Enhanced Membrane-tethered Mucin 3 (MUC3) Expression by a Tetrameric Branched Peptide with Conserved TFLK Motif Inhibits Bacteria Adherence

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*Running title: TFLK-containing 10-mer BP enhances MUC3 expression

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Background: A tetrameric branched peptide (BP) containing TFLK-motif was designed to induce MUC3 expression and inhibit bacteria adherence. Results: TFLK-containing 10-mer BP induced MUC3 expression and dramatically inhibited bacteria adherence to the HT-29-Gal cells. Conclusion: This peptide acts through transcriptional mechanisms. Significance: This peptide has potential therapeutic value against gastrointestinal infection.

SUMMARY

We investigated whether a synthetic tetrameric branched peptide based on the conserved TFLK-motif from M-SAA3 is more efficient than the monomeric peptide at up-regulating MUC3 expression and examined the possible mechanism(s) and biological significance of this process. We used standard solid-phase methods to synthesize a tetrameric branched peptide (sequence GWLTFLKAAG) containing a trilysine core, termed the TFLK-containing 10-mer BP. The aberrant expression of transcription factors (TF) was analyzed using a transcription factor protein/DNA array. MUC3 and relevant TFs were detected using real-time PCR and (or) Western blots. The luciferase assay, EMSA and ChIP assays were used to analyze the activity of the human MUC3 promoter. The bacterial adherence assay was used to evaluate the in vitro inhibition of enteropathogenic Escherichia coli (EPEC) or enterohemorrhagic Escherichia coli serotype O157:H7 (EHEC O157:H7) adherence to HT-29-Gal cells after treatment with the TFLK-containing 10-mer BP. In HT-29-Gal cells, the TFLK-containing 10-mer BP induced higher levels of MUC3 expression than the M-SAA3 derived N-terminal 10-mer monomeric peptide, and MUC3 expression was activated through transcriptional mechanisms, including the induction of multiple transcription factors and further binding with their cis-elements between nucleotides -242 and -62 within MUC3 promoter. Interestingly, the TFLK-containing 10-mer BP dramatically inhibited EPEC and EHEC O157:H7 adherence to the HT-29-Gal cells compared with the controls. This finding
suggests a potential therapeutic use for this peptide to prevent gastrointestinal infection.

Mucosal surfaces are protected against xenobiotics and pathogenic microorganisms by a variety of innate and adaptive mechanisms. For example, mucins prevent the noxious interaction of the epithelial cells with microbial pathogens and toxic chemicals by providing a physicochemical barrier through specific mucin-microorganism interactions and cell-signaling regulation (1, 2). By limiting the adherence or invasion of microorganisms to the epithelial cell surface, mucin effectively hampers the ability of the bacteria and viruses to colonize and invade the cells, blocks the spread of the pathogens along the mucosal surfaces, and limits the amount of microbial-produced toxins that reaches the mucosal cells (3). Intestinal mucins provide protection against the adherence of enteric pathogens by steric hindrance or through specific bacterial- or viral-binding domains. The attachment of microbes to intestinal mucins depends on the composition and quantity of mucins expressed and secreted and on the motility and fluid flow of the intestinal tract (4). If the rate of pathogenic microbial adherence and colonization exceeds the rate of removal, disease is a likely result. These events contribute to the development of localized intestinal necrosis that promotes further growth and the translocation of pathogenic microbes that can lead to shock, sepsis, and sometimes death. Human MUC3 and rodent Muc3 are membrane-bound mucins that are moderately expressed in the colon but are found abundantly in both goblet cells and enterocytes in the small intestine (5). MUC3 binds numerous enteropathogenic bacteria and viruses and prevents their attachment to the intestinal cell surfaces (4). Recombinant Muc3 and recombinant human MUC17 mucins stimulate cell migration in human colon cell lines and protect cells from apoptosis (6, 7). Although known inducers of MUC3 expression are limited, a previous in vitro study demonstrated that co-incubation of the probiotic bacteria Lactobacillus spp. with human intestinal epithelial cells induced MUC3 expression and inhibited enteropathogenic Escherichia coli (EPEC) adherence (8, 9).

Numerous studies have shown that compared with formula-fed infants, infants fed colostrum are approximately 7–10 times less likely to suffer gastrointestinal diseases (10, 11). Colostrum is a complex source of nutrients, immune factors, and bioactive substances that facilitates the successful transition of the mammalian neonate to extrauterine life (12, 13). However, despite the considerable number of studies investigating the stimulatory and inhibitory effects of colostrum, the numerous colostrum components and the underlying mechanisms that benefit the neonate are unknown.

In vitro studies demonstrated that the human mammary-associated serum amyloid A3 (M-SAA3) derived N-terminal peptide increased MUC3 transcriptional expression in human intestinal epithelial cells and decreased EPEC adherence to these cells (14-16). M-SAA3 is secreted at highly elevated levels in colostrum. Each of these colostrum-derived M-SAA3 proteins contains a conserved TFLK motif positioned within the first eight residues of the N-terminus, regardless of the species. This TFLK motif has not been reported for any serum-derived serum amyloid A isoform in mammals and appears to be in part responsible for the bioactivity that results in the up-regulation of MUC3 transcriptional expression by human intestinal cells. The discovery of elevated M-SAA3 levels in the colostrum and the reduction of gastrointestinal diseases in infants fed colostrum implies a possible role for M-SAA3 in promoting the health of the neonate.
Recently, multimeric peptides have been described in the literature as an obvious solution to increase peptide size and affinity while maintaining the original amino acid sequence (17-19), and this approach is a key strategy to increase recognition surfaces and/or stability. Increasing the peptide size also increases the difficulty of identifying peptides with high affinity and selectivity; however, multimeric peptides comprising simple branched structures and a short arm may offer an optimal compromise between molecular size and the ease of preparation, yield and purity of the final product. This strategy of combinatorial chemistry prompted us to explore whether the multimeric branched peptide based on the conserved TFLK-motif from M-SAA3 induces increased expression of MUC3 in intestinal epithelial cells and exerted a more significant effect.

In this study, we demonstrate that compared to the M-SAA3 derived N-terminal 10-mer monomeric peptide containing the TFLK motif (residues 1-10, sequence GWLTFLKAAG), the TFLK-containing 10-mer BP stimulates higher levels of human MUC3 expression in intestinal epithelial cells and dramatically inhibits the adherence of EPEC and EHEC O157:H7. Furthermore, we explored the mechanism(s) of transcriptional regulation of human MUC3 mucin by the TFLK-containing 10-mer BP in HT-29-Gal cells.

**EXPERIMENTAL PROCEDURES**

*Peptide synthesis* — In this study, the 10-mer branched peptide based on the conserved TFLK-motif from M-SAA3 (referred as TFLK-containing 10-mer BP) was synthesized using the solid phase method at the Chinese Peptide Company (Hangzhou, China). The introduction of three symmetrically protected lysines into the peptide produces a trilysine core, which is required for producing tetramERICALLY branched peptides. This method utilizes standard solid-phase methods and an Applied Biosystems (Foster City, CA) Model 430A synthesizer. The syntheses were performed on a 0.25-mmol scale on a Wang resin using the N-(9-fluorenyl) methoxycarbonyl (Fmoc) synthetic scheme. The peptides were purified using analytical and preparative HPLC columns packed with C18-bonded silica gel and were characterized using amino acid compositional analysis and mass spectrometry. The M-SAA3 derived N-terminal 10-mer monomeric peptide was synthesized simultaneously and used as a control.

*Cell line and culture* — To examine MUC3 expression, the human intestinal epithelial cell line HT-29 (ATCC) was cultured in glucose-free McCoy’s 5a medium (Gibco-BRL, Bethesda Research Laboratories, Gaithersberg, MD, USA). The culture medium was supplemented with 5 mM galactose, 10% heat-inactivated qualified FBS, 100 U/mL penicillin G, 100 mg/mL streptomycin sulfate, and 0.25 mg/mL amphotericin B (Gibco-BRL). The glucose-free and galactose-substituted cell culture medium induces the differentiation of the HT-29 cells, referred to as HT-29-Gal cells (a mature and differentiated form of the HT-29 cell line) (20, 21). The synthetic branched peptide was added to the HT-29-Gal cells in culture for 0.5, 1, or 3 h. The cultures were incubated at 37 °C in a humidified atmosphere with 5% CO₂. The cells were passaged following the release of the adherent cells using trypsin-EDTA (Gibco-BRL).

*Real-time PCR analysis* — The HT-29 cells were grown to 85% confluency in culture flasks containing either glucose-containing culture medium or the glucose-free, galactose-containing medium. The synthetic branched peptides were added to the culture medium and were incubated for 1 h at 37 °C in a humidified atmosphere with 5% CO₂. The total RNA was isolated from the
cells using Trizol (TaKaRa Biotechnology Co. Ltd., Dalian, China). The RNA was subjected to real-time PCR, or was stored at 70°C. The mRNA was reversed transcribed into cDNA using the PrimeScript RT reagent kit (TaKaRa Biotechnology Co. Ltd., Dalian, China). To determine the fold-changes in the expression of each gene, real-time PCR was performed using the first strand cDNA, the forward and reverse primers (supplementary Table 1), and the SYBR premix Ex Taq™ Green II (Takara, Japan). The alteration of the MUC3 mRNA was quantified using real time PCR (CFX 96, BioRad). The reaction and signal detection were measured using the BioRad CFX manager software (BioRad, USA). The expression levels were calculated as the relative expression ratio compared to the expression levels of the control β-actin and GAPDH genes. The real-time PCRs were performed independently in triplicate.

Western blotting — The cell pellets or nuclear proteins were extracted using NE-PER nuclear and cytoplasmic extraction reagents (Thermo Scientific Pierce Protein Research Products, USA), were lysed in SDS sample buffer and were subjected to SDS/PAGE (7.5 or 8% gels). Next, the proteins were transferred to PVDF membranes (Millipore, Bedford, MA, USA). The membranes were blocked for 1 h in 5% nonfat dry milk in TBST (TBS containing 0.05% Tween-20) and were hybridized at 4 °C overnight in TBST containing the following primary antibodies: anti-MUC3 (1:6,000, Abcam, Cambridge, UK), anti-Sp1 (1:1,500), anti-CDX2 (1:1,500), anti-CREB1 (1:1,000), or anti-USF1 (1:1500) (the anti-Sp1, anti-CDX2, anti-CREB1, and anti-USF1 antibodies were from Santa Cruz Biotechnology Inc., California, USA). Lamin B1 and GAPDH were detected using the anti-lamin B1 (1:5,000) and anti-GAPDH antibodies (1:5,000) (Santa Cruz Biotechnology Inc., California, USA) and were used as the internal controls. The membranes were washed three times in TBST for 5 min each, and the secondary detection was performed using 1:50,000 dilutions of HRP-conjugated goat anti-rabbit or goat anti-mouse antibodies. The membranes were washed three times for 10 min each. The blots were processed using the SuperWest Pico chemiluminescent reagent (Pierce, Rockford, IL, USA) and were analyzed using ChemiDox™ XRS with Image Lab™ software.

Transfections and luciferase assays — Fragments of the MUC3 5′-flanking sequence were amplified using PCR and were cloned into the luciferase reporter vector, pGL3-Basic (Promega, Madison, WI, USA). Briefly, primers with containing XhoI and Hind III adapters were used to amplify the MUC3 promoter sequence from intestinal tissue DNA. The primer pairs used to produce each promoter fragment are listed in supplementary Table 2. The products were ligated into the pGL3-Basic vector that was digested with XhoI and Hind III. The 5′ serial deletions of the 1,499-bp MUC3 promoter region were generated using the Erase-ABase system (Promega, Madison, WI, USA) in accordance with the manufacturer's recommendations. The plasmids for the transient transfections were purified using the EndoFree Plasmid Maxi kit (Qiagen, Valencia, CA). The day before the transfection, the cells were plated on 24-well plates at a density of 5×10⁴ cells per well. The MUC3 promoter-luciferase constructs were transfected into the cells using Lipofectamine™ 2000 (Invitrogen, Groningen, Netherlands). To normalize for the transfection efficiency, the cells were simultaneously cotransfected with a pRL-TK vector expressing the Renilla luciferase enzyme (pRL, Promega). The cells were harvested after 24 h, and the luciferase activity was measured using the Dual-Luciferase Reporter Assay System (Promega) using a single sample luminometer. The cells were also transfected with the pRL-TK vector and the
MUC3 activity is presented as the percentage of the pGL3-control activity. For the synthetic branched peptide stimulation, the cells were exposed to the peptide at a concentration of 50 μg/mL for one hour 12 h after transfection, and the stimulation was repeated three times. The differences between the untreated cells and the synthetic branched peptide-treated cells were determined using Student’s t test.

**Transcription factor (TF) protein/DNA array**—The protein/DNA array (Combo Array, Panomics/Affymetrix) was performed at the Shanghai KANCHENG Biochip Company (Shanghai, China) in accordance with the protocols recommended for the protein/DNA array system. The nuclear protein extracts were semi-quantitatively assayed by measuring the DNA binding activity of 345 TFs. The nuclear protein extracts were prepared from the HT-29-Gal cells and the HT-29-Gal cells treated with the TFLK-containing 10-mer BP as previously described (22). Briefly, 5 μg of nuclear protein pooled from four HT-29-Gal cell extracts or from four TFLK-containing 10-mer BP-treated HT-29-Gal cell extracts were used for binding to a mixture of biotin-labeled TF-specific DNA probes in solution. The unbound DNA probes were washed away, and the TF-bound probes were denatured and hybridized to the array membrane. After the addition of streptavidin-HRP, signals were generated using enhanced chemiluminescence (ECL) and exposure to Hyperfilm ECL (Amersham Biosciences, Piscataway, NJ, USA). Various exposure times were used to generate signals over a large dynamic range. The signal intensities were quantified using a Bio-Rad calibrated GS-800 scanner and the Quantity One software (Bio-Rad, Hercules, CA, USA). Only the non-saturated signals were used for further analysis. The TF binding activity was considered significant when at least a two-fold signal difference was observed between the HT-29-Gal cells and the HT-29-Gal cells treated with the TFLK-containing 10-mer BP.

**Electrophoretic mobility-shift assay (EMSA)**—The nuclear protein extract was prepared from the HT-29-Gal cell line. The following double-stranded DNA probes were used: the wild-type Sp1 consensus oligonucleotide (5’-ATTGATCGGGCCGGGCAGC-3’), the wild-type CREB1 consensus oligonucleotide (5’-AGAGATTGCGCTACGAGAGTACAG-3’), the wild-type USF-1 consensus oligonucleotide (5’-CACCGGGTCAGTGGGCCTACACC-3’) (positive control); the mutated Sp1 oligonucleotide (5’-ATTGATCGGTCGGGGCCAGC-3’), the mutated CREB1 oligonucleotide (5’-AGAGATTGCCTGGCTACGAGAGTACAG-3’), and the mutated USF-1 oligonucleotide (5’-CACCGGGTCAGTGGGCCTACACC-3’). The probes were purchased from Santa Cruz Biotechnology Inc. An isotopic electrophoretic mobility-shift assay (EMSA) was performed as described previously (23), using 5 μg of the nuclear cell extracts, 50 ng of the probe, and 1 μg of salmon testes DNA (Sigma-Aldrich, St Louis, MO, USA) in a 5 × binding reaction buffer (100 mM Hapes, pH 7.9; 250 mM KCl; 2.5 mM DTT; 0.25 mM EDTA; 5 mM MgCl2; and 25% glycerol). After 20 minutes, the samples were separated on a 10% polyacrylamide gel (39:1 polyacrylamide:bisacrylamide) at a constant temperature of 4°C in 1 × Tris-borate-EDTA buffer (45 mA for 4.5 hours). Similarly, for the supershift assay, 8 ng/μl of specific antibody (anti-Sp1, anti-CREB1, or anti-USF-1) or the control antibody (anti-ER) was incubated for 20 minutes in the HT-29-Gal cell nuclear protein extracts before the addition of the probes. The samples were size-separated by electrophoresis on
a 10% polyacrylamide minigel (39:1 polyacrylamide:bisacrylamide) at room temperature in 0.5 × Tris-borate-EDTA buffer.

Chromatin immunoprecipitation assay (ChIP) — ChIP assays were performed using a ChIP assay kit according to the manufacturer's instructions (Upstate, Lake Placid, NY). Soluble chromatin was prepared from HT-29-Gal cells treated with or without TFLK-containing 10-mer BP. Chromatin was immunoprecipitated with antibodies against CDX2, Sp1 or CREB1, respectively. The final DNA extracts were amplified by PCR using primer pairs that included a CDX2, Sp1 or CREB1 consensus sequence in the human MUC3 promoter (supplementary Figure 1). The primer sequences and the amplified PCR products that included CDX2 (ChIP1 and ChIP2), CREB1 (ChIP3) and or Sp1 (ChIP4) consensus sequence in the MUC3 promoter were presented in supplementary Table 3.

Bacterial strains, growth conditions and bacterial adherence assay — For the in vitro inhibition of EPEC or EHEC O157:H7 (purchased from the Chinese Center for Food Safety Risk Assessment, Beijing, China) adherence to the HT-29 intestinal cells and the HT-29-Gal cells, the cells were grown to near confluence in glucose-free, galactose-containing McCoy's 5a culture medium in 12-well tissue culture plates. The cells were washed and the antibiotic-free culture medium containing the M-SAA3 derived N-terminal 10-mer monomeric peptide (50 μg/ml or 200 μg/ml), the TFLK-containing 10-mer BP (50 μg/ml or 200 μg/ml) or no peptide was added 1 h prior to incubation with the bacteria. After the 1 h incubation with or without stimulation with the peptides, the EPEC or EHEC O157:H7 bacteria (10⁶ CFU in 0.1 mL PBS, pH 7.4) were added to each well and the cells were incubated for an additional 3 h at 37 °C. The cells were washed with chilled PBS four times for 5 min each to remove the non-adherent EPEC or EHEC O157:H7 bacteria and to inhibit further growth of the adhering bacteria. The cells with adherent bacteria were released from the polystyrene wells using trypsin-EDTA. The bacteria were quantified by determining the CFU following the plating of serial dilutions of the bacteria on MacConkey agar (BD Biosciences, New Jersey, USA) and incubation overnight at 37 °C. The number of viable EPEC or EHEC O157:H7 bacteria that remained on the TFLK-containing 10-mer BP-treated HT-29-Gal cells was compared to the number of bacteria adhered to the HT-29-Gal cells not exposed to peptides or exposed to the M-SAA3 derived N-terminal 10-mer monomeric peptide and HT-29 cells. The individual experiments were performed in triplicate and each experiment was repeated at least twice.

RESULTS

Schematic representation of TFLK-containing 10-mer BP and M-SAA3 derived N-terminal 10-mer monomeric peptide — The M-SAA3 derived N-terminal 10-mer monomeric peptide (residues 1-10, sequence GWLTFLKAAG) is shown in Figure 1A and a schematic representation of the tetramERICally branched polyopeptide based on the conserved TFLK-motif from M-SAA3 (TFLK-containing 10-mer BP) is shown in Figure 1B. The four amino acids (the TFLK motif) are underlined and are conserved within the first eight N-terminal amino acid residues of M-SAA3.

TFLK-containing 10-mer-BP enhances MUC3 expression in HT-29-Gal cells — To determine the ideal concentration of the TFLK-containing 10-mer BP for the activation of MUC3 expression in the HT-29-Gal cells, we analyzed the MUC3 mRNA levels by real-time PCR in the HT-29-Gal cells stimulated with different concentrations of the TFLK-containing 10-mer BP (50, 100, 200, 400 and 800 μg/ml) and compared it to the
untreated HT-29-Gal cells at three time points (0.5, 1 and 3 hours). We found that 200 μg/ml of the TFLK-containing 10-mer BP produced the most efficient stimulation of MUC3 mRNA expression at 1 hour point (p<0.01) (Figure 1C).

As shown in Figure 1D, the MUC3 mRNA levels in the HT-29-Gal cells following stimulation with the TFLK-containing 10-mer BP for 1 hour (200 μg/ml) was significantly higher than the untreated HT-29-Gal cells (p<0.01) and the HT-29-Gal cells stimulated with the M-SAA3 derived N-terminal 10-mer monomeric peptide (200 μg/ml) (p<0.01). The MUC3 protein level in HT-29-Gal cells after stimulation with the TFLK-containing 10-mer BP was significantly higher than in the untreated HT-29-Gal cells and the HT-29-Gal cells after stimulation with the M-SAA3 derived N-terminal 10-mer monomeric peptide (p<0.01) (Figure 1E). There was no significant difference in the MUC3 protein levels between the untreated HT-29-Gal cells and the HT-29-Gal cells after stimulation with the M-SAA3 derived N-terminal 10-mer monomeric peptide (p>0.05) (Figure 1E).

Characterization of the human MUC3 promoter — As shown in Figure 2, the human MUC3 promoter region produced a high luciferase activity in the HT-29-Gal cells in the presence and absence of the TFLK-containing 10-mer BP. The data indicated that this region of the human MUC3 promoter (from nucleotides -999 to +57) contains the regulatory elements responsible for the expression of human MUC3. To locate the specific elements involved in MUC3 expression, we generated the following series of truncations of the promoter region: pGL3-M4 (-775 to +57), pGL3-M3 (-623 to +57), pGL2-M2 (-242 to +57) and pGL3-M1 (-62 to +57) (Figure 2A). Truncation of the upstream region to within 242 bp of the putative transcription start sites (TSS) produced the strongest effect on the luciferase activity; however, further truncation of the upstream region to within 62 bp of the TSS resulted in a significant loss of activity, indicating that multiple elements between nucleotides -242 and -62 contribute to the activity of the promoter (Figure 2B). The truncation of the upstream region to within 999, 775 and 623 bp of the TSS produced a modest effect on the promoter activity compared with truncation of the upstream region to within 242 bp of the TSS. Therefore, multiple elements between nucleotides -999 and -242 exerted an inhibitory effect on the activity of the promoter (Figure 2B).

TFLK-containing 10-mer BP transcriptionally activated the MUC3 promoter in HT-29-Gal cells — The TFLK-containing 10-mer BP had a significant effect on increasing the luciferase activity in HT-29-Gal cells for the promoter truncations pGL3-M1 (p<0.05), pGL3-M2 (p<0.01), pGL3-M4 (p<0.05), and pGL3-M5 (p<0.05) (Figure 2B). The data indicated that multiple elements between -999 and -775, -775 and -623, -242 and -62, and -62 and +57 contributed to the activity of the human MUC3 promoter when stimulated with TFLK-containing 10-mer BP in the HT-29-Gal cells.

Transcriptomic responses to the TFLK-containing 10-mer BP — The Protein/DNA Combo array analysing the transcriptomic changes in the HT-29-Gal cells in response to the TFLK-containing 10-mer BP indicated the transcription factors (81 in total) that were upregulated 2.0-fold in the TFLK-containing 10-mer BP-treated HT-29-Gal cells versus the untreated HT-29-Gal cells are shown in supplementary Table 4. The transcription factors (12 in total) that were down-regulated 2.0-fold in the TFLK-containing 10-mer BP-treated HT-29-Gal cells versus the untreated HT-29-Gal cells are shown in supplementary Table 5. The following eleven transcription factors: AP-1,
GAS/ISRE, Pbx1, AP-2, GATA, CREB1, USF-1, RIP3a1, Pax3, Sp1 and PRDII-BF1, were up-regulated 10-fold.

Analysis of MUC3 promoter indicating that Sp1, CREB1 and CDX2 might be responsible for the up-regulation of MUC3 — The genomic region flanking the MUC3 promoter (from -999 to +57) containing the consensus sequences for possible TF-binding sites predicted using TFSEARCH (ver 1.3) (http://mbs.cbrc.jp/research/db/TFSEARCH.html) are depicted in supplementary Figure 1. These elements, including CREB1, Nkx-2, CDX2, SRY, deltaE, HSF2, C/EBP, GATA, Sox5, HNF-2, AML-1a, CDXA, c-Rel, IK-2/3, N-Myc, USF, and Sp1, are present in the promoter region (from -999 to +57) of the human MUC3 gene.

We focused on the Sp1, CREB1, USF1, and CDX2 elements for the following reasons: the Sp1, CREB1 and USF1 cis-elements are located between -242 and -64 of the TSS of MUC3 and were significantly induced following stimulation with the TFLK-containing 10-mer BP, and CDX2 is critical for the development of the intestinal epithelium (24-27).

TFLK-containing 10-mer BP activates CDX2, CREB1, Sp1 and USF1 in HT-29-Gal cells accompanied by the up-regulation of MUC3 — The mRNA levels of CDX2, CREB1, Sp1 and USF1 quantified by real-time PCR analysis were significantly increased (3.2-, 2.3-, 4.4- and 3.0-fold, respectively) in the HT-29-Gal cells after stimulation with the TFLK-containing 10-mer BP, and CDX2 is critical for the development of the intestinal epithelium (24-27).

ChIP assays indicated Sp1, CREB1 and CDX2 binding to the MUC3 promoter and Sp1 and CREB1 binding to the MUC3 promoter increased in HT-29-Gal cells treated with TFLK-containing 10-mer BP — To determine if CDX2, Sp1, and CREB1 can bind directly to the potential sites in the human MUC3 promoter and their bindings increased due to TFLK-containing 10-mer BP...
treatment in HT-29-Gal cells, we performed chromatin immunoprecipitation (ChIP). As shown in Figure 5A, two CDX2, one CREB1 and one Sp1 potential binding sites were located in the proximal promoter of the human MUC3 gene. The binding of these transcription factors in their consensus sites within the MUC3 promoter was exactly present when compared with the negative controls. The binding of CDX2 to its potential sites was not increased in HT-29-Gal cells after treatment with TFLK-containing 10-mer BP, however, the binding of CREB1 and Sp1 to their potential sites was increased in HT-29-Gal cells after treatment with TFLK-containing 10-mer BP compared with untreated HT-29-Gal cells (Figure 5B).

_TFLK-containing 10-mer BP significantly inhibited the adherence of EPEC and O157 EHEC O157:H7 to HT-29-Gal cells_—To evaluate the effect of the TFLK-containing 10-mer BP on the adherence of E. coli (EPEC) and E. coli (EHEC) O157:H7 to human intestinal epithelial cells, the TFLK-containing 10-mer BP (50 μg/mL or 200 μg/mL) or the M-SAA3 derived N-terminal 10-mer monomeric peptide (50 μg/mL or 200 μg/mL) were added to the culture medium of HT-29-Gal or HT-29 cells, and the effect of the peptides on the adherence of the bacteria was examined. As shown in Fig. 6A, pretreatment of the HT-29-Gal cells with 200 μg/mL of the TFLK-containing 10-mer BP produced the greatest reduction in adherence of the E. coli (EPEC) compared to the untreated HT-29-Gal cells and the HT-29-Gal cells pretreated with 50 μg/mL TFLK-containing 10-mer BP or 50 μg/mL of the M-SAA3 derived N-terminal 10-mer monomeric peptide (p<0.01) and the HT-29-Gal cells pretreated with 200 μg/mL of the M-SAA3 derived N-terminal 10-mer monomeric peptide (p<0.05) (Figure 6B). The results suggested that the TFLK-containing 10-mer BP of the N-terminal human serum amyloid A3 is more efficient than the M-SAA3 derived N-terminal 10-mer monomeric peptide at inducing MUC3 expression, and leading to the significant inhibition of E. coli (EPEC) and E. coli (EHEC) O157:H7 adherence to intestinal epithelial cells.

Comparing the adherence of the E. coli (EPEC) and E. coli (EHEC) O157:H7 to the HT-29 and HT-29-Gal cells, it was interesting that the HT-29-Gal cells (a more mature and differentiated form of the HT-29 cell line) exhibited a significant reduction in the adherence of both the E. coli (EPEC) and E. coli (EHEC) O157:H7 strains. Therefore, the differentiation of the intestinal epithelial cells is associated with the adherence of E. coli (EPEC) or E. coli (EHEC) O157:H7.

**DISCUSSION**

Based on the study by Mack et al., the M-SAA3 N-terminal 10-mer peptide (GWLTFLKAAG) containing the intact TFLK-motif, increased MUC3 mRNA expression (14). M-SAA3 is a constituent of colostrum in humans and other mammalian species and plays a protective gastrointestinal role through the overexpression of intestinal mucin. It is likely that the presence of M-SAA3 in human colostrum is one of the reasons why colostrum-fed infants are approximately 7-10 times less likely to suffer gastrointestinal diseases than formula-fed infants (10). A study by Domènech et al. produced the recombinant goat
milk serum amyloid A isoform 3 encompassing the TFLK region and showed that it was responsible for the up-regulation of mucins in the intestine; however, the protein did not promote macrophage phagocytosis (28). The experiments ruled out the possibility of determining the serum-derived forms of SAA that can opsonize Gram-negative bacteria facilitating their phagocytosis by circulating macrophages or intestinal epithelial cells (28). The exact mechanism of the intact TFLK-motif in the M-SAA3-mediated induction of MUC3 expression and the inhibition of bacterial adherence remains unknown. Therefore, in our study, we investigated a more efficient design for a polypeptide based on the conserved TFLK-motif from M-SAA3 that would induce increased MUC3 mucin expression in the intestinal epithelial cells and would be a more effective treatment to prevent enteric pathogen adherence. We also investigated the biological mechanism of the induction of human MUC3 expression by the TFLK-containing 10-mer BP.

Combinatorial chemistry has had a profound effect on the approach of medical chemists to drug discovery, especially on the strategies for the synthesis and screening of synthetic peptides developed over the last two decades (29, 30). The linear, cyclic, multimeric and branched structures have been used as scaffolds into which natural or non-natural amino acids have been inserted to increase the chemical diversity or to achieve a more rigid conformation and more stable structures (17). Several reports have demonstrated that increasing the copy number of a precursor peptide by multimerization is a simple and effective way to enhance immunogenicity, affinity, and stability. It is known that the multimerization of a bioactive monomeric peptide essentially leads to the multiplication of the interacting groups/sites of the starting unit but does not create new recognition surfaces. Instead, multimerization provides new molecules with an increased avidity deriving from the multiple identical and cooperative sites. The tetrameric branched tripeptide 4-23-5 binds to the vascular endothelial growth factor receptor-1 (VEGFR-1) and inhibits the formation of the structures similar to capillaries that are formed by primary human endothelial cells. The monomeric and dimeric variants were inactive, and the trimeric variants were partially active (31). Pini et al. reported that the tetrabranched peptide based on the nonnatural antimicrobial peptide KKIRVRLSA is a good candidate for the development of a new antibacterial drug (32). Based on the finding by Mack et al. showing that the M-SAA3 N-terminal 10-mer peptide induces human MUC3 transcriptional expression and inhibits EPEC adherence, we generated a synthetic tetrameric structure of branched polylysine based on the conserved TFLK-motif from M-SAA3 (GWLTFLKAAG). This tetrameric peptide is more efficient at inducing human MUC3 expression in the HT-29-Gal cells than the M-SAA3 derived N-terminal 10-mer monomeric peptide at both transcriptional and protein levels. The results indicated that the synthetic branched peptide offers the numerous advantages over biotherapeutics because it is easier and cheaper to produce, is more stable, and is generally free of biological contaminants. Furthermore, the tetrameric peptide offers more opportunities for delivery.

Although the function of the M-SAA3 N-terminal 10-mer peptide in inducing MUC3 expression and inhibiting EPEC adherence is clear, the mechanism behind the biological phenomena is not understood. This polypeptide, as an exogenous stimulator, induces the intestinal epithelial cells to express MUC3 mucin; however, how it interacts with the cell surface (possibly
through a receptor) and how it transmits activating signals into the cells is unknown. Remarkably, the results from the transcription factor protein/DNA array comparing the HT-29-Gal cells stimulated with the TFLK-containing 10-mer BP with the untreated HT-29-Gal cells indicated that at least 81 TFs were up-regulated 2.0-fold (supplementary Table 4), and only 12 TFs were down-regulated 2.0-fold (supplementary Table 5). Our study of the MUC3 promoter and previous studies of the MUC3 or Muc3 promoters (33, 34) showed that the TFLK-containing 10-mer BP produced the highest luciferase activity in HT-29-Gal cells when using the pGL3-M2 construct (Figure 2); therefore, the multiple cis-elements in the -242 to -64 region of the TSS of human MUC3 contribute to the activity of the human MUC3 promoter. The N-myc, USF1, CREB1, MZF1 and Sp1 cis-elements were shown to bind to the -242 to -64 region of the human MUC3 TSS. N-myc was not induced by the TFLK-containing 10-mer BP (supplementary Table 4); however, the levels of USF1, CREB1 and Sp1 increased more than 10-fold following stimulation with the TFLK-containing 10-mer BP, and the MZF-1 levels increased only 2.65-fold. The ChIP results in Figure 5 suggest that the enhanced activity of the pGL3-M2 construct results in part from enhanced binding of Sp1 and CREB1 to the indicated sites in the proximal promoter. Together the results from Figure 2, Figure 4, and Figure 5 suggest a general role for CDX2 in MUC3 up-regulation, and a specific role for Sp1 and CREB1 in MUC3 up-regulation following stimulation with TFLK-containing 10-mer BP. To our knowledge, this is the first study to report a mechanism for the biological function of the TFLK-motif.

Notably, the induction of human MUC3 mucin by the TFLK-containing 10-mer BP led to the dramatic inhibition of EPEC or EHEC O157:H7 adherence in HT-29-Gal cells. This finding suggests that the TFLK-containing 10-mer BP has potential applications for molecular therapeutic approaches in the prevention of gastrointestinal infection. Furthermore, the TFLK-containing 10-mer BP was more efficient at inhibiting adherence of the bacteria than the M-SAA3 derived N-terminal 10-mer monomeric peptide. EPEC is an important cause of infant diarrhea in developing countries (35). This bacteria produces a characteristic intestinal histopathological lesion on enterocytes known as ‘attaching and effacing’ (A/E), and these two steps are mediated by a type-III secretory system. EHEC O157:H7 causes illnesses ranging from mild diarrhea to severe diseases such as hemorrhagic colitis and hemolytic uremic syndrome (36). EHEC O157:H7, which produces Shiga toxin (Stx), is the major EHEC O157:H7 serotype responsible for public health problems worldwide (37). We propose that human MUC3 production induced by the TFLK-containing 10-mer BP improves the intestinal defense mediated by epithelial cells and thereby protects the host against lethal infection.

Collectively, our results suggest that the TFLK-containing 10-mer BP based on the conserved TFLK-motif from M-SAA3 is a new specific compound design that has potential applications for clinical uses to prevent gastrointestinal infection through the multiple transcription factor-mediated up-regulation of human MUC3 mucin.
REFERENCE


**FOOTNOTE**

*This work was supported by The National Natural Science Foundation of China (30800519 and 81170340)*
FIGURE LEGENDS

Figure 1. Schematic representation of the M-SAA3 derived N-terminal 10-mer monomeric peptide (A) and the branched polypeptides based on the TFLK motif (B) and their induction of MUC3 expression in HT-29-Gal cells. TFLK-containing 10-mer BP increased MUC3 mRNA expression levels in HT-29-Gal cells dose-dependently compared with non-treated HT-29-Gal cells (C). TFLK-containing 10-mer BP (10TFLK-BP) (200 μg/mL) induced MUC3 mRNA expression levels in HT-29-Gal cells at significantly higher levels than in untreated HT-29-Gal cells (p<0.01) and in HT-29-Gal cells pretreated with the M-SAA3 derived N-terminal 10-mer monomeric peptide (10TFLK) (D) (p<0.01). Western blot experiments demonstrated similar results for MUC3 protein levels, the anti-MUC3 antibody used in the study recognized the epitope which was located at the carboxyl terminal domain of human MUC3 and just after the proteolytic cleavage site (LRNGSIVV) and can only recognized the cleaved C-terminal fragment of human MUC3, so the Mt of MUC3 protein by Western blot was in 96kDa (E) (p<0.01). The M-SAA3 derived N-terminal 10-mer monomeric peptide (10TFLK) did not increase MUC3 protein levels comparing with untreated HT-29-Gal cells (p>0.05) (E), real-time PCR experiments, however, demonstrated a significant difference between these samples (p<0.05) (D), ‘∗’ represents a significant difference (p<0.05), ‘∗∗’ represents a significant difference (p<0.01).

Figure 2. Transcriptional regulation of human MUC3 by the TFLK-containing 10-mer BP. (A) Schematic representation of the human MUC3 promoter constructs. The five deletion luciferase constructs that were generated to identify the sites of transcriptional regulation within the MUC3 promoter responded to the TFLK-containing 10-mer BP (10TFLK-BP). (B) HT-29-Gal cells were transfected with the different luciferase constructs and were treated or not treated with the TFLK-containing 10-mer BP. The cells were harvested after 48 h and the relative luciferase activity was determined using the Dual-Luciferase Reporter Assay System on a single sample luminometer. The data represent the mean±SEM of three independent experiments. ‘∗’ represents a significant difference (p<0.05), ‘∗∗’ represents a significant difference (p<0.01).

Figure 3. Expression of multiple transcription factors following stimulation with the TFLK-containing 10-mer BP. The mRNA levels of CDX2, CREB1, SP1, USF1 and MUC3 in HT-29-Gal cells treated with the TFLK-containing 10-mer BP (10TFLK BP) (200 μg/mL), untreated HT-29-Gal cells and untreated HT-29 cells were determined using real-time PCR (A). The protein levels of CDX2, CREB1, SP1 and USF1 in HT-29-Gal cells treated with the TFLK-containing 10-mer BP (10TFLK BP) (200 μg/mL), untreated HT-29-Gal cells and untreated HT-29 cells were determined using Western blotting (B). The protein levels of each transcription factor relative to the nuclear protein internal control, lamin B1, in three independent experiments, are shown in panel C. ‘∗’ represents a significant difference (p<0.05), ‘∗∗’ represents a significant difference (p<0.01).

Figure 4. TFLK-containing 10-mer BP-treated HT-29-Gal cells demonstrate the increased binding activities of Sp1, CREB1 and CDX2 to the MUC3 promoter. Representative EMSA of the Sp1, CREB1 and CDX2 response elements in the human MUC3 promoter using the nuclear extract from HT-29 cells (HT-29), HT-29-Gal cells (HT-29-G) and HT-29-Gal cells treated with 200 μg/mL
TFLK-containing 10-mer BP (TFLK), the complexes formed with Sp1, CREB1 or CDX2 probes and the nuclear extract from HT-29-Gal cells treated with 200 μg/mL TFLK-containing 10-mer BP were more than that formed with Sp1, CREB1 or CDX2 probes and the nuclear extracts from non-treated HT-29 cells and untreated HT-29-Gal cells (A). The supershift EMSA was performed to confirm that the induced Sp1, CREB1 or CDX2 complexes were specific and contained Sp1, CREB1 or CDX2, the antibodies against Sp1, CREB1 or CDX2 led to the supershift (marked as arrow) compared with the relative complexes (B and C). Competitive inhibition using the mutated Sp1, CREB1 or CDX2 probes led to the failure of complexes forming. The wild types of Sp1, CREB1 or CDX2 probes, antibodies against Sp1, CREB1 or CDX2 (and nonimmune serum, IgG), and mutants (Sp1 mutation, mSp1; CREB1 mutation, mCREB1; and CDX2 mutation, mCDX2) were used in the supershift EMSAs.

**Figure 5.** CDX2, Sp1, or CREB1 bind to the MUC3 promoter in response to TFLK-containing 10-mer BP treatment. Chromatin was isolated from HT-29-Gal cells that were either untreated or treated with 200 μg/mL TFLK-containing 10-mer BP. Chromatin was then subjected to immunoprecipitation using antibodies for IgG (negative control), CDX2, Sp1, CREB1, or RNA Poly-merase II (positive control). “Input” represents 10% of the DNA used in the immunoprecipitation. Four loci were tested in the MUC3 promoter (two CDX2, one CREB1 and one Sp1 consensus sequences), “ChIP1”, “ChIP2”, “ChIP3” and “ChIP4”. The binding sites for first CDX2 (-960 to -951), second CDX2 (-523 to -505), CREB1 (-148 to -136) and Sp1 (-92 to -84) in the MUC3 promoter region were presented (A). Final DNA extracts were amplified by PCR using primers that included CDX2 (ChIP1 and ChIP2), CREB1 (ChIP3) and or Sp1 (ChIP4) consensus sequence in the MUC3 promoter (B).

**Figure 6.** EPEC and EHEC O157:H7 adherence to HT-29-Gal cells following stimulation with the TFLK-containing 10-mer BP or the M-SAA3 derived N-terminal 10-mer monomeric peptide. HT-29-Gal cells or HT-29 cells received a single dose of the TFLK-containing 10-mer BP (10TFLK-BP) at 50 or 200 μg/mL or the M-SAA3 derived N-terminal 10-mer monomeric peptide (10TFLK) at 50 or 200 μg/mL. The cells were incubated at 37 °C for 1 h prior to the addition of 10^6 CPU/well of EPEC or EHEC O157:H7. After incubation for 3 h, the unbound bacteria were removed by four washes with chilled PBS (pH 7.4). The EHEC O157:H7 (A) or EPEC (B) adhering to the HT-29-Gal or HT-29 cells were quantified by determining the CFU/well. The results are expressed as the mean±SEM of two independent experiments that were run in triplicate. ‘*’ represents a significant difference (p<0.05), ‘**’ represents a significant difference (p<0.01).
Figure 1

A  N-terminal 10-mer of M-SAA3 containing the TFLK motif (residues 1-10):

GWLTFLKAAG

B  Schematic representation of branched polypeptides Based on TFLK-motif

GWLTFLKAAG

GWLTFLKAAG

GWLTFLKAAG

C  Normalized Fold Expression of MUC3 mRNA at Cq

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D  Normalized Fold Expression of MUC3 mRNA at Cq

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E  MUC3/GAPDH protein density (% of control)

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** p < 0.01
*  p < 0.05
Figure 2

A

-999  -775  -57  -623  -242  -62

B

pGL3-M6  pGL3-M4  pGL3-M3  pGL3-M2  pGL3-M1  pGL3

Relative luciferase activity

[] treated with 10TFLK-BP
[] untreated

** **

* *
Figure 3

A

B

C

untreated HT-29
untreated HT-29-Gal
HT-29-Gal cells treated with 10TFLK BP

Relative expression levels of mRNA

0 1 2 3 4 5 6 7

CDX2 CREB MUC3 SP1 USF1

40kDa
43kDa
106kDa
43kDa
67kDa

CDX2 CREB SP1 USF1 Lamin B1

untreated HT-29
untreated HT-29-Gal
HT-29-Gal cells treated with 10TFLK BP

CDX2 CREB-1 SP1 USF1
Figure 4

A) Nuclear protein

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Complex →

Free probe →

B) Nuclear protein

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Antibody

|       | - | - | - | Sp1 | IgG | - | - | CDX2 | IgG | - | - | CREB | IgG |
|       | - | - | - |      |      | - | - |      |      | - | - |      |      |

Supershift → Complex →

Free probe →

C) Nuclear protein

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Competitor

|       | - | - | - | mSp1 | - | - | mCDX2 | - | - | mCREB |
|       | - | - | - |      | - | - |      | - | - |      |

Complex →

Free probe →
Figure 5

A

\[
\begin{align*}
\text{Binding sites} & \quad \text{CDX2} \quad \text{CDX2} \quad \text{CIRB} \quad \text{Sp1} \\
\text{ChIPs} & \quad \text{CDX2} \quad \text{ChIP1} \quad \text{CDX2} \quad \text{ChIP2} \quad \text{CIRB} \quad \text{ChIP3} \quad \text{Sp1} \quad \text{ChIP4}
\end{align*}
\]

B

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<td>ChIP4</td>
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Figure 6

A

B

Adherent bacteria (CFU/well)

E. coli (EHEC) O157:H7

HT-29  HT-29-Gal  HT-29-Gal  HT-29-Gal  HT-29-Gal  HT-29-Gal

10TFLK -  -  50  -  200  -

10TFLK-BP -  -  -  50  -  200

Adherent bacteria (CFU/well)

E. coli (EPEC)

HT-29  HT-29-Gal  HT-29-Gal  HT-29-Gal  HT-29-Gal  HT-29-Gal

10TFLK -  -  50  -  200  -

10TFLK-BP -  -  -  50  -  200

**

**

**

**

**

*
Enhanced Membrane-tethered Mucin 3 (MUC3) Expression by a Tetrameric Branched Peptide with Conserved TFLK Motif Inhibits Bacteria Adherence
Qiong Pan, Yin Tian, Xiaohuan Li, Jun Ye, Yun Liu, Lili Song, Yongtao Yang, Rong Zhu, Yonghong He, Lei Chen, Wensheng Chen, Xuhu Mao, Zhihong Peng and Rongquan Wang

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