

Duality of β -glucan microparticles: antigen carrier and immunostimulants

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Abstract: Designing efficient recombinant mucosal vaccines against enteric diseases is still a major challenge. Mucosal delivery of recombinant vaccines requires encapsulation in potent immunostimulatory particles to induce an efficient immune response. This paper evaluates the capacity of β -glucan microparticles (GPs) as antigen vehicles and characterizes their immune-stimulatory effects. The relevant infectious antigen FedF was chosen to be loaded inside the microparticles. The incorporation of FedF inside the particles was highly efficient (roughly 85%) and occurred without antigen degradation. In addition, these GPs have immunostimulatory effects as well, demonstrated by the strong reactive oxygen species (ROS) production by porcine neutrophils upon their recognition. Although antigen-loaded GPs still induce ROS production, antigen loading decreases this production by neutrophils for reasons yet unknown. However, these antigen-loaded GPs are still able to bind their specific β -glucan receptor, demonstrated by blocking complement receptor 3, which is the major β -glucan receptor on porcine neutrophils. The dual character of these particles is confirmed by a T-cell proliferation assay. FedF-loaded particles induce a significantly higher FedF-specific T-cell proliferation than soluble FedF. Taken together, these results show that GPs are efficient antigen carriers with immune-stimulatory properties.

Keywords: β -glucan microparticles, FedF, antigen delivery vehicle, immunostimulants

Introduction

Mucosal vaccination is the most efficient way to protect human and animals against enteric infections. However, the development of effective mucosal vaccine subunits encounters multiple challenges, such as antigen damaging and limited immunogenicity.¹⁻⁴ The encapsulation of antigens in microparticles is a promising approach to overcome these problems, as they can protect the antigens against degradation as well as carry potent adjuvants or immune modulators to enhance the immunogenicity.⁵⁻⁷ Frequently used particle systems for oral immunization are those based on poly(lactic-co-glycolic acid); however, low antigen encapsulation efficiency and antigen degradation during their formation limit their use in commercial vaccines.⁸ On the contrary, β -glucan microparticles (GPs) are emerging microparticles known for their safety, immunogenicity, and high antigen encapsulation efficiency.⁹⁻¹⁸ These promising antigen carriers are derived from the cell wall of *Saccharomyces cerevisiae* (Baker's yeast) and are composed of >85% β -1,3-D-glucan polymers (β -glucans), ~2% chitin, and <1% lipids and proteins, with the rest being mostly ash and moisture.¹⁵ The main component of these particles, the β -glucans, is a "microbe-associated molecular pattern", which is very interesting in vaccine development for its immunostimulating characteristics.^{19,20} We have previously reported that particulate β -glucans are mainly recognized by complement receptor 3 (CR3) on porcine innate immune cells and require the signaling molecule

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FAK to carry out their immunostimulatory responses.²¹ However, the receptor involved in the immunostimulating characteristics of GPs is not yet known. Many articles have described the strong potency of these GPs in eliciting durable immune responses in mice.^{9,10,14,22,23} Incubation of mouse bone marrow-derived dendritic cells with ovalbumin (OVA)-loaded GPs resulted in phagocytosis, upregulation of maturation markers, rapid proteolysis of OVA, and proliferation of OVA-reactive transgenic CD8⁺ OT-I and CD4⁺ OT-II T-cells.¹⁴ However, the immunological effect of these particles on porcine innate immune cells has not been investigated yet. Moreover, besides the incorporation of model antigens, no clinically relevant antigens were incorporated in these GPs. We have incorporated FedF, the tipadhesin of F18 fimbriae, inside the particles, since this antigen is one of the most important virulence factors of F18⁺ *Escherichia coli*. *E. coli* carrying the F18 fimbriae colonizes the small intestine of pigs and cause postweaning diarrhea (enterotoxigenic *E. coli* [ETEC]) and edema disease (Shiga toxin-producing *E. coli* [STEC]) in pigs, resulting in economic losses to the pig production industry.²⁴ The bacteria adhere to the intestinal epithelial surface with their fimbriae and produce enterotoxins, which induce secretion of electrolytes and water. Porcine ETEC strains can produce five fimbrial types (F4, F18, F41, F5, and F6), of which F4 and F18 are most frequently associated with ETEC-induced diarrhea.²⁵ The F18 fimbriae are polymeric proteins composed of several subunits: FedA is the structural subunit, while FedF is the tipadhesin mediating the binding of the bacteria to the epithelium. The FedF subunit of F18 fimbriae is located in a dedicated single-copy adhesin at the distal tip of the fimbriae.²⁶ Tipadhesins are usually composed of two immunoglobulin (Ig)-like domains: an N-terminal lectin or receptor binding module and a C-terminal pilin. As it needs a complementary β -strand from the FedE subunit, using donor strand complementation is necessary to achieve a stable recombinant expression of FedF.²⁷ To protect pigs against F18⁺ ETEC infections, local intestinal immunity is required. However, oral immunization with F18 fimbriae does not elicit protection in a piglet model, probably due to the presence of immunodominant epitopes in the FedA subunit.²⁸ We reasoned that oral immunization of piglets with purified FedF subunit would enable induction of protective immunity, but only if FedF is encapsulated inside microparticles so as to deliver higher amounts of antigens to the gut-associated lymphoid tissue and to induce strong immune responses by mimicking pathogen dimensions.

So, we incorporated FedF inside GPs for oral administration to induce intestinal FedF-specific antibody responses.

In this study, we investigated the capacity of these GPs as both antigen vehicles and immunostimulants. Porcine immune cells were used as the translational model, because β -glucan recognition and signaling in porcine innate immune cells is closer related to humans than murine cells.^{21,29,30}

Experimental section

Generation of FedF

FedF was generated using the method of De Kerpel et al.²⁷ Briefly, the *fedF*₁₅₋₁₆₅ gene from F18 *E. coli* 107/86 strain was cloned in the pDEST14 vector under a T7 promoter. The gene was C-terminal truncated and then transformed into *E. coli* strain C43 (DE3). The C43 (DE3) cells were grown, induced with 1 mM isopropyl β -D-1-thiogalactopyranoside, and incubated for 2 hours. Cells were subjected to osmotic shock and the periplasmic extract was loaded onto a Source 3S column (Amersham plc, Amersham, UK) for cation exchange in 20 mM Tris (pH 7.5). Proteins were eluted with 140 mM NaCl and further purified using gel filtration on a Superdex-75HR column in 10 mM 4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid (pH 7.0) and 150 mM NaCl. Finally, the protein was dialyzed against 0.9% saline.

Development of GPs and protein encapsulation

To generate GPs, *S. cerevisiae* (Baker's yeast) was treated with alkali and acid and dried at 20°C–25°C following alcohol and acetone washing steps.^{14,15} The resulting GPs were hollow and porous biomimetic 2–5 μ m particles consisting of >85% β -1,3-D-glucan polymers (β -glucans), ~2% chitin, and <1% lipids and proteins, with the rest being mostly ash and moisture.¹⁵ Subsequently, these GPs were loaded with FedF as previously described.¹⁴ Briefly, 10 mg of dry GPs were swollen with 5 mg FedF dissolved in 0.9% saline (1.25 mL) for 2 hours at 4°C, followed by lyophilization. These dry GP-FedF preparations were reswollen with 300 μ L aqua dest to maximize FedF diffusion into the GPs. Upon lyophilisation, FedF was trapped inside the GPs by adding 350 μ L of 25 mg/mL transfer RNA (derived from torula yeast) in Tris, EDTA, and NaCl buffer (50 mM Tris HCl with 2 mM ethylenediaminetetraacetic acid and 0.15 M NaCl, pH 8.2) for 30 minutes at 50°C. To complete the complexation reaction, another 500 μ L of 10 mg/mL transfer RNA was added to the particles. The suspension was centrifuged, washed four times in 0.9% saline, and stored at –20°C (2.5 mg/mL).¹⁴ To calculate the amount of FedF trapped inside the GPs, the unbound FedF protein in the wash

fractions was measured by a bicinchoninic acid assay against a bovine serum albumin standard (0.05–1 mg/mL).

Western blotting

Incorporation of FedF inside the particles was confirmed using a protein gel electrophoresis after digesting the GPs with β -1,3-D-glucanase from *Helix pomatia* (Sigma-Aldrich Co., St Louis, MO, USA). FedF-loaded GPs were first incubated in β -glucanases (1 mg/mL) diluted in 150 mM sodium acetate buffer (Sigma-Aldrich Co.) for 24 hours at 37°C. Proteins, released in the supernatants, were separated via sodium dodecyl sulfate–polyacrylamide gel electrophoresis consisting of a 12% separating gel and a 4% stacking gel. The gels were prepared by dissolving 30% acrylamide in Tris (pH 8.8), 10% sodium dodecyl sulfate, 10% ammonium persulfate, and tetramethylethylenediamine. To load the samples, the proteins were mixed in equal volume ratios with the loading buffer containing β -mercaptoethanol and heated at 100°C for 5 minutes. Electrophoresis was performed at 28 mA for 1 hour. After separation, the proteins were transferred to a 0.45 μ m polyvinylidene fluoride membrane (GE Healthcare, Stockholm, Sweden) at 40 V for 2 hours. Next, the membrane was blocked (1 hour at room temperature) with 5% milk powder diluted in phosphate-buffered saline +0.1% Tween®20 (Sigma-Aldrich Co.) to prevent antibodies from nonspecifically binding to the membrane. Monoclonal anti-mouse FedF-specific antibodies (10 μ g) were added overnight to the blot, followed by horseradish peroxidase-conjugated rabbit anti-mouse IgG (1/2,000; Dako, Denmark A/S Glostrup, Denmark) for 1 hour at room temperature. The bands were visualized using enhanced chemiluminescence (ECL) Western blotting substrate (Thermo Fisher Scientific, Waltham, MA, USA) and ChemiDOC™ MP imaging system (Bio-Rad Laboratories Inc., Hercules, CA, USA).

Oxidative burst by porcine neutrophils

All animal experiments were approved by the animal care and ethics committee of the Faculty of Veterinary Medicine, Ghent University (Gent, Belgium; EC2013/62). Piglets (between 4 and 20 weeks old) were housed under standard conditions as blood donors. Peripheral blood was collected on heparin from the jugular vein of four pigs. Subsequently, neutrophils were isolated by density gradient centrifugation on a discontinuous Percoll gradient (68% and 75%; GE Healthcare) as described previously.³¹ To examine the immunostimulating characteristics of the particles, the ability of the GPs to induce an oxidative burst response by porcine neutrophils was investigated. The production of reactive oxygen

species (ROS) was measured by the chemiluminescence assay described by Donne et al³² with some modifications. Neutrophils were seeded in a 96-well plate at 2.0×10^5 cells/well in Roswell Park Memorial Institute (RPMI) without phenol red. The plates were incubated at 37°C for 2 hours in a humidified atmosphere with 5% CO₂ to allow the cells to adhere to the plastic surface. Subsequently, the supernatant was replaced by 175 μ L luminol (0.5 mM). After 5 minutes of background measurement at 37°C, 25 μ L of the indicated GPs (400, 200, or 100 μ g/mL) were added. Stimulation of the cells with phorbol myristate acetate (PMA; 50 μ g/mL) was used as a positive control. ROS production was then measured during 120 minutes in the integration mode. All stimulations were performed in duplicate. The ROS production is expressed as relative light units.

Determination of the β -glucan receptor in porcine neutrophils

To analyze the role of dectin-1 and CR3 in the immunostimulating characteristics of these GPs, the production of ROS by neutrophils was measured using the chemiluminescence assay (as already described) with some modifications. Neutrophils were seeded in a 96-well plate at 2.0×10^5 cells/well. The plates were incubated at 37°C for 2 hours in a humidified atmosphere with 5% CO₂ to allow the cells to adhere to the plastic surface. Then, the cells were pretreated with the β -glucan receptor inhibitors for 1 hour at 37°C. To inhibit dectin-1, laminarin (1 mg/mL) was added to the cells. To inhibit CR3-mediated responses, monoclonal antibodies (mAbs; mouse IgG₁ isotypes) against CD18 (5 μ g/mL) and CD11R3 (15 μ g/mL) (Bio-Rad Laboratories Inc.) were added. These antibodies were dialyzed against phosphate-buffered saline to remove sodium azide. An isotype-matched mAb control (IgG₁) was added to the cells to measure background inhibition. Subsequently, luminol was added to the cells (0.5 mM diluted in Hanks' balanced salt solution + Ca²⁺ Mg²⁺), and, after 5 minutes of background measurement at 37°C, 25 μ L of GPs (200 μ g/mL) was added. Stimulation of the cells with PMA (50 μ g/mL) was used as a positive control. ROS production was then measured after 120 minutes in the integration mode. All stimulations were performed in duplicate. ROS production is expressed as relative light units.

Generation of monocyte-derived dendritic cells

Peripheral blood mononuclear cells (PBMCs) were isolated by density gradient centrifugation on Lymphoprep (Nycomed Pharma AS, Thermo Fisher Scientific).

Furthermore, monocytes were enriched from the PBMCs to a purity of >95% by positive immunomagnetic bead selection (MACS; Miltenyi Biotec, Bergisch Gladbach, Germany) using the anti-SIRP α monoclonal antibody (clone 74-12-15a³³). These cells were seeded in a 24-well plate at a density of 0.5×10^6 cells/mL in monocyte-derived dendritic cell (MoDC) medium (Dulbeccos' Modified Eagle Medium supplemented with 10% fetal calf serum, 1% penicillin–streptomycin, recombinant porcine 1/200 GM-CSF³⁴ and 5 ng/mL IL-4). After 3 days incubation at 37°C in a humidified atmosphere at 5% CO₂, fresh medium supplemented with GM-CSF and IL-4 at the same concentration to cells was added to generate MoDCs as previously described.³⁵

Antigen presentation assay

To investigate whether incorporation of FedF inside the particles is necessary to induce FedF-specific cellular immune responses, a coculture of MoDCs and CD6⁺ T-cells was set up. So, T-cells isolated from pigs immunized intramuscularly with 250 μ g purified FedF in incomplete Freund's adjuvant were used.³⁶ CD6⁺ T-cells were enriched from PBMCs to a purity of >95% by positive immunomagnetic selection with the α -CD6 mAb (IgG₁, clone a38b2³⁷) and goat anti-mouse microbeads (MACS, Miltenyi Biotec). MoDCs were stimulated with 8 μ g FedF or FedF-loaded GPs (8 μ g FedF encapsulated) for 24 hours or left untreated (control). These stimulated MoDCs were subsequently cocultured at

titrated numbers with 1.0×10^5 autologous CD6⁺ T-cells for 5 days, and then the [3H] methyl-thymidine (1 μ Ci/well; Amersham plc) incorporation (18 hours) was measured using a β -scintillation counter (PerkinElmer Inc., Waltham, MA, USA).

Statistics

Data are presented as the mean \pm standard error of the mean. Data were analyzed with GraphPad Prism 5 (GraphPad Software, Inc., La Jolla, CA, USA) using the paired *t*-test for comparison of two data and using a repeated measures analysis of variance (post hoc: Tukey) for comparison of multiple data sets. Data were considered significant at $P < 0.05$.

Results and discussion

Efficient loading of FedF inside GPs

To assess if loading of FedF into GPs affects the antigenicity, we performed Western blotting. FedF was incorporated inside the GPs with a loading efficiency of $84.53\% \pm 2.89\%$, thereby confirming the high loading efficiency of the GPs (Figure 1A). Furthermore, we examined by gel electrophoresis the correct incorporation of FedF inside the particles. Therefore, we first destroyed the β -glucan particles by β -glucanases, after which the supernatants with the released antigens were loaded on a protein gel electrophoresis and detected by monoclonal FedF-specific antibodies (clone IMM04³⁸). Figure 1B illustrates the protein FedF in lane 1

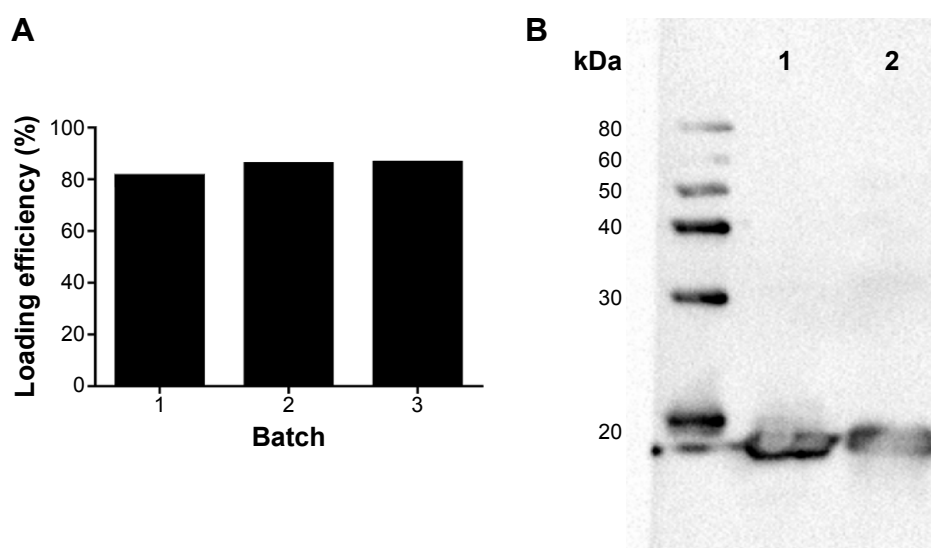


Figure 1 FedF was efficiently incorporated inside GPs.

Notes: (A) The loading efficiency of FedF inside GPs was determined by BCA. Therefore, the unbound FedF protein in the wash fractions was measured by BCA against a BSA standard (0.05–1 mg/mL). (B) FedF incorporation inside the GPs was analyzed by Western blotting after digesting the particles with β -glucanases. FedF was detected with specific monoclonal antibodies against FedF. Lane 1 illustrates FedF, while the FedF released from the GPs upon digestion is shown in lane 2.

Abbreviations: GPs, β -glucan microparticles; BCA, bicinchoninic acid assay; BSA, bovine serum albumin.

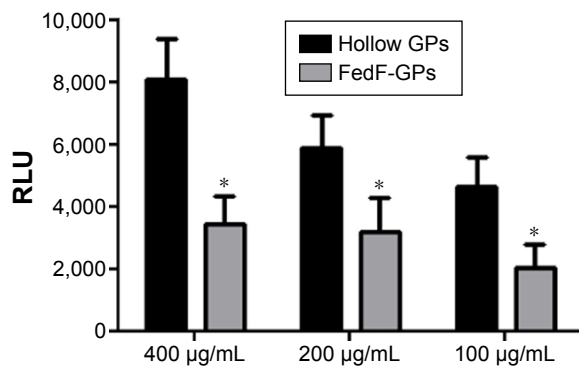


Figure 2 FedF-loaded GPs still possess their adjuvant function.

Notes: Neutrophils (2×10^5 cells) were stimulated with GPs at the indicated concentrations. ROS production was determined via chemiluminescence. Data are shown as the mean RLU \pm SEM of four pigs. * $P < 0.01$.

Abbreviations: GPs, β -glucan microparticles; RLU, relative light units; ROS, reactive oxygen species; SEM, standard error of the mean.

and the FedF released from the digested GPs in lane 2. As shown, both FedF and the released FedF migrate near the expected molecular weight of 18 kDa. No degradation is visible, indicating the correct incorporation of FedF inside the particles. On the basis of these results, we conclude that FedF is efficiently incorporated inside the particles.

FedF-loaded GPs still possess immune-stimulatory effects

To assess the effect of antigen loading on the inherent immunostimulating characteristics of the GP, we determined the oxygen radical production of porcine neutrophils after stimulation with antigen-loaded particles. Figure 2 demonstrates that FedF-loaded GPs possess immune-stimulatory effects. Interestingly, hollow GPs could stimulate ROS

production by neutrophils more than antigen-loaded particles, indicating that antigen loading partially masks the β -glucans of the particles from recognition by their receptors, dectin-1, and CR3.²¹

The immune-modulatory effects of GPs are mainly mediated by binding to the α -subunit of CR3

We further investigated which β -glucan receptor is involved in the particle-induced activation of porcine neutrophils. As shown in Figure 3, the GPs-induced ROS production by neutrophils was significantly inhibited in the presence of anti-CD11R3 as compared to its isotype-matched control. In contrast to the α -subunit of CR3, the ability of CD18-specific mAbs to block ROS production by porcine neutrophils was negligible (β -subunit of CR3). Interestingly, blocking dectin-1 by laminarin decreased ROS production as well, albeit not as much as CD11R3-specific mAbs. Although incorporation of antigens inside the particles seems to influence the recognition of GPs by dectin-1, CD11R3 is still the most important receptor for the GP-induced ROS production in porcine neutrophils.

T-cell proliferation is significantly increased when FedF is incorporated inside GPs

To assess whether GPs could serve as an immune stimulatory antigen delivery system, we investigated the FedF-specific T-cell proliferation after stimulation of MoDCs with FedF-GPs. As illustrated in Figure 4, FedF-loaded GPs promoted the antigen presentation capacity of MoDCs

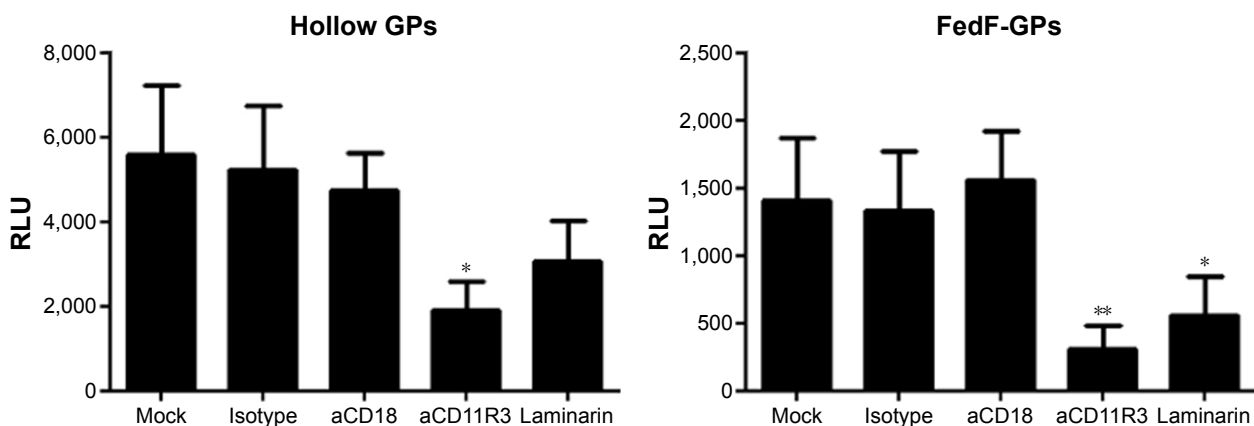


Figure 3 CR11R3 is involved in the recognition of GPs by neutrophils.

Notes: Neutrophils (2×10^5 cells) were incubated with laminarin (1 mg/mL), aCD18 (5 μ g/mL), aCD11R3 (15 μ g/mL), or isotype control (IgG; 15 μ g/mL). Subsequently, the cells were stimulated with 200 μ g/mL of the indicated GPs. ROS production was determined via chemiluminescence. Values shown are mean RLU \pm SEM of four pigs. Values of the negative control have been subtracted from the values represented on the graph. * $P < 0.05$; ** $P < 0.01$.

Abbreviations: IgG, immunoglobulin G; GP, β -glucan microparticle; RLU, relative light units; ROS, reactive oxygen species; SEM, standard error of the mean.

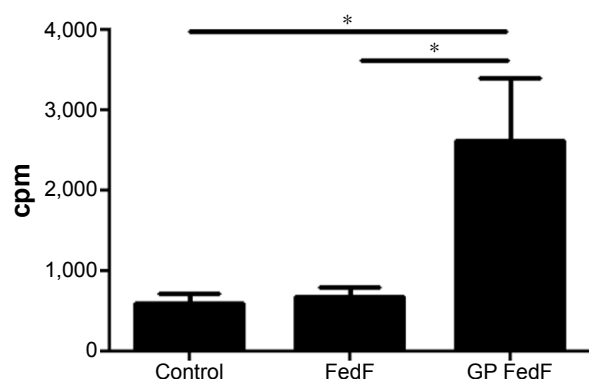


Figure 4 FedF-specific T-cell proliferation is significantly increased when FedF is incorporated inside GPs.

Notes: Encapsulation of FedF in GPs promotes antigen presentation to T-cells by MoDCs. Autologous CD6⁺ T-cells were cocultured for 5 days with immature, FedF (8 µg) or FedF-loaded GPs (8 µg FedF encapsulated)-stimulated MoDCs. The data represent the mean cpm ± SEM (n=4). The spontaneous proliferation of the CD6⁺ responder cells equaled 78 ± 14 cpm, while for ConA-stimulated cells, the proliferation amounted to 15,203 ± 5,623 cpm. Control: immature MoDCs. *P < 0.05.

Abbreviations: GPs, β-glucan microparticles; MoDCs, monocyte-derived dendritic cells; cpm, counts per minute; SEM, standard error of the mean.

to porcine CD6⁺ T-cells as compared to immature and FedF-stimulated MoDCs. Surprisingly, FedF stimulation of dendritic cells did not elicit T-cell proliferation, suggesting that FedF has an inhibitory effect on MoDCs. Clearly, the encapsulation of FedF inside immunostimulatory particles avoids this inhibitory effect and hence is necessary to induce T-cell proliferation.

Conclusion

GPs are emerging particles known for their high antigen-loading capacity and safety. In this study, we evaluated the incorporation of a clinically relevant infectious disease antigen FedF, the tipadhesin of F18 fimbriae, inside these particles. The results clearly demonstrate that FedF is incorporated intact inside the particles with a high loading efficiency (84.5%). Besides antigen carrier activity, these particles are able to stimulate the innate immune system by binding to the α-subunit of CR3. To confirm the duality of these particles, we demonstrated in a T-cell proliferation assay that FedF inside the particles is processed by MoDCs and subsequently presented to T-cells, resulting in their activation and proliferation. Taken together, FedF-loaded GPs are promising vaccine candidates in the protection of pigs against F18⁺ ETEC and STEC infections, and this study motivates exploring this vaccine in vivo.

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Disclosure

The authors report no conflicts of interest in this work.

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