L-fucose ameliorates the carcinogenic properties of *Fusobacterium nucleatum* in colorectal cancer

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Abstract. *Fusobacterium nucleatum* (Fn) is considered a promoting factor in colorectal cancer (CRC); however, only a few studies have investigated therapies against Fn. L-fucose is a natural monosaccharide that has prebiotic potential. The present study aimed to investigate the effect of L-fucose on the carcinogenic properties of Fn. The HCT116 and SW480 colon cancer cell lines were treated with Fn and Fn+L-fucose (Fnf), respectively. The Cell Counting Kit-8 (CCK-8), colony formation, Transwell migration and invasion and wound healing assays were performed to assess the proliferative, migratory and invasive abilities of the cells, respectively. Western blot was performed to detect the protein levels of jak/stat3 pathway components and EMT. The results of the CCK-8, colony formation, Transwell and wound healing assays demonstrated that treatment with Fn significantly enhanced the proliferative, migratory and invasive abilities of HCT116 and SW480 colon cancer cells. Notably, these effects were significantly reversed following addition of L-fucose. Furthermore, L-fucose inhibited the carcinogenic properties of Fn to activate the stat3 pathway and epithelial-to-mesenchymal transition. Taken together, the results of the present study suggest that L-fucose ameliorates the carcinogenic properties of *Fn in vitro*, and thus may serve as a novel therapeutic target for flora-related colon cancer.

Introduction

Colorectal cancer (CRC) is the fourth most common cancer worldwide (6.1% of total cases in 2018) (1). Despite advancements in its diagnosis and treatment strategies, CRC remains the leading cause of cancer-associated mortality worldwide (9.2% of the total cancer-associated deaths in 2018) (1). Thus, better understanding the molecular mechanism of CRC progression and identifying novel effective therapeutic strategies remains essential.

Gut microbiome is closely associated with the development of CRC (2). Recently, the Gram-negative anaerobic bacteria, *Fusobacterium nucleatum* (Fn), has gained great interest. Previous studies have demonstrated that Fn is enriched in CRC tissues compared with adjacent normal tissues, and associated with poor prognosis (3,4). In addition, Fn promotes the development of CRC by activating autophagy of cancer cells, protecting tumors from immune attack and creating an inflammatory microenvironment (5,6). Although it has been reported that using berberine and targeting Fn Fap2 may decrease Fn potentiation of CRC (7,8), methods for suppressing the carcinogenic properties of Fn remain largely unknown.

L-Fucose (Fucose), a natural monosaccharide present in foods or bodies, plays an important role in sustaining the gut homeostasis (9). Our previous studies demonstrated that fucose alleviates dextran sulfate sodium-induced acute and chronic colitis by regulating immune responses and affecting the intestinal microenvironment (10,11). Notably, fucose has been reported to affect microbial metabolic pathways, and decrease pathogen virulence (12). However, the role of fucose in CRC remains unknown.

Thus, the present study aimed to investigate the effects of fucose on the functional regulations of Fn and determine its underlying molecular mechanism in CRC. Taken together, the results of the present study suggest that fucose may ameliorate the carcinogenic properties of Fn.

Materials and methods

Cell lines, bacterial strain culture and groups. The HCT116 and SW480 human colon cancer cell lines were purchased from the American Type Culture Collection and maintained in high glucose DMEM supplemented with 10% fetal bovine serum (FBS) and 100 U/ml streptomycin/penicillin (all purchased from Gibco; Thermo Fisher Scientific, Inc.), at 37°C in 5% CO₂. Fn (25586) was purchased from the American Type Culture Collection and cultured at Wuhan Research Institute of First
Light Industry (Wuhan, China). The methods of bacterial pellets and conditioned medium were performed, as previously described (13). The following groups were classified: Control group, Fn group (supernatant of Fn was added to the cells) and 0.5% fucose+Fn (Fnf) group. Fucose (Sigma-Aldrich; Merck KGaA) was added when Fn was cultured and the supernatant was subsequently added to the cells. In total, 100 nM 3I-201 (Selleck Chemicals), an inhibitor of stat3, was incubated for 1 h prior to bacteria treatment to access the role of stat3 pathway.

Cell Cell Potentiation (EMD Millipore) and blocked with 5% of protein (40 µg/lane) were separated by 10% SDS-PAGE.

The separated proteins were subsequently transferred onto PVDF membranes (EMD Millipore) and blocked with 5% of Biotechnology). Total protein was quantified using the BCA assay (Thermo Fisher Scientific, Inc.) and equal amounts of Biotechnology) supplemented with phenylmethyl sulfonyl fluoride protease inhibitor (Beyotime Institute of Biotechnology). Western blotting.

membranes were precoated with Matrigel at 37˚C for 4 h. The colon cancer cells were stimulated with Fn or Fnf for 12 h at 37˚C and subsequently seeded into 6-well plates at a density of 500 cells/well. Cells were cultured for 14 days at 37˚C in 5% CO₂. Following incubation, cells were washed with PBS for three times, fixed with 4% paraformaldehyde for 15 min, and stained with 0.5% crystal violet for 15 min both at room temperature. Cell colonies were counted using ImageJ software (National Institutes of Health).

Wound healing assay. Cells from each group were respectively seeded into 6-well plates at a density of 5x10⁵ cells/well. Cells were washed with PBS three times, fixed with 4% paraformaldehyde for 30 min and stained with 0.5% crystal violet. Cells from each group were respectively seeded into 6-well plates at a density of 5x10⁵ cells/well. Cells were washed with PBS three times, fixed with 4% paraformaldehyde for 30 min and stained with 0.5% crystal violet. Cells from each group were respectively seeded into 6-well plates at a density of 5x10⁵ cells/well. Cells were washed with PBS three times, fixed with 4% paraformaldehyde for 30 min and stained with 0.5% crystal violet. Cells from each group were respectively seeded into 6-well plates at a density of 5x10⁵ cells/well. Cells were washed with PBS three times, fixed with 4% paraformaldehyde for 30 min and stained with 0.5% crystal violet. Cells from each group were respectively seeded into 6-well plates at a density of 5x10⁵ cells/well. Cells were washed with PBS three times, fixed with 4% paraformaldehyde for 30 min and stained with 0.5% crystal violet.

Transwell migration and invasion assays. For the cell migration assay, cells from each group were resuspended with serum-free DMEM medium and plated in the upper chambers of 24-well Transwell plates at a density of 1x10⁵. DMEM medium supplemented with 10% FBS was plated in the lower chambers. Following incubation for 24 h at 37˚C, the migratory cells were fixed with 4% paraformaldehyde for 30 min and stained with 0.5% crystal violet for 15 min both at room temperature. Cells were observed under a light microscope (magnification x200; Olympus Corporation). For the cell invasion assay, Transwell membranes were precoated with Matrigel at 37˚C for 4 h.

Western blotting. Total protein was extracted from cells or tissues using RIPA lysis buffer (Beyotime Institute of Biotechnology) supplemented with phenylmethyl sulfonyl fluoride protease inhibitor (Beyotime Institute of Biotechnology) and phosphatase inhibitor (Beyotime Institute of Biotechnology). Total protein was quantified using the BCA assay (Thermo Fisher Scientific, Inc.) and equal amounts of protein (40 µg/lane) were separated by 10% SDS-PAGE. The separated proteins were subsequently transferred onto PVDF membranes (EMD Millipore) and blocked with 5% BSA at room temperature for 1 h. The membranes were incubated with primary antibodies (all 1:1,000 dilution) against: GAPDH (ABclonal Biotech Co., Ltd.; cat. no. AC001), ACTB (ABclonal Biotech Co., Ltd.; cat. no. AC006), phospho-stat3 (Tyr705; Cell Signaling Technology, Inc.; cat. no. 9145S), stat3 (ABclonal Biotech Co., Ltd.; cat. no. A11216), phosphor-jak2 (Tyr1007/1008; ABclonal Biotech Co., Ltd.; cat. no. AP0531), jak2 (ABclonal Biotech Co., Ltd.; cat. no. A7694), β-catenin (Cell Signaling Technology, Inc.; cat. no. 8480S), E-cadherin (Cell Signaling Technology, Inc.; cat. no. 3195S), N-cadherin (GeneTex, Inc.; cat. no. GTX127345) and Vimentin (Cell Signaling Technology, Inc.; cat. no. 5741S) overnight at 4˚C. Following the primary incubation, membranes were incubated with HRP-labelled secondary antibodies (AntGene; cat. no. ANT020; 1:2,000) at room temperature for 1 h. Protein bands were visualized using enhanced chemiluminescent reagents (Beyotime Institute of Biotechnology).

Statistical analysis. Statistical analysis was performed using SPSS 25.0 software (IBM Corp.) and GraphPad Prism 7.0 software (GraphPad Software, Inc.). All experiments were performed in triplicate and data are presented as the mean ± standard error of the mean. One-way analysis of variance and Tukey’s post hoc test were used to compare differences between multiple groups. P<0.05 was considered to indicate a statistically significant difference.

Results

Fucose ameliorates the pro-proliferative characteristic of Fn on colon cancer cells. To investigate whether there were differences between the impacts that Fn and Fnf exerted on colon cancer cells, the CCK-8 assay was performed on HCT116 and SW480 cells following treatment for 12 h. The results demonstrated that cells treated with Fn proliferated faster compared with the control group (HCT116 control, 0.42±0.01; Fn, 0.60±0.02, P<0.0001; SW480 control, 0.36±0.02; Fn, 0.42±0.01, P=0.005; Fig. 1A). However, following treatment with fucose, Fn exerted a weaker pro-proliferative ability (HCT116 Fn, 0.60±0.02; Fnf, 0.54±0.01, P=0.014; SW480 Fn, 0.42±0.01; Fnf, 0.38±0.01, P=0.009; Fig. 1A). The colony formation assay was subsequently performed to assess the long-term effect on cell proliferation. The results demonstrated that the number of HCT116 colonies treated with Fn increased (control, 52.0±7.0; Fn, 125.3±11.3; P=0.005), while less colonies were observed in the Fnf group (Fn, 125.3±11.3; Fnf, 81.0±9.9; P=0.042; Fig. 1B and C). Similar results were observed in SW480 cells (control, 96±10; Fn, 170±5; Fnf, 131.5±6.5; P=0.022 and P=0.043; Fig. 1B and C). Collectively, these results suggest that L-fucose may ameliorate the pro-proliferative ability of Fn on colon cancer cells.

Fucose ameliorates the pro-migratory ability of Fn on colon cancer cells. The migratory ability of colon cancer cells is a functional characteristic to assess aggressiveness (14). The Transwell migration assay was performed to assess the effect of Fn and Fnf on the migratory ability of HCT116 and SW480 cells. The results demonstrated that treatment with Fn significantly enhanced the migratory ability of HCT116 and SW480 cells (HCT116 control, 119.30±3.84; Fn, 148.30±4.41, P=0.008;
SW480 control, 123.00±4.93; Fn, 216.50±3.30; P<0.0001; Fig. 2A), while the migratory ability decreased in cells treated with Fnf (HCT116 Fn, 148.30±4.41; Fnf, 128.80±2.29; P=0.008; SW480 Fn, 1216.50±3.30; Fnf, 157.70±2.96, P=0.002; Fig. 2A).

The results of the wound healing assay demonstrated that treatment with Fn increased the migratory rate in HCT116 cells compared with both the control cells and Fnf treated cells (control, 0.152±0.006; Fn, 0.313±0.020; Fnf, 0.221±0.011; P=0.002 and P=0.018; Fig. 2B). Similar trends were observed in SW480 cells (control, 0.190±0.011; Fn, 0.410±0.016; Fnf, 0.259±0.029; P<0.0001 and P=0.004; Fig. 2C). Taken together, these results suggest that L-fucose may inhibit the pro-migratory ability of Fn on colon cancer cells.

Fucose ameliorates the pro-invasive ability of Fn on colon cancer cells. The Transwell invasive assay was performed to assess the effect of Fn and Fnf on the invasive ability of HCT116 and SW480 cells. The number of HCT116 and
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SW480 cells that passed through the membrane was significantly higher in the *Fn* group compared with the control group (HCT116 control, 111.50±15.16; *Fn*, 234.00±14.70, P=0.001; Fig. 3A and B; SW480 control, 500.00±44.77; *Fn*, 857.80±50.50, P=0.002; Fig. 3C and D). As expected, the number of cells in the *Fnf* group significantly decreased compared with the *Fn* group (HCT116 *Fn*, 234.00±14.70; *Fnf*, 137.30±27.53, P=0.021; Fig. 3A and B; SW480 *Fn*, 857.80±50.50; *Fnf*, 676.00±51.14, P=0.042; Fig. 3C and D). Collectively, these results suggest that L-fucose may ameliorate the pro-invasive ability of *Fn* on colon cancer cells.

**Fucose ameliorates the carcinogenic property of *Fn* to activate stat3 and EMT.** The effect of fucose on *Fn* potentiation on activation of the pathway that promotes CRC progression was assessed in the present study. Stat3 plays a crucial role in tumorigenesis, and progression and invasion of cancer cells (15). Thus, the expression and phosphorylation of stat3 were detected via western blot analysis. As presented in Fig. 4, the protein levels of p-stat3 and p-jak2 significantly increased following treatment with *Fn* (for p-stat3/stat3, HCT116 control, 1.000±0.020; *Fn*, 1.851±0.063; P=0.002; SW480 control, 1.000±0.006; *Fn*, 1.759±0.075; P=0.004. For p-jak2/jak2, HCT116 control, 1.000±0.127; *Fn*, 2.272±0.098;
P=0.016; SW480 control, 1.000±0.052; Fn, 1.400±0.057; P=0.002). However, the protein levels significantly decreased following treatment with fucose (for p-stat3/stat3, HCT116 Fn, 1.851±0.063; Fnf, 1.490±0.095; P=0.044; SW480 Fn, 1.759±0.075; Fnf, 1.318±0.031; P=0.005. For p-jak2/jak2, HCT116 Fn, 2.272±0.098; Fnf, 1.327±0.157; P=0.036; SW480 Fn, 1.400±0.057; Fnf, 1.082±0.108; P=0.040).

Stat3 activation is upstream of epithelial-to-mesenchymal transition (EMT) in CRC, which is associated with tumor progression (16). The present study assessed the specific protein markers of EMT, which exist in different types of tumors, such as prostate and colon cancer, and contribute to tumor metastasis (17). As presented in Fig. 5A and B, the expression levels of N-cadherin, β-catenin and vimentin were higher in HCT116 cells treated with Fn compared with the control group (N-cadherin control, 1.00±0.01; Fn, 1.34±0.03, P=0.007; β-catenin control, 1.00±0.02; Fn, 1.44±0.03, P=0.006; vimentin control, 1.00±0.01; Fn, 2.19±0.09; P=0.006). As expected, the expression levels of N-cadherin, β-catenin and vimentin significantly decreased following treatment with fucose (N-cadherin Fn, 1.34±0.03; Fnf, 1.20±0.01, P=0.048; β-catenin Fn, 1.44±0.03; Fnf, 1.26±0.03, P=0.042; vimentin Fn, 2.19±0.09; Fnf, 1.67±0.07, P=0.045). E-cadherin was expressed at low levels in HCT116 cells treated with Fn.
compared with the other two groups, which demonstrated the change of the EMT pathway (E-cadherin control, 1.00±0.04; Fn, 0.37±0.03; Fnf, 0.72±0.07; P=0.005 and P=0.040). Similar findings were observed in SW480 cells (Fig. 5C and D).

The stat3 inhibitor, S3I-201, was used to assess the association between stat3 activation and EMT following treatment with Fn. Western blot analysis demonstrated that inhibition of stat3 activation significantly suppressed the activation of EMT in both HCT116 and SW480 cells (Fig. 5E-H). Taken together, these results suggest that fucose inhibits the carcinogenic properties of Fn to activate the stat3 pathway and EMT.

**Discussion**

Recently, several studies have focused on the molecular mechanisms of carcinogenesis of Fn (5,18,19); however, methods for resisting the carcinogenic properties remain largely unknown. The present study revealed a distinct role of fucose in ameliorating the carcinogenic properties of Fn in *vivo*.

The results of the present study demonstrated that Fn significantly promoted proliferation, migration and invasion of colon cancer cells. Fn was first observed in oral cavity contributing to periodontal diseases (20). Recently, increasing evidence suggests an association between Fn and CRC. For example, Mima *et al* (21) reported that Fn was detected in 76/598 (13.0%) colorectal carcinomas (stages I-IV) and 19/558 (3.4%) adjacent non-tumor tissues. Furthermore, highly enriched *Fusobacterium* in CRC tissues is associated with microsatellite instability-high status (22). Fn enrichment may augment myeloid-derived immune cells in CRC, which can inhibit T-cell proliferation and induce apoptosis (23). In addition, Fn activates toll-like receptor 4 signaling and downstream PAK1 and NF-κB in CRC, thus increasing the proliferative and invasive abilities (18,19). Consistent with these findings, the results of the present study demonstrated the carcinogenic properties of Fn.

Currently, as our research team discovered that fucose is testified to competent in different types of diseases, including colitis, renal ischemia/reperfusion injury and high-fat diet-induced obesity and hepatic steatosis (10,24,25), it has also been demonstrated to impact the microbial ecosystem (11). Notably, fucose may decrease pathogen virulence through certain bacterium's metabolic pathway, such as *Salmonella enterica* (12). Thus, the present study assessed whether L-fucose can restrain the ability of Fn to promote CRC progression. The results demonstrated that Fn exhibited less tendency to facilitate the proliferation, migration and invasion of colon cancer cells following treatment with L-fucose. To the best of our knowledge, the present study was the first to demonstrate that L-fucose suppresses flora associated with colon cancer.

Mechanistically, previous studies have demonstrated that Fn induces stat3 expression in macrophages, resulting in M2 polarization and increased tumor-immune cytokine secretion, which alters the tumor microenvironment and promotes colorectal tumor development (8,26). The expression of stat3 signaling was assessed in the present study. Given that stat3 activation can promote EMT (27), and EMT is an important factor to drive carcinogenesis (16), the change in the expression levels of EMT markers was assessed in the present study. The results demonstrated that the protein expression levels of the EMT markers significantly increased following treatment with Fn, suggesting that Fn may promote EMT in colon cancer cells. In addition, activation of stat3 signaling and EMT weakened following treatment with L-fucose.

However, there were some limitations in this study. On the one hand, *in vivo* experiments could further confirm the conclusion and make the study more complete. On the other hand, further studies are needed to explore the mechanism that how fucose impair the carcinogenic properties of Fn. One hypothesis is that it may alter its metabolism.
Figure 5. Fucose ameliorates the carcinogenic properties of Fn to activate epithelial-to-mesenchymal transition. (A) Representative N-cadherin, E-cadherin, β-catenin and vimentin immunoblots of HCT116 cells. Cells were treated with supernatant of Fn or Fnf for 12 h. ACTB was used as the loading control. (B) Relative protein levels are presented in histograms. (C) Representative N-cadherin, E-cadherin, β-catenin and vimentin immunoblots of SW480 cells. Cells were treated with supernatant of Fn or Fnf for 12 h. ACTB was used as the loading control. (D) Relative protein levels are presented in histograms. (E) Representative N-cadherin, E-cadherin, β-catenin and vimentin immunoblots of HCT116 cells. Cells were treated with or without S3I-201 (100 nM) and supernatant of Fn. (F) Relative protein levels are presented in histograms. (G) Representative N-cadherin, E-cadherin, β-catenin and vimentin immunoblots of SW480 cells. Cells were treated with or without S3I-201 (100 nM) and supernatant of Fn. (H) Relative protein levels are presented in histogram. Data are presented as the mean ± standard error of the mean of at least three repeated experiments. *P<0.05, **P<0.01. Fn, Fusobacterium nucleatum; Fnf, Fn+L-fucose; Con, control.
In conclusion, the results of the present study demonstrated that L-fucose ameliorated the carcinogenic properties of Fn by suppressing its ability to activate stat3 and EMT of colon cancer cells in vitro. Thus, L-fucose may serve as a novel therapeutic strategy of microflora-related colon cancer.

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Availability of data and materials

The datasets used and analyzed during the present study are available from the corresponding author upon reasonable request.

Authors' contributions

CD performed the experiments, analyzed the data and drafted the initial manuscript. XT, WW and WQ helped analyze the data and revised the manuscript for important intellectual content. XF and XD helped culture the bacteria. XH and CH designed the present study, provided funding and obtained the grants. All authors have read and approved the final manuscript.

Ethics approval and consent to participate

Not applicable.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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