c-Fos mediates α1, 2-fucosyltransferase 1 and Lewis y expression in response to TGF-β1 in ovarian cancer

YINGYING HAO¹, LIANCHENG ZHU¹, LIMEI YAN¹, JUANJUAN LIU¹, DAWO LIU¹, NA GAO², MINGZI TAN¹, SONG GAO¹ and BEI LIN¹

¹Department of Obstetrics and Gynecology, Shengjing Hospital Affiliated to China Medical University, Shenyang, Liaoning 110004; ²Department of Obstetrics and Gynecology, First Affiliated Hospital of Dalian Medical University, Dalian, Liaoning 116011, P.R. China

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Abstract. FUT1 is a key rate-limiting enzyme in the synthesis of Lewis y, a membrane-associated carbohydrate antigen. The aberrant upregulation of FUT1 and Lewis y antigen is related to proliferation, invasion and prognosis in malignant epithelial tumors. A c-Fos/activator protein-1 (AP-1) binding site was found in the FUT1 promoter. However, the mechanisms of transcriptional regulation of FUT1 remain poorly understood. TGF-β1 is positively correlated to Lewis y. In the present study, we investigated the molecular mechanism of FUT1 gene expression in response to TGF-β1. We demonstrated that c-Fos was highly expressed in 77.50% of ovarian epithelial carcinoma cases and was significantly correlated with Lewis y. Using luciferase activity and chromatin immunoprecipitation (ChIP) assay, we further revealed that c-Fos interacted with the FUT1 promoter in ovarian cancer cells and transcriptional capacity of the heterodimer formed by c-Fos and c-Jun was stronger than that of the c-Fos or c-Jun homodimers. Then, we demonstrated that TGF-β1 induced dose-dependent c-Fos expression, which was involved in TGF-β1-induced ovarian cancer cell proliferation. In addition, inhibition of MAPK activation or TGF-β1 receptor by pharmacological agents prevented TGF-β1-induced c-Fos and Lewis y expression. Silencing of c-Fos prevented TGF-β1-induced Lewis y expression. Collectively, the results of these studies demonstrated that TGF-β1 regulated FUT1 and Lewis y expression by activating the MAPK/c-Fos pathway.

Introduction

Ovarian cancer is a common malignant tumor of the female reproductive system, and is the major cause of cancer-related deaths in women. In the US, in 2013 alone, there were 22,240 new cases of ovarian cancer and 14,030 deaths due to this disease (1). Due to the inconspicuous behavior of ovarian cancer and lack of effective measures for early diagnosis, ~75% of epithelial ovarian cancer patients are diagnosed at an advanced stage (II or IV) (2). Even with standard ovarian cancer cytoreductive surgery and chemotherapy, the 5-year survival rate is still lower than 30% due to drug resistance and relapse (3). An important feature of the malignant transformation of tumors is the change in cell surface glycosylation, which affects the function of adhesion molecules, alters mutual interactions between cells and the matrix and subsequently leads to the chemoresistance of cells (4-6).

Glycosylated proteins have been used as markers in cancer diagnosis and in evaluation of therapeutic effects (7,8). Lewis y antigen, a type 2 carbohydrate antigen, exhibited high expression in malignant epithelial tumors and was related to the prognosis of the disease (9-11).

The expression of cancer-associated carbohydrate antigens was modified by abnormal control by glycosyltransferase. FUT1 is a key enzyme for Lewis y synthesis (12-16). Overexpression of FUT1 led to a marked increase in the expression of Lewis y and promoted the proliferation, invasion (17) and drug-resistance (6,18) of ovarian cancer cells transfected with the FUT1 gene. Knockdown of FUT1 expression attenuated cell proliferation in a HER2-overexpressing cancer cell line (16). Therefore, understanding the molecular mechanism of FUT1 expression in ovarian cancer is critical for early diagnosis and searching for better treatment options.

AP-1 is a classic nucleus transcription factor, including c-Fos, Fos-B, Fra-1 and Fra-2 of the Fos family and c-Jun, Jun-B and Jun-D of the Jun family. They bind to DNA target sequences in the form of homologous or heterologous dimers, which regulate gene expression in response to a variety of stimuli, including cytokines, growth factors, bacterial and viral infections. As an important downstream target of the MAPK signaling pathway, AP-1 is essential for normal cell differentiation and survival. However, overexpression of c-Jun and c-Fos also promoted the malignant transformation of cells (19). Previous investigations revealed that silencing of c-Fos sensitized glioma cells to radiation and c-Fos overexpression was correlated with poor prognosis in malignant...
normal ovarian tissue (resected from cervical cancer patients, 30 cases of benign ovarian tumors, and 20 cases of clear-cell carcinoma), 30 cases of borderline ovarian epithelial, 10 cases of ovarian endometrioid carcinoma and 10 cases of adenocarcinoma and 30 cases of serous cystadenocarcinoma, epithelial ovarian cancer (including 30 cases of mucinous cysto
from China Medical University (Shenyang, China), from paraffin specimens obtained from the Department of Collection of human samples.

Materials and methods

Collection of human samples. We selected 160 resected paraffin specimens obtained from the Department of Gynecology and Obstetrics of Shengjing Hospital Affiliated to China Medical University (Shenyang, China), from 2000 to 2013. All tissues were re-diagnosed by pathologists. According to the pathological results, there were 80 cases of epithelial ovarian cancer (including 30 cases of mucinous cystadenocarcinoma and 30 cases of serous cystadenocarcinoma, 10 cases of ovarian endometrioid carcinoma and 10 cases of clear-cell carcinoma), 30 cases of borderline ovarian epithelial tumors, 30 cases of benign ovarian tumors, and 20 cases of normal ovarian tissue (resected from cervical cancer patients during the same period). The mean age of the patients was 47 years (15-73 years) and the median age was 44 years. The age of ovarian cancer patients ranged from 38 to 83 years with a median age of 51 years. Borderline patients ranged from 22 to 55 years with a median age of 35 years. Benign patients ranged from 15 to 72 years with a median age of 44 years. Normal ovarian patients ranged from 37 to 52 years with a median age of 42 years. The age difference between patients in the different groups was not significant (P>0.05). The ovarian cancer group was pathologically classified as follows, 41-well-differentiated cases, 18 moderately differentiated cases and 21 poorly differentiated cases. Pathology stage: according to the standards of the International Federation of Gynecology and Obstetrics: 56 cases were stage I and II; 24 cases were stage III and IV. Of these, 15 cases had pelvic lymph node metastasis. All cases were primary tumors with complete clinical and pathological data and without preoperative radiotherapy or chemotherapy. The present study was approved by the hospital Ethics Committee.

Ethical approval. All procedures performed in the study involving human participants were in accordance with the Ethical Standards of the Institutional and/or National Research Committee and with the 1964 Helsinki Declaration and its later amendments or comparable ethical standards.

Cell culture. The human ovarian cancer cell lines CAVO3, SKOV3, ES-2 and human embryonic kidney 293 (293) were purchased from the Shanghai Institute of Life Sciences of the Chinese Academy of Sciences. The cells were conventionally cultured in Roswell Park Memorial Institute formulation (RPMI)-1640 and McCoy's 5A medium (Gibco by Invitrogen) with 10% fetal bovine serum (HyClone, Logan, UT, USA).

Transient transfection and luciferase assay. A luciferase reporter vector of the FUT1 promoter, pGL4-FUT1, including one binding site for AP-1 (-3,000 to -1) was constructed as previously described (22). Transfections were performed using Lipofectamine 2000 (Invitrogen, Carlsbad, CA, USA) according to the manufacturer's instructions. Cells were transferred to 24-well plates at a density of 4x10^4 cells/well 1 day before transfection. pcDNA3-c-Fos, pCMV6-c-Jun, pcDNA3-c-Fos/pCMV6-c-Jun or vector pCMV6 and pcDNA3 vector (1 µg/ml) (OriGene, Beijing, China) along with the human FUT1 promoter reporter gene were co-transfected in 293, SKOV3, CAV03 and ES-2 cells. Forty hours after transfection, the activity of the promoters was detected by the Dual-Luciferase Assay System (Promega, Fitchburg, Wl, USA). The luciferase data of the FUT1 promoter reporter gene was detected by the Dual-Luciferase Assay System (Promega, Fitchburg, WI, USA). The luciferase data of the FUT1 promoter reporter gene was calculated and normalized with Renilla luciferase activity. All transfections were carried out in triplicate.

Immunohistochemistry analysis. Histological sections from each group of ovarian tissues was 4 µm. Each tissue had two serial sections. The expression of c-Fos and Lewis y in paraffin sections were detected via immunohistochemical strepavidin-peroxidase staining. The sections were dewaxed and rehydrated by rinsing with xylene followed by graded
ethanol washing. Endogenous peroxidase activity was blocked with 3% hydrogen peroxide in methanol for 15 min before antigen retrieval by high pressure treatment in 10 mmol/L citrate phosphate buffer for 1.5 min. The sections were then incubated with a protein block in 10% normal goat serum for 10 min. The sections were incubated overnight at 4°C with polyclonal rabbit anti-human c-Fos (1:200; Santa Cruz Biotechnology, Inc., Santa Cruz, CA, USA) and anti-Lewis y antibodies (1:200; Abcam, Cambridge, UK). After being washed with phosphate-buffered saline (PBS), the sections were incubated with horseradish peroxidase-labelled secondary antibodies for 2 h at 37°C. Two cases of strongly positive sections and 2 cases of negative sections served as the positive control and negative control for c-Fos and Lewis y antigen, respectively. Rabbit IgG (BIOS, Beijing, China) was used as the negative control. In addition, the blank control was incubated with PBS instead of a primary antibody. The empirical procedure was performed base on the manufacturer's protocol of UltraSensitive™ S-P (mouse/rabbit) IHC and DAB kits (both from MaiXin Bio, Fuzhou, China).

Five high power fields (magnification, x400) were randomly selected in each section according to the staining intensity and the percentage of positive cells. The degree of staining was defined as follows: 0 points represented no staining; 1 point represented faint yellow; 2 points represented yellowish brown; 3 points represented brown. Percentage of stained cells was as follows: 0 points represented <5%; 1 point represented 5-25%; 2 points represented 26-50%; 3 points represented 51-75%; 4 points represented >75%. When both scores were multiplied: 0-2 indicated negative (-); 3-4 indicated weakly positive (+); 5-8 indicated moderately positive (++) and 9-12 indicated strongly positive (+++).

Chromatin immunoprecipitation (ChIP) assay. ChIP experiments were conducted according to the ChIP kit instructions (Upstate, Charlotteville, VA, USA). CAVO3 cells were fixed in 1% formaldehyde for 10 min for crosslinking reaction which was quenched with 125 mM glycine. The nuclei were pelleted by centrifugation at 3,000 rpm for 5 min at 4°C and sonicated to chromatin fragments between 100 and 1,000 bp. The sonicated lysate was centrifuged at 10,000 rpm for 5 min at 4°C. Supernatant (500 µl) was incubated with anti-c-Jun (1:50; Cell Signaling Technology, Inc., Danvers, MA, USA), anti-c-Fos (1:50) antibody followed by an isolation procedure using protein A/G magnetic beads (both from Santa Cruz Biotechnology, Inc.). A normal rabbit IgG was used as a control. The crosslinking was reversed by incubation at 65°C for 10 h. Primers were designed according to the binding site of AP-1 in the FUT1 promoter and the sequence was as follows: F, 5'-CTAGCCTCAAGGTCTTGTC-3' and R, 5'-GCAAGATGAGGAAACTGTGC-3'. The PCR conditions were as follows, 98°C for 5 min, 98°C for 30 sec, 60°C for 20 sec and 72°C for 5 min for 30 cycles. PCR products were resolved by electrophoresis on a 1% agarose gel and visualized after ethidium bromide staining.

Western blotting. Protein was extracted with lysis buffer [150 mM NaCl, 1% w/v NP-40, 0.1% w/v SDS, 2 µg/ml aprotinin, 1 mM phenylmethylsulfonyl fluoride (PMSF)]. Protein (50 µg) was subjected to 10% sodium dodecyl sulphate-polyacrylamide gel electrophoresis (SDS-PAGE), and then transferred to nitrocellulose membranes. The membranes were blocked for 2 h at room temperature with 5% non-fat dry milk in Tris-buffered saline containing 0.1% Tween-20 (TBS-T) and subsequently incubated overnight at 4°C with the primary antibodies: anti-c-Fos (1:500; Santa Cruz Biotechnology, Inc.), β-actin (1:20,000; Sigma, St. Louis, MO, USA), anti-Lewis y (1:100; Abcam), anti-phospho-p38 (1:500), anti-total p38 (1:500), anti-phospho-JNK (1:500), anti-total JNK (1:500), anti-phospho-ERK (1:500), anti-total ERK (1:500) (all from Cell Signaling Technology, Inc.). The samples were then washed 3 times for 15 min with TBS-T and incubated with appropriate horseradish peroxidase-conjugated IgG (1:5000; Sigma) for 2 h at 37°C. The immunocomplex bands were detected with an enhanced chemiluminescence HRP substrate for western blotting (Pierce, Rockford, IL, USA) using the Molecular Imager system GDS8000b (UVP, Upland, CA, USA).

Interfering RNA transfection. When the cells reached 60% confluence, scramble siRNA, c-Jun siRNA and c-Fos siRNA obtained from Santa Cruz Biotechnology, Inc. were transfected into the cell line, respectively, using Lipofectamine RNA iMAX transfection reagent (Invitrogen). After 72 h of transfection, proteins were harvested for subsequent tests.

Cell proliferation assay. Twenty-four hours after CAVO3 cells had been transfected with pcDNA3-c-Fos, pcDNA3 or c-Fos siRNA, the transfected cells were inoculated into a 96-well plate at a density of 2x10³ cells/ml (100 µl/well). Methyl thiazolyl tetrazolium (MTT; 20 µl) (5 mg/ml) was added 24, 48, 72 and 96 h later and after another 4 h. Dimethyl sulfoxide solution (150 µl) was added to each well to dissolve the crystals. The absorbance in each well was determined with enzyme-linked immunosorbent analyzer at 550 nm.

Colonies formation. Cells in the logarithmic growth phase were titrated into single cells and inoculated into a 6-cm culture dish (200 cells/well) and the medium was discarded after 24 h when the cells adhered. The c-Fos interference, c-Fos overexpression and the corresponding control group were established. Using the control group as the reference standard, the original culture medium was discarded when visible cell clusters in the control group reached 50 cells under the microscope, and the cells were fixed using methanol. An appropriate amount of Giemsa pigment was added for 5 min to stain the nucleus. The dish was then inverted, placed under the microscope for clone counting, and cell clusters containing >50 cells were considered one clone, followed by calculation of the clone formation rate.

Statistical analysis. Quantitative data were expressed in terms of the means ± SD. Qualitative data were expressed in terms of the composition ratio. SPSS 16.0 software (SPSS, Inc., Chicago, IL, USA) was used for statistical analysis. The Pearson χ² and likelihood ratio tests were used to analyze the correlation between the expression of Lewis y and c-Fos and clinicopathological factors. Spearman rank correlation analysis was employed to analyze the correlation between Lewis y and c-Fos. A P-value <0.05 indicated statistical significance.
Results

Overexpression of c-Fos in ovarian epithelial carcinoma and its correlation with clinicopathological characteristics.

To determine the role of c-Fos in the progression of ovarian epithelial carcinoma, we examined c-Fos expression and its correlation with clinicopathological characteristics in 140 primary ovarian tumor samples and 20 normal ovarian tissue samples using immunohistochemistry. The 140 primary ovarian tumor samples included 30 mucinous and serous cystadenoma (benign), 30 borderline mucinous and borderline serous cystadenoma (borderline), 60 mucinous and serous cystadenocarcinoma, 10 endometrioid and 10 clear cell carcinoma (malignant). We observed that c-Fos was hardly expressed in the benign (Fig. 1A and B) and normal ovarian tissues (data not shown), only expressed in the nucleus of borderline mucinous cystadenoma (Fig. 1C), moderately expressed in the nucleus and the cytoplasm of borderline serous cystadenoma (Fig. 1D) and significantly overexpressed in both the nucleus and the cytoplasm of different malignant ovarian epithelial tumor cells, including mucinous (Fig. 1E) and serous cystadenocarcinoma (Fig. 1F), endometrioid (Fig. 1G) and clear cell carcinoma (Fig. 1H). The degree of the staining were defined as follows, negative (-), weakly positive (+), moderately positive (++) and strongly positive (+++) (see Materials and methods for details) and the cases were further divided into high (++) or (+++) and low (- or +) c-Fos expression groups. The expression rate of c-Fos was positively associated with the degree of ovarian tumor malignancy (P<0.05) (Table I).

The high expression rates of c-Fos in mucinous and serous ovarian cystadenocarcinoma were 83.33% (25/30) and 80% (24/30), respectively, which were higher than that in ovarian endometrioid carcinoma and clear cell carcinoma.
(60%, 6/10; and 70%, 7/10), but no significant difference was detected (P>0.05). The high expression rates of c-Fos in ovarian cancer tissues with high, middle and low differentiation were 63.41% (26/41), 88.89% (16/18) and 95.24% (20/21), respectively, and the high expression rate of c-Fos gradually increased with the decreasing degree of differentiation (P<0.05). The expression of c-Fos in advanced cancer (stage III-IV) was 91.67% (22/24), which was significantly higher than that in early-stage ovarian cancer (stage I-II) (71.43%, 40/56) (P<0.05). The expression rate of c-Fos was not correlated with lymph node metastasis (P>0.05) (Table II).

The expression rates of Lewis y in mucinous (Fig. 2E) and serous ovarian cancer tissues (Fig. 2F) were 86.67% (26/30) and 90% (27/30), respectively, which were higher than those in ovarian endometrioid (60%, 6/10) and clear cell carcinoma (80%, 8/10), but the difference was not statistically significant (P>0.05). The expression rate of Lewis y in high, middle and low differentiation ovarian cancer were 75.61% (31/41), 83.33% (15/18) and 100% (21/21), respectively, with the high expression rate significantly increased with decreasing degree of differentiation (P<0.05). The expression of Lewis y was not correlated with lymph node metastasis (P>0.05) (Table II).

| Characteristics | Lewis y Low n (%) | Lewis y High n (%) | P-value |
|-----------------|------------------|-------------------|---------|
| Malignant group (n=80) | 18 (22.50) | 62 (77.50) | <0.01<sup>a,b,c</sup> |
| Borderline group (n=30) | 20 (66.67) | 10 (33.33) | 0.013<sup>d</sup>, 0.091<sup>e</sup> |
| Benign group (n=30) | 25 (83.33) | 5 (16.67) | 0.687<sup>f</sup> |
| Normal group (n=20) | 18 (90.00) | 2 (10.00) | |

<sup>a</sup>Compared with borderline group. <sup>b,c</sup>Compared with benign group. <sup>d,e,f</sup>Compared with normal tissue group. n, the number of cases.
Of the 80 cases with ovarian cancer, 55 cases had high expression of both c-Fos and Lewis y, and 6 cases had double low expression. The expression of c-Fos and Lewis y in ovarian

Table III. Expression of Lewis y in different ovarian tissues.

| Characteristics   | Low n (%) | High n (%) | P-value |
|-------------------|-----------|------------|---------|
| Malignant group (n=80) | 16 (16.25) | 84 (83.75) | <0.01<sup>a,b,c</sup> |
| Borderline group (n=30) | 12 (40.00) | 18 (60.00) | 0.038<sup>d</sup>, <0.01<sup>e</sup> |
| Benign group (n=30) | 20 (66.67) | 10 (33.33) | 0.003<sup>f</sup> |
| Normal group (n=20) | 20 (100.00) | 0 (0.00) |         |

<sup>a</sup>Compared with borderline group. <sup>b</sup>Compared with benign group. <sup>c</sup>e<sup>f</sup>Compared with normal tissue group. n, the number of cases.

Table IV. Correlation between Lewis y and c-Fos in ovarian carcinoma.

| Lewis y expression | Low (n) | High (n) | P-value | Correlation coefficient |
|--------------------|---------|----------|---------|-------------------------|
| c-Fos expression   | 6       | 12       | 0.026<sup>g</sup> | 0.250                  |
| c-Fos expression   | 7       | 55       |         |                         |

<sup>g</sup>Indicate statistical significance. n, the number of cases.
cancer exhibited significant correlation, with a correlation coefficient of 0.250, P<0.05 (Table IV).

**Binding of c-Fos to TPA response element (TRE) of the FUT1 promoter enhances the activation of FUT1 transcription by c-Jun.** We previously constructed the FUT1 (-3,000 to -1) promoter luciferase reporter gene vector, which had one binding site for AP-1, and revealed that c-Jun transactivated FUT1 via the AP-1 binding site (22). The FUT1 promoter activity was first tested in 293 cells, and further confirmed in 3 ovarian cancer cell lines SKOV3, CAVO3 and ES-2, with the strongest enzyme activity achieved in 293 cells (Fig. 3A). Furthermore, to investigate the effect of c-Fos in the regulation of FUT1 transcription, we co-transfected ovarian cancer cells with the transcription factor expression vector c-Fos, c-Jun, c-Fos/c-Jun or empty vector (pCMV6 and pcDNA3) along with the human FUT1 promoter reporter gene. Compared with the empty vector transfection, c-Fos did not significantly affect the activity of the FUT1 promoter, whereas c-Jun transcription factor expression vector increased promoter activity. Compared with the single transfection of c-Jun expression vector, co-transfection with c-Fos and c-Jun expression vector significantly increased promoter activity, and the activity was increased 3.5-fold in SKOV3 cells (Fig. 3B). These results revealed that the transcriptional activation ability of the heterodimer formed by c-Fos and c-Jun was significantly enhanced in comparison with that of the homodimer with two c-Juns.

To determine whether c-Fos and c-Jun interacted with TRE of the FUT1 promoter (-1,908 to -1,914), ChIP assay was performed by transfection of c-Fos or c-Jun siRNA into CAVO3 cells. MNase cleavage was used to cut cell chromatin into 100-1,000 bp fragments, with an indwelling portion of the cell suspension used as the DNA input positive control, rabbit IgG precipitation as the negative control. In the input and c-Fos, c-Jun-specific antibody precipitation groups, PCR amplification obtained FUT1 promoter fragments of 147 bp, while the corresponding fragment was not amplified in the IgG precipitation group (Fig. 3C). The precipitated FUT1 promoter fragment expression was significantly decreased after c-Fos or c-Jun siRNA interference. These results revealed that c-Fos and c-Jun were bound to the FUT1 promoter region containing the AP-1 binding site.

**TGF-β1 activates c-Fos via the MAPK signaling pathway, regulates FUT1 transcription, and promotes Lewis y expression.** TGF-β1 is positively correlated to Lewis y in ovarian cancer (35). To determine whether TGF-β1 increased AP-1 expression, we assessed c-Jun and c-Fos expression in response to TGF-β1 in 3 ovarian cancer cell lines. The ovarian cancer cells SKOV3, CAVO3 and ES-2 were treated with various concentrations of TGF-β1 (0.2, 1, 2, 5 and 10 ng/ml) for 5 min,
Figure 4. TGF-β1 promotes cell proliferation via c-Fos in ovarian cancer cells. (A) Cell viability was detected using MTT cell proliferation assay in CAVO3 cells. CAVO3 cells were transfected with pcDNA3-c-Fos, pcDNA3 or c-Fos siRNA for 24 h, followed by pretreatment with TGF-β1. (B and C) Colony formation assessed cell proliferation ability. The cells and processing were the same as shown in a. The data are expressed as the mean ± SD of 3 independent experiments; *P<0.05, compared with the control.

Figure 5. TGF-β1 activates c-Fos through the MAPK/AP-1 pathway. Upon TGF-β1 treatment, western blotting was employed to determine c-Jun and c-Fos expression in 3 ovarian cells. (A) SKOV3, (B) CAVO3, (C) ES-2. c-Fos and c-Jun expression is related to TGF-β1 in a concentration manner in ovarian cells pretreated with various concentrations of TGF-β1 (0.2, 1, 2, 5 and 10 ng/ml) for 5 min. (D) Western blot analysis was carried out to determine the expression of c-Fos, Lewis y and the downstream signal elements of the MAPK signaling pathway, JNK, p38 and ERK, and the change in the phosphorylation level of JNK, p38 and ERK. CAVO3 cells were treated with pcDNA-c-Fos, c-Fos siRNA, inhibitor of TGF-β1 receptor (LY2109761) and JNK (SPF600125) upon TGF-β1 stimulation.
and subsequently the change in c-Jun and c-Fos expression was detected by western blotting (Fig. 5A-C). TGF-β1 induced the expression of c-Jun and c-Fos in all 3 types of cells. In CA VO3 and SKOV3 cells, the expression of c-Fos and c-Jun was strongly dependent on the concentration of TGF-β1 used, and was highest when 10 ng/ml TGF-β1 was added. These results indicated that TGF-β1 induced the expression of c-Fos and c-Jun in ovarian cancer cells in a dose-dependent manner.

TGF-β1 and the members of TGF-β-independent MAPK signaling pathway act as key determinants of carcinoma cell behavior. AP-1 transcriptional activity is regulated by the activation of the MAPK signaling pathway, involving extracellular-regulated kinase (ERK), c-Jun N-terminal kinase (JNK) and p38 pathways (38,39). To study the role of MAPK pathway in mediating TGF-β1-induced Lewis y expression, we applied specific JNK inhibitor (SP600125) and TGF-β receptor inhibitor (LY2109761). Upon TGF-β1 stimulation, CA VO3 was treated with LY2109761 and SP600125, respectively. Western blot analysis revealed that the expression of c-Fos, Lewis y, the phosphorylation levels of p38, JNK and ERK decreased significantly (Fig. 5D).

Furthermore, to address whether the increased expression of c-Fos contributed to Lewis y expression, we transfected CA VO3 cells with c-Fos and c-Fos-siRNA upon TGF-β1 stimulation. c-Fos transfection promoted TGF-β1-induced Lewis y expression and phosphorylated (p)-p38 and p-JNK. Consistent with these results, the silencing of c-Fos prevented TGF-β1-induced Lewis y expression and also suppressed p-p38 and p-JNK (Fig. 5D). Whereas, silencing or overexpression of c-Fos did not exert any effect on p-ERK. These results indicate that as a downstream effector of MAPK pathway, c-Fos is also capable of modulating p-p38 and p-JNK, but not p-ERK in response to TGF-β1. Collectively, our findings support that MAPK pathway plays a critical role in the mechanism of TGF-β1-induced Lewis y expression in ovarian cells (Fig. 6).

Discussion

Lewis y antigen, a tumor-associated carbohydrate antigen, is overexpressed in malignant epithelial tumors and closely associated with the prognosis of the disease. Previously, in ovarian cancer cells, we identified an AP-1 binding site in the promoter of FUT1, which was a rate-limiting enzyme of Lewis y. AP-1, a dimer composed of Fos (c-Fos, Fos-B, Fra-1 and Fra-2) and Jun (c-Jun, Jun-B and Jun-D) family members, modulates gene transcription and cellular pathophysiological functions. It has been demonstrated that MMP9 promoter was activated by the IL-1β/p38/c-Fos pathway, which was correlated with gastric adenocarcinoma metastasis (40). Furthermore, c-Fos overexpression enhanced the epithelial-mesenchymal transition (EMT) state in head and neck squamous cell carcinoma (41). In the present study, we observed high levels of expression of c-Fos in 140 ovarian epithelial tumor cases. Overexpression of c-Fos in ovarian cancer was as high as 77.5% (62/80) and its expression rate and intensity increased with decreasing degree of differentiation. In malignant ovarian epithelial tumors, high expression of c-Fos was observed in both the nucleus and the cytoplasm. c-Fos expression was only detected in the nucleus in benign and normal ovarian tissues. These findings indicated that c-Fos expression was aberrantly upregulated in malignant ovarian tumors and significantly correlated with malignant behaviors.

Figure 6. Pathway of c-Fos-mediated Lewis y in response to TGF-β1. TGF-β1 activates c-Fos/AP-1 through the MAPK pathway. As a transcriptional factor, c-Fos together with c-Jun upregulated FUT1 expression by binding to its promoter, consequently leading to the enhanced synthesis of Lewis y.
TGF-β1 plays an important role in carcinogenesis and is associated with the proliferation and progression in advanced malignancies. TGF-β1 promotes tumor growth via regulating downstream targets of its signaling pathway. It was reported that c-Fos overexpression is positively correlated with carcinoma growth (42). Knockdown of c-Fos greatly suppressed tumor cell proliferation and invasion and downregulated CD44 and cyclin D1 expression (43). In the present study, we examined whether c-Fos was involved in TGF-β1-induced cell proliferation in ovarian cancer cells. These findings indicated that c-Fos expression was increased and contributed to the TGF-β1-induced tumor promoting effect in ovarian epithelial cells.

We previously revealed that c-Jun transactivated FUT1 through the AP-1 binding site in ovarian cancer cells (22). Thus, we wondered how c-Fos plays its role and how do the different components of AP-1 interact with each other in the regulation of FUT1 transcription. The underlying mechanism is still unclear. To answer this question, we performed ChIP assays to determine whether c-Fos and c-Jun bind to the FUT1 promoter. Furthermore, using luciferase activity analysis, we observed that the homodimer of c-Fos hardly influenced the activity of FUT1 promoter. Additionally, using luciferase activity analysis, we observed that the homodimer of c-Fos hardly influenced the activity of the FUT1 construct, suggesting that c-Fos activation alone was insufficient for the activation of FUT1 promoter activity. The activity of FUT1 promoter interacted with heterodimer of c-Fos and c-Jun increased 3.5-folds, compared with interaction with c-Jun alone, further illustrating the interaction and coordination between c-Fos and c-Jun. It was reported that c-Fos could not bind to TRE, but the DNA binding force of the heterodimer formed by c-Fos and c-Jun was 25 times that of the homodimer with the two c-Juns (44). However, DebRoy et al reported that the p38 pathway induced stoma interaction molecule 1 (STIM1) expression through c-Fos rather than c-Jun in endothelial cells (45). Collectively, these results revealed that c-Fos and c-Jun are essential in regulating FUT1 promoter activity in ovarian cancer cells.

In the present study it was also demonstrated that in the absence of TGF-β1 or stimulation with low concentrations of TGF-β1, the c-Fos protein expression level in ovarian cancer cells was significantly lower than that of c-Jun, however with increased TGF-β1 concentration, the protein expression levels of c-Fos significantly increased, reaching or even exceeding the expression level of c-Jun. The protein concentration and activity of c-Fos are extremely low under normal circumstances, while c-Jun has strong potential transcriptional competence. When cells are stimulated, levels of c-Fos protein are temporarily and quickly increased. Fos/Jun heterodimers instead of stable homodimers are formed, with rapid increased DNA binding and transcription-inducing ability. Following degradation of c-Fos which has a short half-life, AP-1 returns to basal levels and an inert state (46,47). Previous studies demonstrated that c-Fos was highly expressed in cervical cancer, and almost never expressed in normal tissues and precancerous lesions (48). c-Fos expression was upregulated following TGF-β1 treatment in immortalized liver cells, which was correlated with cell migration and invasion (49). Therefore, we proposed that increased TGF-β1 promoted expression of c-Fos and the formation of heterodimers, and upregulated FUT1 transcription. Inflammatory cytokines regulated the expression of FUTs involved in the biosynthesis of tumor-associated sialylated Lewis antigen in pancreatic cancer cells (50). To elaborate whether c-Fos is activated with TGF-β1 treatment and whether c-Fos promotes FUT1 and Lewis y synthesis in ovarian epithelial cancer, we focused on the MAPK signaling pathway. Inhibition of the JNK pathway with pharmacologic inhibitor resulted in marked reduction in c-Fos and Lewis y expression in ovarian cancer cells upon TGF-β1 stimulation. The JNK pathway inhibitor not only suppressed JNK phosphorylation, but also downregulated the phosphorylation level of p38 and ERK. Therefore, the transcriptional regulation is likely to involve the crosstalk between the JNK, p38 and ERK pathways in response to TGF-β1 stimulation. To determine whether there is a relationship between c-Fos and Lewis y expression, we knocked down c-Fos and examined the resultant expression of Lewis y. The result revealed that Lewis y expression was positively related to c-Fos expression. In addition, we observed that knockdown of c-Fos also diminished the phosphorylation of JNK and p38. It has been reported that TGF-β1 promoter regions had an AP-1 binding site, where c-Jun and c-Fos are essential (51-54). TGF-β1 can activate its own mRNA transcription, promoting its own secretion. The expression of TGF-β1 was significantly increased in osteosarcoma tissues of c-Fos transgenic mice and c-Fos transfected osteosarcoma cell lines (55). These data indicated that there may be mutual regulation between TGF-β1 and AP-1 through the JNK and p38 pathways. As our results revealed, the MAPK pathway plays a vital role in Lewis y expression induced by c-Fos in response to TGF-β1.

In conclusion, we demonstrated overexpression of c-Fos and its positive correlation with Lewis y in ovarian epithelial carcinoma. The transcriptional activity of the heterodimer formed by c-Fos and c-Jun was stronger than that of the homodimer with two c-Juns in FUT1 promoter activation. Notably, we revealed that TGF-β1 activated c-Fos via the MAPK pathway, promoting FUT1 transcription, and enhancing Lewis y biosynthesis in ovarian cancer cells.

In conclusion, the results of our studies in combination with those of other researchers, suggest that in ovarian cancer cells, TGF-β1 activates c-Fos through the MAPK signaling pathway, regulates FUT1 transcription, and eventually promotes Lewis y expression.

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