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Short communication

Presence of antibodies against SARS-CoV-2 spike protein in bovine whey IgG enriched fraction

Satoshi Oshiro, Tatsuya Tada, Naeko Mizutani, Keiji Funatogawa, Jun-ichiro Sekiguchi, Masao Takahashi, Teruo Kirikae

Department of Microbiology, Juntendo University School of Medicine, Tokyo, Japan
Tochigi Prefectural Institute of Public Health and Environmental Science, Utsunomiya, Tochigi, Japan
Microbiology Research Division, Kohjin Bio Co., Ltd., Saitama, Japan
Aotearoa Co., Ltd., Tokyo, Japan

ABSTRACT

Bovine whey IgG enriched fraction contains antibodies against various human bacterial pathogens. It contains antibodies against some viral antigens, including human respiratory syncytial virus and influenza virus. We investigated whether the IgG enriched fraction has cross-reactivity with IgG antibodies against SARS-CoV-2 spike (S) and nucleocapsid (N) proteins. The full-length and partial-length SARS-CoV-2 S, N, a recombinant protein of the receptor binding domain (RBD) and nine peptides covering the receptor binding motif (RBM) of S were prepared. Direct enzyme-linked immunosorbent assays were conducted using these recombinant proteins and peptides as coating antigens and revealed the IgG enriched fraction contained antibodies against partial-length S [amino acids (aa) 177–512, 288–512, 348–578, 387–516 and 408–664], full-length N (aa 1–419) and partial-length N (aa 1–120, 111–220, 1–220 and 210–419), two RBD peptides, covering aa 427–446 and 502–520 of S, and recombinant RBD of S. These results indicate IgG enriched fraction contains antibodies against SARS-CoV-2.

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1. Introduction

Bovine milk products contain IgG against several human bacterial pathogens, and rotavirus that cause gastrointestinal tract infections (Ulfman, Leusen, Savelkoul, Warner, & van Neerven, 2018), and bovine IgG was reported to bind to human respiratory syncytial virus (RSV) and influenza virus (Hartog et al., 2014). Bovine colostrum preparations obtained from cows immunised with antigens of several human gastrointestinal tract infections was called “hyperimmunised milk” (Golay, Ferrara, Felber, & Schneider, 1990). It is characterised by high antibody activities against specified pathogens. Clinical trials demonstrated that immune cow colostrum shortened the duration of gastrointestinal tract infections (Ulfman et al., 2018). Second-generation milk products obtained from colostrum derived from healthy non-immunised pasture fed cows provided immunity against Salmonella infection in calves (Funatogawa et al., 2002; Griffiths, 1969; Royal, Robinson, & Duganzich, 1986). An immunoglobulin preparation from non-immunised cows contained high levels of antibodies and neutralising activity against verotoxin of Escherichia coli O157:H7 (Funatogawa et al., 2002; Lissner, Schmidt, & Karch, 1996). Several reports that indicate bovine IgG has antibodies against various bacterial antigens and activates the human immune system to repel pathogens have been reviewed (Ulfman et al., 2018). Bovine IgG fraction was reported to protect mice against food-borne infections with enterohaemorrhagicol E. coli O157:H7 and Salmonella enterica serovar Enteritidis (Funatogawa et al., 2019). This IgG fraction partially protected mice against respiratory tract infection with Mycobacterium avium (Funatogawa, Tada, Kuwahara-arai, Kirikae, & Takahashi, 2019). However, it is unclear whether bovine IgG recognises human viral pathogens except for rotavirus, RSV, and influenza virus.

The novel coronavirus, severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), has been causing a coronavirus disease (COVID-19) pandemic since 2019 (WHO, 2019). Sequence data analysis of human coronaviruses, including SARS-CoV-2 suggests they have an animal origin, especially bat (Cui, Li, & Shi, 2019). Here, we report IgG antibodies against SARS-CoV-2 spike protein...
were used. The cows were not administered insecticidal drugs, antibiotics, or growth hormones. The powder included (w/w); 84.7% protein, <8.4% fat, 7.0 mg g⁻¹ lactoferrin and 327.7 mg g⁻¹ immunoglobulin.

### 2. Material and methods

#### 2.1. Construction and purification of recombinant SARS-CoV-2 spike protein (S) and nucleocapsid protein (N)

A partial-length of SARS-CoV-2 S gene (2055 bp) and the full-length of SARS-CoV-2 N gene (1260 bp) were synthesised based on SARS-CoV-2 isolate 2019-nCoV WHU01, complete genome (accession no. MN988668). Five sequences of SARS-CoV-2 S gene (529–1536 bp, 862–1536 bp, 1042–1734 bp, 1159–1548 bp and 1222–1992 bp), corresponding to amino acid (aa) 177–512, 288–512, 348–578, 387–516 and 408–664, and five sequences of SARS-CoV-2 N gene (1–360 bp, 330–660 bp, 1–660 bp, 628–1260 bp and 1–1280 bp), corresponding to amino acid (aa) 1–120, 111–220, 1–220, 210–419 and 1–419, were cloned into pET28a expression vectors (Novagen, Inc., USA) using primers listed in Table 1. The constructed plasmids were used to transform E. coli BL21-CodonPlus (DE3)-RIP (Agilent Technologies, USA). Recombinant SARS-CoV-2 proteins were purified using Ni-NTA agarose, according to the manufacturer's instructions (Qiagen, Germany), and dissolved at 3 mg mL⁻¹ in phosphate-buffered saline (PBS). A recombinant protein covering the RBD of SARS-CoV-2 spike protein was purchased from Sino Biological Inc, USA. For use in direct enzyme-linked immunosorbent assays (ELISAs), these recombinant proteins were diluted to 10 μg mL⁻¹ in PBS-0.1% Tween 20 and used as coating antigens.

#### 2.2. Peptides of SARS-CoV-2 S

Nine peptides of SARS-CoV-2 S, corresponding to aa 382–401, 397–416, 427–446, 442–461, 457–476, 472–491, 487–506 and 502–520, were synthesised by Eurofins Genetics Inc, Japan. These peptides were dissolved at 3 mg mL⁻¹ in distilled water free of endotoxin (LONZA, Switzerland) and diluted to 10 μg mL⁻¹ in PBS containing 0.1% Tween 20 for use as coating antigens in ELISAs and as inhibitors in competitive inhibition ELISA.

#### 2.3. Bovine whey IgG enriched fraction

Bovine whey IgG enriched fraction (IgG₃0/₄/C₀) was obtained from milk of pasture fed, non-immunised healthy New Zealand cows by New Zealand Dairy Group by centrifugation and dissolved at 3 mg mL⁻¹ in phosphate-buffered saline (PBS). A bovine IgG enriched fraction (0.3 g C₀) was obtained from milk of pasture fed, non-immunised healthy New Zealand cows by New Zealand Dairy Group by centrifugation and dissolved at 3 mg mL⁻¹ in phosphate-buffered saline (PBS) and then incubated for 1 h at room temperature, then washed three times with PBS containing 0.1% Tween 20 between each of the following steps: blocking the wells with SuperBlock™ Blocking Buffer in PBS (Thermo Scientific, MA, USA) (150 μL well⁻¹) for 30 min at room temperature; addition of 100 μL well⁻¹ of IgG (0.003, 0.03, 0.3, 3 and 30 μg mL⁻¹ in PBS containing 0.1% Tween 20) bovine IgG enriched fraction for 1 h at room temperature; addition of 100 μL well⁻¹ of peroxidase-conjugated anti-bovine IgG (whole molecule) (1:10,000 dilution) (Sigma-Aldrich, St Louis, MO, USA) for 30 min at room temperature; addition of 50 μL TMB Peroxidase ELA Substrate Kit (Bio-Rad, Hercules, CA, USA) for 10 min. After 10 min incubation, 50 μL of 1% sulphuric acid was added to stop the peroxidase reaction. Absorbance at 450 nm and 620 nm as reference were measured using infinite F50 microplate readers (TECAN, Switzerland).

Competitive inhibition ELISAs were performed by incubating the bovine IgG enriched fraction (0.3 μg mL⁻¹) with one of the three synthesized peptides covering RMB of S (aa 382–401, 427–446 and 502–520) at concentrations of 0.001, 0.01, 0.1, and 10 μg mL⁻¹ in PBS containing 0.1% Tween 20 in 1.5-mL tubes overnight at 4 °C. The remaining free IgG against SARS-CoV-2 S was measured by direct ELISA, using plates coated with recombinant SARS-CoV-2 S (aa 288–512).

#### 2.5. Endotoxin assay

Endotoxin concentrations in the solutions of the recombinant proteins and peptides were quantitatively measured using QCL-1000 Limulus Amebocyte Lysate, according to the manufacturer's instructions (LONZA, Switzerland). Solutions of recombinant proteins and peptide (3 μg mL⁻¹ each) were diluted to 10 μg mL⁻¹ in endotoxin-free water and endotoxin was measured.
3.2. IgG enriched fraction containing antibodies against SARS-CoV-2

Reactivity against SARS-CoV-2 S was higher than that against SARS-CoV-2 N (Fig. 2). The IgG enriched fraction strongly recognised the 288–512 region compared with other regions of SARS-CoV-2 S (Fig. 2). There were few differences in reactivities against recombinant proteins from SARS-CoV-2 S and N among the two lots tested, except for reactivities against SARS-CoV-2 N (aa 1–220), i.e., the lot prepared in 2018 have more reactivity against SARS-CoV-2 N (aa 1–220) than that in 2019 (Fig. 2). The results of ELISA indicate that the bovine IgG enriched fraction contained IgG antibodies against SARS-CoV-2 S and N.

3.3. Epitope mapping of RBD in SARS-CoV-2 spike protein

ELISA using a recombinant RBD protein revealed that the IgG enriched fraction contained antibodies against the RBD in the S protein of SARS-CoV-2 (Fig. 3). ELISA using peptides covering the RBD revealed that this fraction contained IgG antibodies against two peptides (aa 427–446 and 502–520), indicating that the IgG enriched fraction contained antibodies against two epitopes of the RBD in S protein (Fig. 3).

3.4. Competitive inhibition ELISA

The ability of two peptides (aa 427–446 and 502–520) to inhibit the binding of IgG antibodies present in the enriched fraction with the S protein of SARS-CoV-2 was evaluated by competitive inhibition ELISA. Both peptides dose-dependently inhibited the binding of two lots of bovine IgG enriched fraction with the S protein of SARS-CoV-2 (aa 288–512) (Fig. 4). The 50% inhibitory concentrations (ID₅₀) of peptide (aa 502–520) (0.09 and 0.11 μg ml⁻¹, respectively) were slightly lower than those of peptide (aa 427–446) (0.20 and 0.24 μg ml⁻¹, respectively) (Fig. 4). A third peptide (aa 382–401), with which the IgG enriched fraction did not react, was unable to inhibit the binding of the two lots in ranges of 0.001–10 μg ml⁻¹ (Fig. 4).

3.5. Endotoxin levels

The solutions of all the recombinant SARS-CoV-2 S and N proteins (10 μg ml⁻¹) were found to be contaminated with significant concentrations of endotoxin, ranging from 7.7 to 1108 EU ml⁻¹ (0.77–110.8 EU μg⁻¹) (Fig. 5). In contrast, endotoxin was not detected in any of the solutions of the three SARS-CoV-2 S peptides (aa 382–401, 427–446 and 502–520) tested (Fig. 5). These results...
together, with those of direct and indirect ELISA indicated that bovine IgG directly and specifically binds to the two peptides (aa 427–446 and 502–520), although we could not exclude that possibility that endotoxin disrupted the binding of bovine IgG to recombinant SARS-CoV-2 S and N proteins.

3.6. Bovine IgG enriched fraction

It is unlikely these pasture-fed healthy New Zealand cows were exposed to or naturally immunised against SARS-CoV-2. The two lots of bovine IgG enriched fraction prepared in 2019 and 2018 were used (2a and 2b, respectively): 0.003 μg mL⁻¹, 0.03 μg mL⁻¹, 0.3 μg mL⁻¹, 3 μg mL⁻¹, 30 μg mL⁻¹. A picture of a representative ELISA result is shown in 2c.
SARS-CoV-2 is an intercellular protein that has a multifunctional RNA-binding protein for viral RNA transcription and replication (Kang et al., 2020; Lan et al., 2020). SARS-CoV-2 N would have relatively lower antigenicity against cows. Accordingly, the bovine IgG enriched fraction may show less potent activity against SARS-CoV-2 N than SARS-CoV-2 S.

**Author contributions**

SO and NM created the research data. SO and TT wrote the draft of the manuscript. All authors read, made significant edits to the first version, and approved the final manuscript.

**Conflict of interest**

S. J. works for Kohjin Bio Co., Ltd.
M.T. works for Aotearoa Co., Ltd.

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prepared in November 2018 and August 2019, respectively, which predate the emergence of SARS-CoV-2 in December 2019 in China. But it cannot be excluded that the cows had been exposed to some unknown coronavirus that share the immunogenicity of SARS-CoV-2.

The present study showed that the bovine IgG enriched fraction binds specifically to the RBD of SARS-CoV-2 S protein, indicating that the IgG enriched fraction would have potential to neutralise SARS-CoV-2. SARS-CoV-2 S forms homotrimers and uses angiotensin-converting enzyme 2 (ACE2) to enter cell (Walls et al., 2020).

SARS-CoV-2 S murine polyclonal antibodies potently inhibited SARS-CoV-2 mediated entry into cells, indicating that cross-neutralising antibodies targeting conserved S epitopes may be induced by vaccination (Walls et al., 2020). A human monoclonal antibody bound a conserved epitope of spike RBD and neutralised SARS-CoV-2 (Wang et al., 2020).

**Fig. 4.** Competitive inhibition ELISA of the bovine IgG enriched fraction (IgG 0.3 μg mL⁻¹), incubated with one of three peptides of S protein of SARS-CoV-2 (aa 427–446; aa 502–520) at concentrations 0.001, 0.01, 0.1, 1, and 10 μg mL⁻¹; the remaining free IgG against the S protein of SARS-CoV-2 was assayed by direct ELISA, using plates coated with the peptide corresponding to aa 288–512 of S protein of SARS-CoV-2. Two lots of bovine IgG enriched fraction prepared in 2019 (●) and 2018 (■) were tested.

**Fig. 5.** Endotoxin levels in solutions of the recombinant proteins and peptides measured quantitatively using QCL-1000 Limulus Amebocyte Lysate; each solution contained 10 μg of recombinant protein or peptide (ND, not detected).