± 0.58</td>
<td>1.25 ± 0.96</td>
</tr>
</tbody>
</table>

**Figure 2.** Mean levels of plasma IgG of 25 adult sheep naturally infected with *Haemonchus contortus*. The serum samples were analyzed against two chemically-synthesized peptides: (a) Glyceraldehyde-3-phosphate dehydrogenase (GAPDH) and (b) Disorganized muscle family member (Dim 1). These synthetic peptides have the same sequences (7 amino acids) displayed on two clones (6 and 17, respectively) selected by biopanning.
Phage display to map *Haemonchus contortus* mimotopes

Discussion

This is the first study using phage display to select *H. contortus* peptide mimotopes. This technology allowed mimotope selection for displayed peptides, using polyclonal serum of sheep naturally infected with *H. contortus*. The molecules bound to pathogen antibody are antigenic mimotopes of the pathogen (GAZARIAN et al., 2011). The ability of selected peptides to act as antigenic mimotopes was confirmed by serum analyzes against chemical peptides (GAPDH and/or Dim 1). We observed that IgG of animals naturally infected were able to recognize and bind to these peptides. The best results were for GAPDH, where only one sample had an IgG titre less than the cut-off.

In this study, after four selection rounds of biopanning, the bioinformatic analysis revealed two phage clones containing peptides with homology to GAPDH (clone 6) and Dim 1 (clone 17) proteins.
of *H. contortus*. Researchers have analyzed the immunogenic potential of many *H. contortus* molecules and immunoproteomic studies have already demonstrated that 23 immunogenic proteins are shared between males and females of *H. contortus*. These molecules include homologous of Dim 1 and GAPDH, among others (YAN et al., 2010).

GAPDH and Dim 1 are important vaccine candidates. In addition to their immunogenic potential, they perform important functions at cellular level. GAPDH is an enzyme crucial for energy production, in glycolysis and glycogenogenesis (NICHOLLS et al., 2012). It has also been related to other intracellular (SIROVER, 2011) and extracellular (GÓMEZ-ARREAZA et al., 2014) functions. According to Han et al. (2012), this molecule is a major therapeutic candidate for vaccines and a target for chemotherapy treatment against several parasites. Dim 1 is a structural protein located in the muscle cell membrane region, around and between dense bodies. It performs a key role in stabilizing the thin filament components of the sarcomere (YAN et al., 2013). Thus, an immunological response generated against these molecules might protect the animals against *H. contortus* infection.

In the in vivo assay, animals of both immunized groups GAPDH and Dim 1 presented IgG titers against M13 phage. The IgG increase followed the same pattern in both GAPDH and Dim 1 groups, with a peak after the last immunization. However, despite the sequence similarities to the immunogenic molecules of *H. contortus* (YAN et al., 2010), the peptides displayed by the phage did not promote specific IgG titers. It is important to note that the sequence displayed on the phage surface consisted of only 7 amino acids in linear conformation, being a small peptide compared to the full phage surface. Cui et al. (2013) observed significant immune responses and protection against *T. spiralis* infection in mice immunized with phage. However, they used a recombinant phage system, which displayed a 202-aa polypeptide.

To date, sheep immunization with phage displaying mimotopes has been reported only for *F. hepatica*. For this, animals were immunized at week 0 and 2 with $1 \times 10^{13}$ phage particles expressing 7-aa mimotopes and challenged with 300 metacercariae, 4 weeks after immunization. The mean worm burden was reduced by 47.61% and the egg viability from 58.92 to 82.11%. The animals produced specific antibodies after the metacercarialie challenge (VILLA-MANCERA et al., 2008).

Another study with a similar design had sheep receiving $1 \times 10^{13}$ phage particles (12-aa mimotopes) and challenged with 300 metacercariae 4 weeks later. The worm burdens and FEC were reduced by 51.7 and 45.7%, respectively (VILLA-MANCERA & MÉNDEZ-MENDOZA, 2012). In our work, the animals were immunized with phage expressing mimotopes of GAPDH or Dim 1 (7aa) but there was no reduction in FEC or worm counts, indicating the absence of a protective effect. A DNA vaccine encoding Dim 1 antigen against *H. contortus* reduced the FEC and worm counts of goats by 47.5 and 51.1% respectively (YAN et al., 2013). Similarly, a GAPDH DNA vaccine reduced FEC and worm count by 34.9 and 37.7%, respectively (HAN et al., 2012). These studies support the immunogenic potential of these molecules.

Eosinophils also have an important function in immune responses against helminth infections, and are often associated with resistance to parasite infections in sheep (PFEFFER et al., 1996; PATNODE et al., 2014). In our work, eosinophils and the other leukogram parameters did not reveal any difference throughout the experimental period. Yan et al. (2014) evaluating a *H. contortus* actin DNA vaccine in goats failed to detect differences in the eosinophil, neutrophil, basophil and monocyte counts before L3 challenge. However, after the challenge of 5000 L3 infective *H. contortus*, only eosinophil counts were significantly increased.

Santos et al. (2014) tested a single infection with 4000 L3, and one week later the challenged groups presented significantly higher eosinophil numbers. However, there was a gradual decline in the average values, until the end of the trial. In the same experiment, they observed an increase in the eosinophil average in serially infected groups (3 times a week, from day 0 – 25 of the trial with 500 L3). Values significantly higher than the control group were observed on days 14, 21, 25, 28, and 32 with the peak of eosinophil counts on day 25. They concluded that the animals developed this high protection after being consecutively challenged with *H. contortus*. The authors reported that immunity against the parasite was not easily elicited other than after the prolonged infection. In our study, we used a single challenge infection with 2500 L3, and that may have been reflected in the failure to see an increase in eosinophil counts.

It is important to emphasize that the development of immunity to *H. contortus* antigens is complex and highly variable. It depends on the host-parasite interaction, which is influenced by many factors, following an intrinsic and extrinsic relationship (TAK et al., 2015; NISBET et al., 2016). In our trial, the two mimotopes selected by phage display failed to protect, and neither induced a distinct immune response in the animals. The peptides reacted with polyclonal antibodies in serum from sheep naturally infected with *H. contortus*, but were not immunogenic when carried by the M13 phage, even though the two selected clones had high sequence similarity to Dim 1 and GAPDH of *H. contortus*. Our data highlight the importance of new studies to evaluate specific peptides from parasites.

**Conclusion**

The phage display library was successfully used to identify *H. contortus* mimotopes. The selected clones revealed relatedness to Dim 1 and GAPDH of *H. contortus* proteins. The peptides were recognized by several sera from sheep naturally infected with *H. contortus*, indicating their potential use as a diagnostic. The sheep immunized with phage displaying the mimotopes did not develop protection against the parasite.

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