

# Tensin 4 facilitates aerobic glycolysis, migration and invasion of colorectal cancer cells through the $\beta$ -catenin/c-Myc signaling pathway

YAN WANG<sup>\*</sup>, YONGDA LU<sup>\*</sup> and CHUNFANG XU

Department of Gastroenterology, The First Affiliated Hospital of Soochow University, Suzhou, Jiangsu 215000, P.R. China

Received December 19, 2023; Accepted May 8, 2024

DOI: 10.3892/ol.2024.14489

**Abstract.** Tensin 4 (TNS4) is overexpressed in multiple cancers, including colorectal cancer (CRC), and is associated with a poor prognosis of patients with CRC. However, the role and underlying mechanisms of TNS4 in CRC have yet to be elucidated. The expression of TNS4 in CRC tissues were analyzed by immunohistochemistry. Cell migration and invasion were assessed *in vitro* using Transwell assay. Western blot and reverse transcription (RT)-quantitative (q)PCR were used to investigate the molecular mechanisms by which TNS4 regulates aerobic glycolysis, migration and invasion of CRC cells. The present study demonstrated that TNS4 was highly expressed in the cancer tissues of patients with CRC and significantly associated with the tumor-node-metastasis stages. TNS4 silencing led to a significant decrease in glucose consumption and lactate production in CRC cells, and knockdown of TNS4 suppressed the migration and invasion of CRC cells via aerobic glycolysis through the  $\beta$ -catenin/c-Myc pathway. Notably, treatment with DASA-58, an activator of glycolysis, or SKL2001, an activator of  $\beta$ -catenin/c-Myc signaling, significantly reversed the effect of TNS4 knockdown on aerobic glycolysis, migration and invasion of CRC cells. Collectively, these results suggest that TNS4 may act as a novel regulator of aerobic glycolysis, migration and invasion of CRC cells by modulating  $\beta$ -catenin/c-Myc signaling, providing a new potential biomarker and therapeutic target in CRC.

## Introduction

As colorectal cancer (CRC) is one of the most commonly diagnosed malignancies and the leading cause of cancer-related

mortality worldwide with >1.85 million cases and 850 000 deaths annually, it has become a major public health concern. In recent years, the 5-year survival rate of patients with CRC has markedly improved with the development of treatment strategies, such as surgery, radiotherapy, chemotherapy and immunotherapy (1,2); however, the effects of the current treatment methods are limited for patients with advanced-stage or metastatic CRC (3). Hence, the identification of novel targets and the further elucidation of the mechanisms underlying the progression of CRC is of utmost urgency.

Tensin 4 (TNS4), also known as COOH-terminal tensin-like, belongs to the tensin focal adhesion family (4). A number of studies have reported that TNS4 is overexpressed in several types of cancer, including breast cancer, pancreatic cancer, lung cancer and CRC (5-8). Furthermore, TNS4 has been reported to be associated with several biological processes, such as epithelial-mesenchymal transition (EMT) (9-11), cell motility (6,12), cell migration (13), metastasis (8) and drug resistance (14,15). For example, TNS4 overexpression promotes EMT, cell motility and the colony formation of CRC cells through Src signaling (9). Liao *et al* (16) noted that TNS4 interacted with  $\beta$ -catenin in colon cancer cells, which enhanced the colony formation, anchorage-independent proliferation and invasiveness of colon cancer cells.

Aerobic glycolysis, also known as the Warburg effect, is a distinctive hallmark of cancer, which confers a proliferation advantage on cancer cells by providing them with energy and biosynthesis building blocks (17). Accumulating evidence has indicated that targeting or modulating aerobic glycolysis may serve as an antitumor therapeutic strategy (18). For example, a previous study reported that the long non-coding (lnc)RNA AGPG promoted tumor cell glycolysis and the proliferation of esophageal squamous cell carcinoma cells by stabilizing PFKFB3 (19). Furthermore, Yu *et al* (20) reported that OTU deubiquitinase, ubiquitin aldehyde binding 2 suppressed CRC cell proliferation and migration and promoted apoptosis and sensitivity to chemotherapeutic drugs by regulating pyruvate kinase M (PKM)2 ubiquitination and glycolysis. However, the role of TNS4 in the glycolysis and progression of CRC remains unclear.

The present study aimed to explore the roles of TNS4 in regulating aerobic glycolysis, migration and invasion of CRC cells and investigate the underlying molecular mechanisms.

---

*Correspondence to:* Professor Chunfang Xu, Department of Gastroenterology, The First Affiliated Hospital of Soochow University, 188 Shizi Road, Suzhou, Jiangsu 215000, P.R. China  
E-mail: xcf601@163.com

<sup>\*</sup>Contributed equally

**Key words:** colorectal cancer, Tensin 4, aerobic glycolysis, migration, invasion,  $\beta$ -catenin

## Materials and methods

**Analysis of TNS4 expression in patients with CRC.** The expression of TNS4 in patients with colon adenocarcinoma or rectum adenocarcinoma was obtained from the University of Alabama at Birmingham Cancer data analysis Portal (UALCAN; <http://ualcan.path.uab.edu/>) and Gene Expression Profiling Interactive Analysis (GEPIA; <http://gepia.cancer-pku.cn/>) databases with data from The Cancer Genome Atlas (TCGA).

**Immunohistochemistry.** A total of 92 pairs of paraffin blocks of CRC tissues and corresponding normal adjacent tissues were obtained from the First Affiliated Hospital of Soochow University (Suzhou, China). The detailed clinicopathological information of the patients is provided in Table SI. Ethics approval was obtained from the Institutional Review Board of the First Affiliated Hospital of Soochow University and written informed consent was obtained from the patients prior to sample collection. Immunohistochemistry was performed as previously described (21). Briefly, tissues were fixed with 4% paraformaldehyde (Beyotime Institute of Biotechnology) at 25°C for 24 h. The specimens were embedded in paraffin. 5- $\mu$ m-thick sections from paraffin-embedded blocks were deparaffinized and rehydrated. After antigen retrieval with 10 mM sodium citrate buffer (pH 6.0, Beyotime Institute of Biotechnology), the sections were incubated with 3% hydrogen peroxide at room temperature for 10 min to block endogenous peroxidase activity and non-specific protein interactions. The sections were subsequently incubated with rabbit anti-human TNS4 antibodies (1:200; cat. no. 11580-1-AP; Proteintech Group, Inc.) overnight at 4°C and then with biotinylated goat anti-rabbit secondary antibody working solution (1:500; cat. no. SA1020; Boster Biological Technology Co. Ltd.) at 37°C for 30 min. Subsequently, the immunodetection was performed using the Dako EnVision detection system (Agilent Technologies, Inc.). These slides were photographed under a fluorescence microscope (Leica, Buffalo Grove, USA). The semi-quantitative immunoreactive score system was adopted to obtain the score of TNS4 immunostaining, as previously described (22).

**Cells and cell culture.** In total, two CRC cell lines (HCT116 and RKO) were purchased from the American Type Culture Collection. Both the HCT116 and RKO cells were cultured in DMEM (Biological Industries; Sartorius AG) containing 10% fetal bovine serum (FBS, Biological Industries; Sartorius AG), 100 U/ml penicillin and 100 mg/ml streptomycin at 37°C in a humidified atmosphere of 5% CO<sub>2</sub>. The CRC cells were treated with glycolysis activator DASA-58 (5  $\mu$ M, Selleck) and  $\beta$ -catenin activator SKL2001 (10  $\mu$ M, Selleck) at 37°C for 48 h.

**Cell transfection and infection.** In total, three commercial TNS4 small interfering (si)RNAs (TNS4 siRNA-1, TNS4 siRNA-2 and TNS4 siRNA-3) were purchased from Guangzhou RiboBio Co., Ltd. The three pairs of synthesized siRNA sequences were as follows: siRNA-1 forward, 5'-CAAUCAAGAGAAGACCATT-3' and reverse, 5'-UGGUCUCUUCUAUGAUUGTT-3'; siRNA-2 forward, 5'-GGGCCAUCUCUCUGUGAUUTT-3' and reverse, 5'-AAUCACAGAGAGAUGGCCCTT-3'; siRNA-3 forward, 5'-GCAAUGACC

UCAUCCGACATT-3' and reverse, 5'-UGUCGGAUGAGG UCAUUGCTT-3'; and siRNA negative control forward, 5'-UUCUCCGAACGUGUCACGUTT-3' and reverse, 5'-ACGUGACACGUUCGGAGAATT-3'. For cell transfection, the HCT116 or RKO cells were transfected using Lipofectamine 2000® (Invitrogen™; Thermo Fisher Scientific, Inc.) according to the manufacturer's instructions with TNS4 siRNA-1, TNS4 siRNA-2 or TNS4 siRNA-3 (100 pmol) at 37°C for 48 h. Scrambled siRNA negative control (100 pmol) was used. After 48 h post-transfection, the cells were harvested for RNA extraction and reverse transcription (RT)-quantitative (q)PCR and Western blot. Besides, after 6 h of transfection, the cells were used for Transwell analysis, glucose consumption and lactate production assay.

TNS4 siRNA-3 had the highest inhibition efficiency of TNS4 in HCT116 and RKO cells. Lentivirus pGLVU6/Puro vectors carrying TNS4 shRNA containing the sequence of TNS4 siRNA-3 or siRNA negative control sequence were manufactured using the 2nd generation system by GenePharma Co., Ltd. The lentiviral TNS4 shRNA plasmid (8  $\mu$ g) was co-transfected with psPAX2 (2  $\mu$ g, cat. no. 12260; Addgene, Inc.) and pMD2.G (6  $\mu$ g, cat. no. 12259; Addgene, Inc.) in HEK293T cells (American Type Culture Collection) in 10 cm-culture dish at 37° for 10 h. After replacing the transfection medium using the culture medium, lentiviral particles in the culture medium was collected every day for 3 days. For cell transduction, the HCT116 and RKO cells in the logarithmic growth period were infected with lentiviral particles (multiplicity of infection=20) at 37°C for 12 h. The transfection efficiency was observed under a fluorescence microscope at 24 h after transfection. After 72 h infection, puromycin (1  $\mu$ g/ml, Beyotime Institute of Biotechnology) was added to screen for stable TNS4 knockdown and negative control cell lines. The screening period is about 8 to 12 days. The stable cells were cultured in complete medium with 0.5  $\mu$ g/ml puromycin at 37°C.

**RNA extraction and reverse transcription (RT)-quantitative (q)PCR.** Total RNA extraction from HCT116 and RKO cells was performed using the Cell Total RNA Kit (cat. no. ES-RN001, Shanghai Yishan Biotechnology Co., Ltd.). Subsequently, 1.0  $\mu$ g total RNA was used for cDNA synthesis using the RTIII All-in-One Mix (cat. no. MR05101, Monad Biotech Co., Ltd.) with the following conditions: 37°C for 2 min, 55°C for 15 min, and 85°C for 5 min. RT-qPCR was performed using the CFX96 Touch Real-Time PCR Detection System (Bio-Rad Laboratories, Inc.) using the ChemoHS qPCR Mix (cat. no. MQ00401, Monad Biotech Co., Ltd.) with SYBR Green. The cycling conditions were as follows: one cycle at 95°C for 5 min, 40 cycles of amplification at 95°C for 10 sec, and 60°C for 30 sec. RT-qPCR was performed three times for each sample.  $\beta$ -actin was used for the normalization of gene expression. All primers used for RT-qPCR are listed in Table SII.

**Western blot analysis.** The HCT116 and RKO cells were lysed using RIPA buffer containing protease inhibitors and phosphatase inhibitors (cat. no. P0013D; Beyotime Institute of Biotechnology). The Enhanced BCA Protein Assay kit (cat. no. P0010; Beyotime Institute of Biotechnology) was used for

protein determination. Subsequently, 10% SDS-PAGE; (cat. no. P0012AC; Beyotime Institute of Biotechnology) was used to separate the total protein (30  $\mu$ g), which then was transferred onto PVDF membranes (Cytiva). After blocking with 5% BSA (cat. no. FMS-WB021; Nanjing Femacs Biotechnology Co., Ltd.) for 1.5 h at room temperature, the membranes were incubated with rabbit anti-human TNS4 (1:1,000; cat. no. 11580-1-AP; Proteintech Group, Inc.) or mouse anti-human/mouse GAPDH (1:1,000; cat. no. 60004-1-Ig; Proteintech Group, Inc.) antibodies at 4°C overnight. The membranes were then incubated with the corresponding HRP-conjugated goat anti-rabbit (1:1,000; cat. no. A0208; Beyotime Institute of Biotechnology) or anti-mouse secondary antibodies (1:1,000; cat. no. A0216; Beyotime Institute of Biotechnology) for 1 h at room temperature. Finally, the protein bands were visualized using an ECL reagent (cat. no. 10100; NCM Biotech) in a ChemiDoc™ MP Imaging System (Bio-Rad Laboratories, Inc.). Additionally, ImageJ 2.0 software (National Institutes of Health) was used to analyze the density of the protein bands.

**Cell migration and invasion assay.** To assess CRC cell migration and invasion, a Transwell chamber was purchased from BD Biosciences (cat. no. 353097) and Matrigel from Corning, Inc. (cat. no. 356234). For the cell migration assay,  $3 \times 10^4$  HCT116 or RKO cells in serum-free medium (400  $\mu$ l) were seeded into the upper chamber (8- $\mu$ m pore size). Complete medium containing with 10% FBS was added to the lower chamber. For the invasion assay, Matrigel (200  $\mu$ g/ml) was used to coat the upper chamber at room temperature for 2 h. For migration or invasion, after cultured in the upper chamber for 24 h or 48 h, the cells on the lower surface of the upper chamber were fixed with 4% paraformaldehyde (Beyotime Institute of Biotechnology) at room temperature for 30 min and stained with crystal violet (Beyotime Institute of Biotechnology) at room temperature for 15 min. Images were captured using a Nikon Eclipse/NI-U fluorescence microscope and the number of migrated/invaded cells was counted.

**Glucose consumption and lactate production assay.** A Glucose Assay kit (cat. no. 361510; Shanghai Robio Biotechnology Co., Ltd.) and a Lactate Assay kit (cat. no. A019-2-1; Nanjing Jiancheng Taihao Biotechnology Co., Ltd.) were used for the detection of glucose consumption and lactate production according to the manufacturer's protocols, respectively.

**Statistical analysis.** Data are presented as the mean  $\pm$  standard deviation. GraphPad Prism 6.0 software (Dotmatics) was used for statistical analyses. The paired or unpaired Student's t-test, or one-way ANOVA and Tukey's test's, were used to analyze the data.  $P < 0.05$  was considered to indicate a statistically significant difference.

## Results

**TNS4 is highly expressed in clinical CRC tissue and is associated with the TNM stage of patients with CRC.** To assess the role of TNS4 in CRC, its expression was evaluated in the tissues of patients with CRC. The mRNA levels of TNS4 were significantly upregulated in both colon and rectal cancer tissues compared with normal tissues, according to data from the UALCAN and GEPIA databases (Fig. 1A and B). Furthermore, a CRC tumor cohort was used to assess the protein expression

of TNS4. Compared with the normal control, the tumor tissues exhibited a significantly higher protein expression of TNS4 (Fig. 1C). Moreover, TNS4 expression in patients with late-stage (III-IV) disease was significantly higher than that in patients with early-stage (I-II) disease (Fig. 1D). These data suggest that TNS4 may function as an oncogene in CRC.

**TNS4 is positively correlated with glycolysis-related genes in patients with CRC.** The GEPIA database was used to further evaluate the association between TNS4 and glycolysis in CRC. It was demonstrated that there were significant positive correlations between TNS4 expression and glycolysis-related genes, such as hexokinase 2 (HK2), lactate dehydrogenase (LDH) B, LDHM, pyruvate kinase M1/2 (PKM) and solute carrier (SLC)2A1, in the cancer tissues of patients with CRC (Fig. 2). These data indicate that TNS4 may function as a key regulator of glycolysis in CRC.

**TNS4 knockdown inhibits glucose consumption and lactate production in CRC cells.** To assess the potential role of TNS4 in glycolysis in CRC cells, TNS4 was knocked down in the HCT116 and RKO cell lines by transfecting them with three non-overlapping TNS4 siRNAs (TNS4 siRNA-1, TNS4 siRNA-2 and TNS4 siRNA-3). Compared with NC, transfection with TNS4 siRNA-1, TNS4 siRNA-2 and TNS4 siRNA-3 markedly decreased the protein expression of TNS4 in the HCT116 and RKO cells (Fig. 3A). Moreover, TNS4 knockdown significantly decreased glucose consumption and lactate production in the HCT116 and RKO cells compared with NC (Fig. 3B and C). Compared with NC, TNS4 silencing also led to a significant decrease in the mRNA expression of multiple glycolysis-related genes, including glucose transporter 1, HK2, LDHA and pyruvate dehydrogenase kinase 1 in the CRC cells (Fig. 3D).

**TNS4 silencing attenuates the migration and invasion of CRC cells.** The present study assessed whether TNS4 knockdown regulated the migration and invasion of CRC cells. Compared with the control group, the migration and invasion of the HCT116 and RKO cells was significantly decreased in the TNS4 knockdown groups (Fig. 4A and B). Furthermore, transfection with TNS4 siRNA-2 and TNS4 siRNA-3 significantly downregulated the mRNA expression of matrix metalloproteinase (MMP)-2 and MMP-9 in the HCT116 and RKO cells (Fig. 4C). These data suggest that TNS4 may serve a critical role in the migration and invasion of CRC cells.

**TNS4 silencing inhibits CRC cell migration and invasion via glycolysis.** Previous studies have demonstrated that the activation of glycolysis serves a key role in the migration and invasion of cancer cells (23,24). As TNS4 knockdown significantly suppressed the glycolysis, migration and invasion of CRC cells in the present study, the ability of TNS4 to modulate CRC cell migration and invasion via glycolysis was subsequently evaluated. The results demonstrated that compared with NC, the glucose consumption and lactate production were significantly reduced in HCT116 and RKO cells following TNS4 knockdown, then with the addition of DASA-58, an activator of glycolysis (25), they significantly increased (Fig. 5A-D). Notably, treatment with DASA-58 significantly reversed the

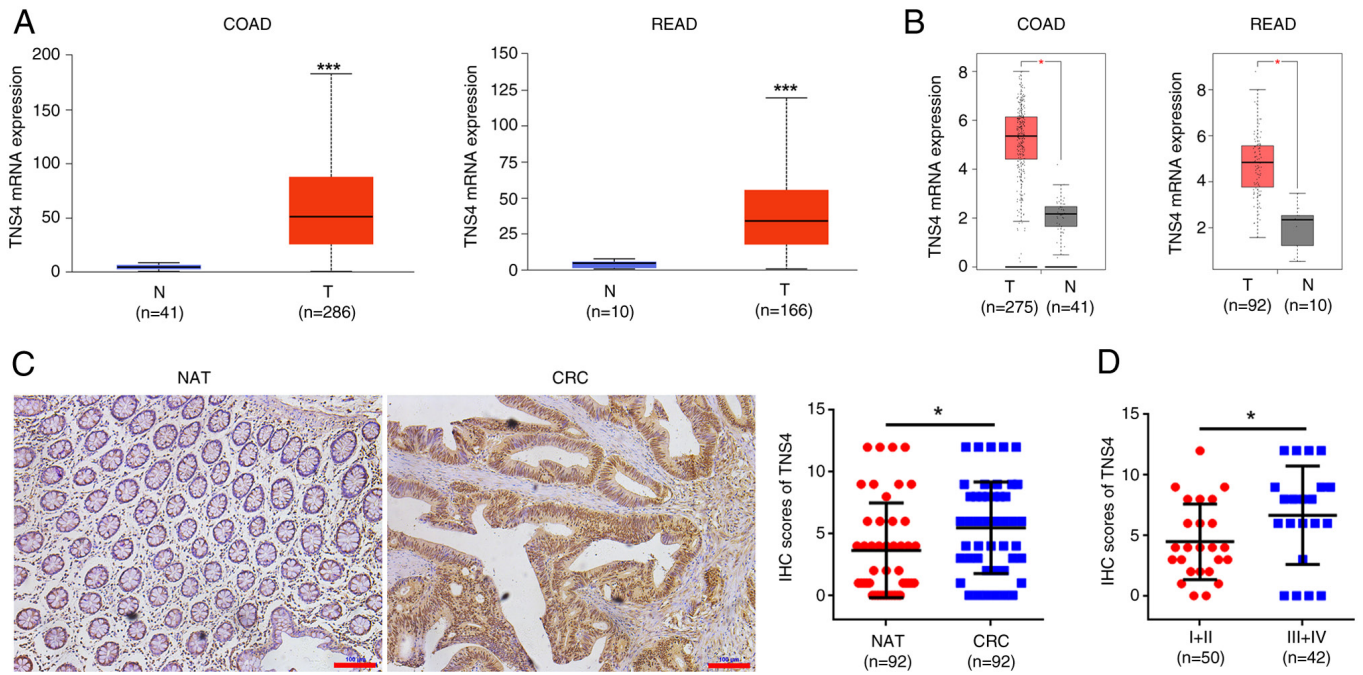


Figure 1. TNS4 is overexpressed in CRC tissue specimens and is associated with tumor-node-metastasis stage. Relative TNS4 mRNA expression in COAD and READ tissue samples from TCGA data based in the (A) University of Alabama at Birmingham Cancer data analysis Portal and (B) Gene Expression Profiling Interactive Analysis databases. T, tumor; N, normal.  $P < 1 \times 10^{-12}$ ;  $P = 1.11 \times 10^{-16}$ . (C) Representative images of IHC of TNS4 in CRC and matched normal tissues from 92 patients with CRC. NAT, nonmalignant adjacent tissues. Scale bar, 100  $\mu$ m. (D) TNS4 protein expression based on the staining index of CRC specimens at different clinical stages.  $P < 0.05$ ;  $***P < 0.001$ . TNS4, Tensin 4; CRC, colorectal cancer; COAD, colon adenocarcinoma; READ, rectum adenocarcinoma; TCGA, The Cancer Genome Atlas; IHC, immunohistochemistry.

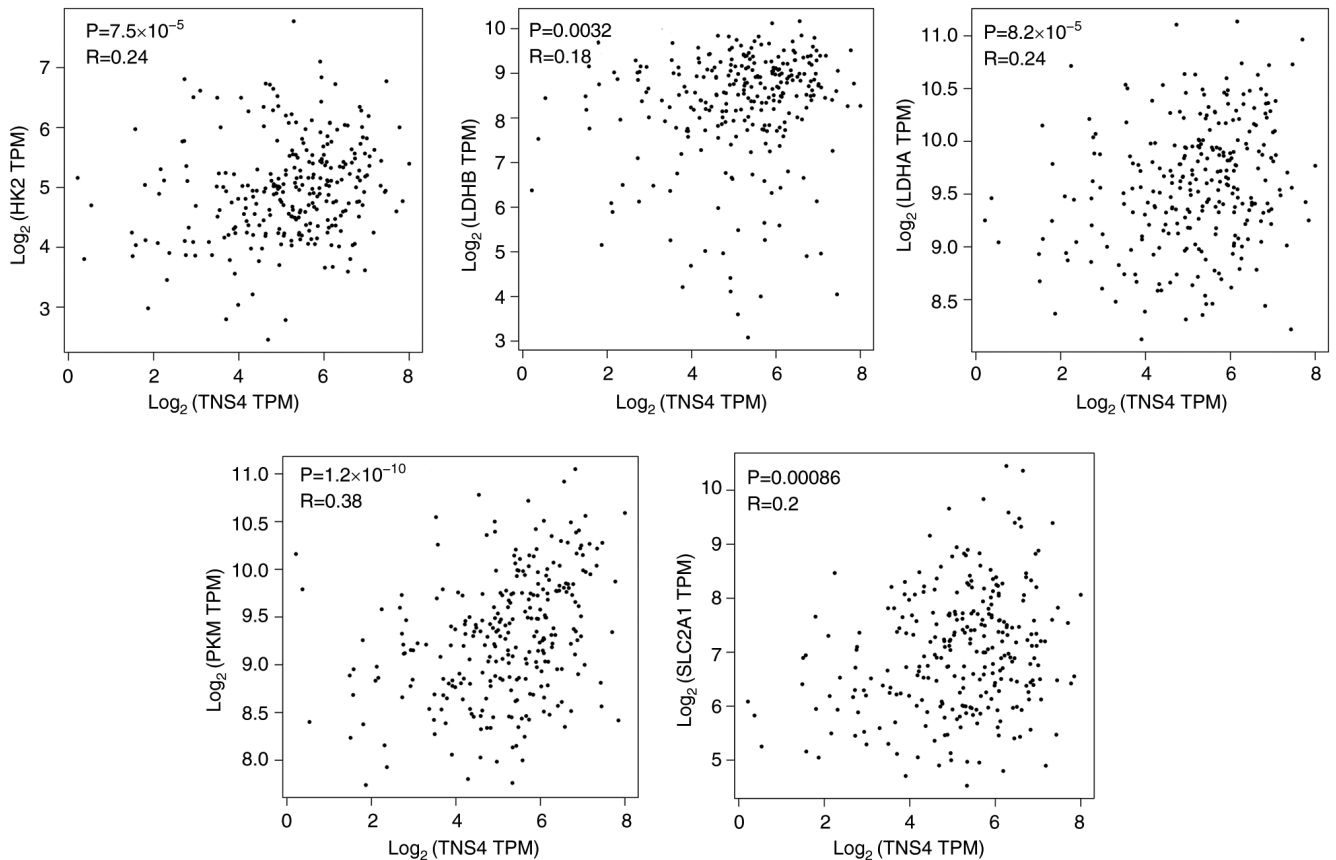


Figure 2. TNS4 is positively correlated with glycolysis-related genes in patients with CRC. The relationship between TNS4 and HK2, LDHB, LDHA, PKM or SLC2A1 in CRC tissue samples from The Cancer Genome Atlas data in the Gene Expression Profiling Interactive Analysis database. TNS4, Tensin 4; CRC, colorectal cancer; HK2, hexokinase 2; LDH, lactate dehydrogenase; PKM, pyruvate kinase M1/2; SLC, solute carrier; TPM, transcripts per million.

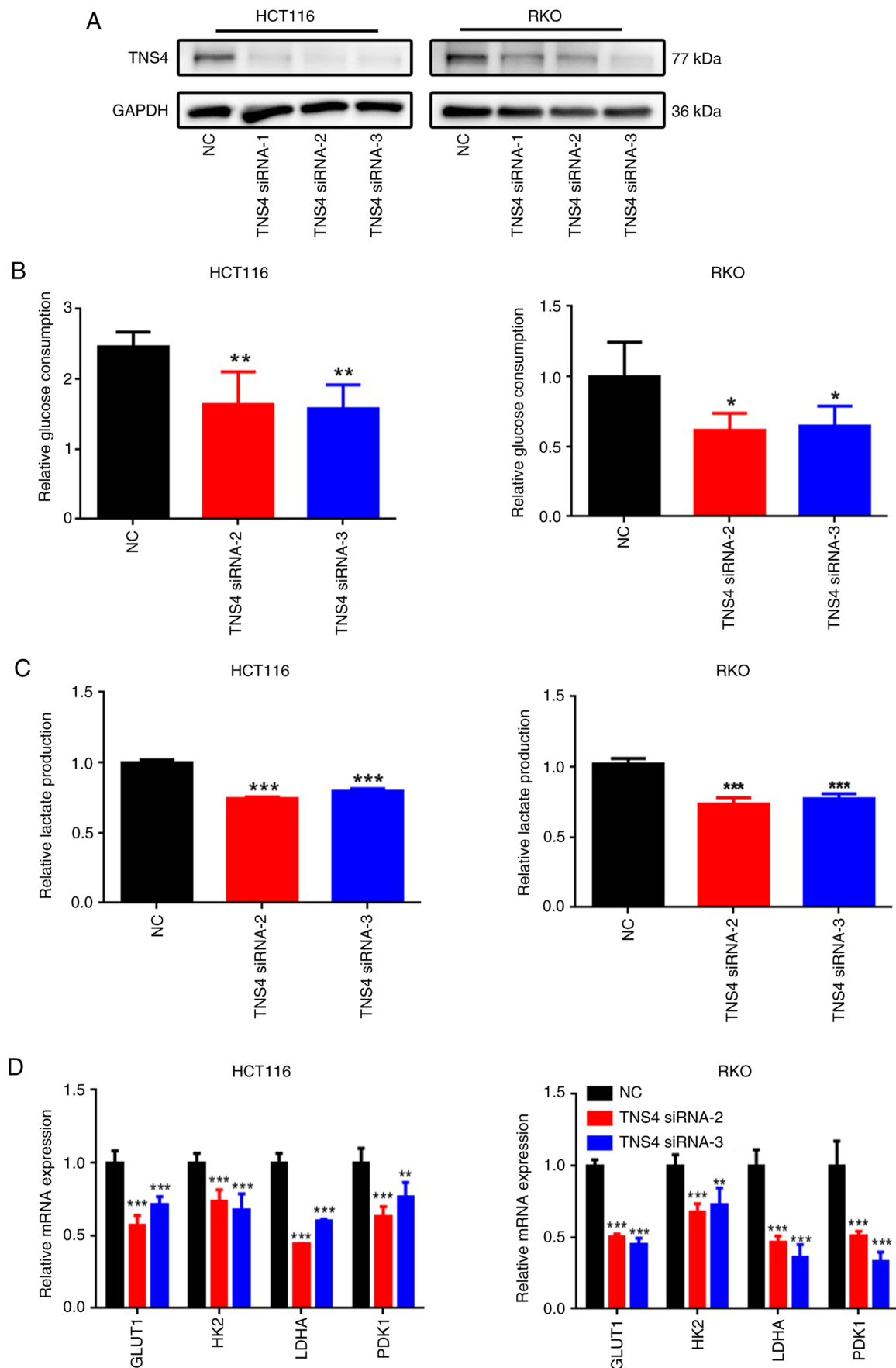


Figure 3. TNS4 depletion inhibits glycolysis in colorectal cancer cells. In both HCT116 and RKO cells transfected with NC, TNS4 siRNA-1, TNS4 siRNA-2 or TNS4 siRNA-3: (A) TNS4 protein levels assessed using western blotting, with GAPDH served as a loading control; (B) glucose consumption and (C) lactate production; and (D) the expression of glycolysis-related genes detected using reverse transcription-quantitative PCR. \* $P < 0.05$ ; \*\* $P < 0.01$ ; \*\*\* $P < 0.001$ . TNS4, Tensin 4; NC, negative control; siRNA, small interfering RNA; GLUT1, glucose transporter 1; HK2, hexokinase 2; LDH, lactate dehydrogenase; PDK1, pyruvate dehydrogenase kinase 1.



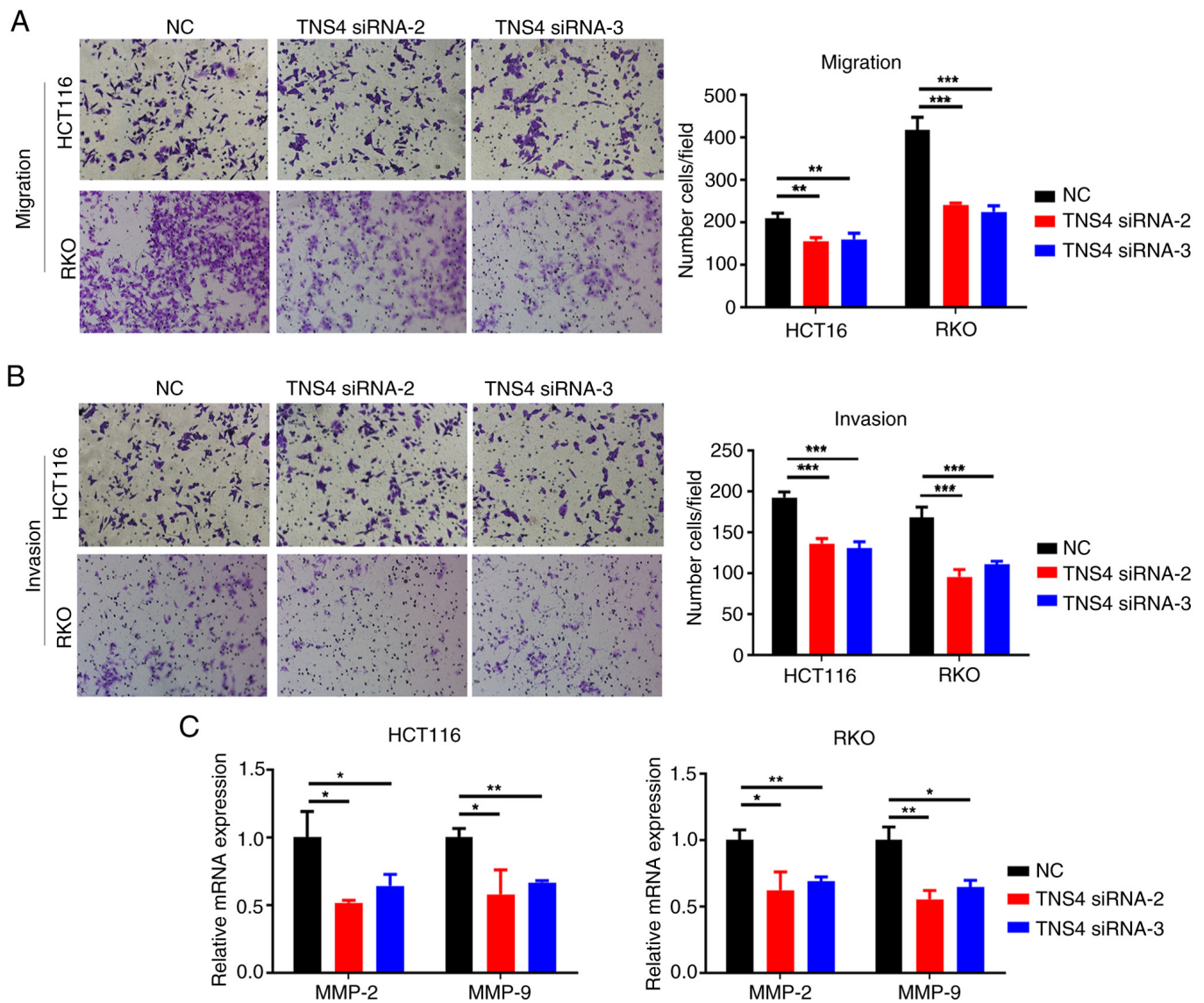


Figure 4. TNS4 depletion suppressed migration and invasion in colorectal cancer cells. (A) Migration and (B) invasion assessed using Transwell assays (magnification 200x), and (C) the expression of MMP-2 and MMP-9 evaluated using reverse-transcription-quantitative PCR in HCT116 or RKO cells transfected with NC, TNS4 siRNA-2 or TNS4 siRNA-3. \*P<0.05; \*\*P<0.01; \*\*\*P<0.001. TNS4, Tensin 4; NC, negative control; siRNA, small interfering RNA; MMP, matrix metalloproteinase.

effects of TNS4 silencing on the migration and invasion of HCT116 and RKO cells (Fig. 5E-G).

*TNS4 knockdown inhibits glycolysis and metastasis of CRC cells via the  $\beta$ -catenin/c-Myc pathway.* To evaluate the mechanism of action of TNS4 in the glycolysis and metastasis of CRC cells, a review of the literature was performed.  $\beta$ -catenin/c-Myc signaling has been reported to be involved in regulating both glycolysis and metastasis in cancers (16,26-30). In addition, TNS4 overexpression has been reported to activate  $\beta$ -catenin signaling in colon cancer (16). Therefore, it was hypothesized that TNS4 knockdown suppressed glycolysis, migration and invasion of CRC cells via the  $\beta$ -catenin/c-Myc pathway. The results of western blot analysis revealed that compared with NC, the knockdown of TNS4 markedly reduced the expression of  $\beta$ -catenin and c-Myc in the HCT116 and RKO cells (Fig. 6A). Furthermore, the addition of SKL2001, an

activator of  $\beta$ -catenin signaling (31), notably reversed the inhibitory effects of TNS4 knockdown on  $\beta$ -catenin and c-Myc expression in the HCT116 and RKO cells (Fig. 6A). In addition, it was demonstrated that compared with NC, the glucose consumption and lactate production were significantly reduced in HCT116 and RKO cells following TNS4 knockdown, then with the addition of SKL2001, they significantly increased (Fig. 6B-E). Additionally, treatment with SKL2001 significantly increased the migration and invasion of the HCT116 and RKO cells following transfection with TNS4 siRNA-2 (Fig. 6F). These results suggest that TNS4 may promote the glycolysis and metastasis of CRC cells via the  $\beta$ -catenin/c-Myc pathway.

## Discussion

There is increasing evidence to indicate that TNS4 is highly expressed in multiple cancer tissues (4). Albasri *et al* (8)

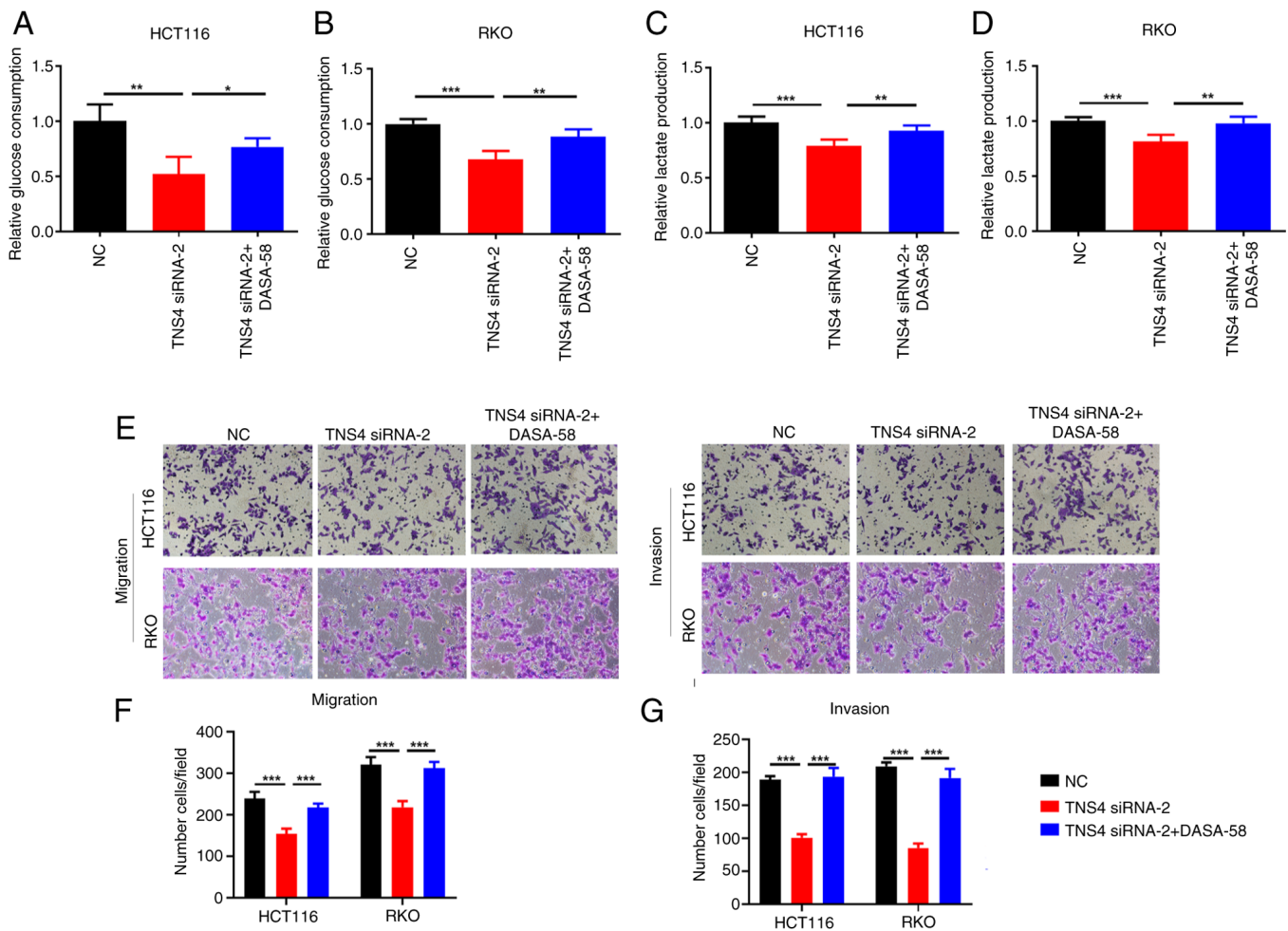


Figure 5. TNS4 promotes migration and invasion in colorectal cancer cells via glycolysis. Glucose consumption assessed in (A) HCT116 and (B) RKO cells co-treated with TNS4 siRNA-2 and DASA-58. Lactate production evaluated in (C) HCT116 and (D) RKO cells co-treated with TNS4 siRNA-2 and DASA-58 assessed using Transwell assays (magnification 200x). (E) Migration and invasion of HCT116 and RKO cells co-treated with TNS4 siRNA-2 and DASA-58 assessed using Transwell assays (magnification 200x). (F) The number of migration cells in (E). (G) The number of invasion cells in (E). \* $P < 0.05$ ; \*\* $P < 0.01$ ; \*\*\* $P < 0.001$ . TNS4, Tensin 4; siRNA, small interfering RNA; NC, negative control.

analyzed the expression of TNS4 in a series of 462 patients with CRC using immunohistochemistry and reported that the expression of TNS4 was markedly increased, and was associated with an advanced Dukes' stage, poor prognosis and distant metastasis. In the present study, it was demonstrated that TNS4 mRNA levels were significantly upregulated in CRC tissues according to data from the UALCAN and GEPIA databases. Furthermore, TNS4 protein expression was significantly increased in the tissue samples of patients with CRC and was positively associated with the TNM stage. These results suggest that TNS4 functions as an oncogene and serves critical roles in the progression of CRC. In addition, TNS4 is an important protein involved in maintaining normal cellular functions, particularly those related to cell adhesion, migration and signaling (4,32). However, as the current study focused on the biological functions of TNS4 in CRC, the role of TNS4 in normal cells by knocking down its expression was not assessed. It is important to explore the roles of TNS4 in maintaining the normal structure and function of the kidney and heart, as well as wound regeneration (32).

Previous studies have demonstrated that TNS4 serves a crucial role in the progression of cancer by regulating several

biological functions, such as EMT, cell motility, apoptosis and tumorigenicity (4,9). Based on data from TCGA, the present study revealed that the expression of TNS4 was significantly associated with glycolysis-related genes (namely, HK2, LDHB, LDHM, PKM and SLC2A1) in the CRC tissue samples, suggesting that TNS4 may be involved in the regulation of aerobic glycolysis in CRC. Aerobic glycolysis, as a key hallmark of cancer, has been demonstrated to be associated with tumor growth, metastasis and drug resistance in CRC (33,34). Shi *et al* (21) reported that B7 homolog 3, an immunoregulatory protein, enhanced the chemoresistance of CRC cells by promoting HK2-mediated aerobic glycolysis. Zhao *et al* (35) reported that lncRNA MIR17HG overexpression resulted in persistent glycolysis and the invasion and liver metastasis of CRC cells. To the best of our knowledge, the present study is the first to demonstrate that TNS4 knockdown significantly inhibits glucose consumption, lactate production and the mRNA expression of multiple glycolysis-related genes in CRC cells. These results suggest that TNS4 is a key regulator of aerobic glycolysis in CRC.

A number of studies have indicated that TNS4 is associated with the metastasis of cancer cells (8,11,36,37). A previous

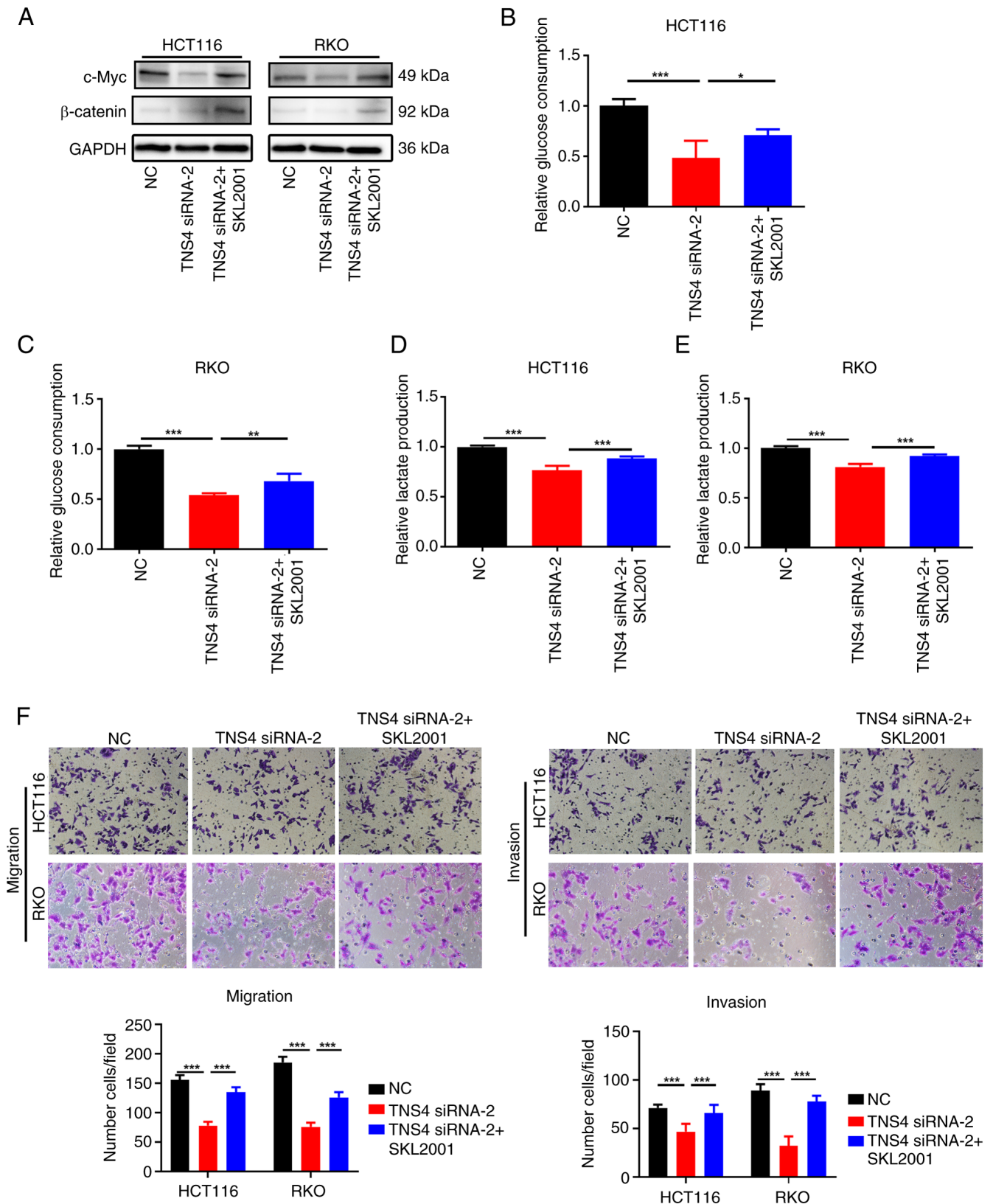


Figure 6. TNS4 promotes glycolysis, migration and invasion in CRC cells through the  $\beta$ -catenin/c-Myc pathway. (A)  $\beta$ -catenin and c-Myc protein levels assessed using western blotting in both HCT116 and RKO cells co-treated with TNS4 siRNA-2 and SKL2001, with GAPDH as the loading control. Glucose consumption assessed in (B) HCT116 and (C) RKO cells, and lactate production evaluated in (D) HCT116 or (E) RKO cells co-treated with TNS4 siRNA-2 and SKL2001. (F) Migration and invasion of HCT116 and RKO cells co-treated with TNS4 siRNA-2 and SKL2001 assessed using Transwell assays (magnification  $\times 200$ ). \* $P < 0.05$ ; \*\* $P < 0.01$ ; \*\*\* $P < 0.001$ . TNS4, Tensin 4; siRNA, small interfering RNA; NC, negative control.

study reported that, in non-small cell lung cancer, TNS4 overexpression induced the EMT process and elevated the

migratory capacity of the cells (12). Asiri *et al* (36) noted that TNS4, as a direct mediator of TGF- $\beta$ 1 signaling, participated



in TGF- $\beta$ 1-induced EMT and cell motility in CRC. Moreover, the TNS4-integrin-linked kinase interaction also modulated cell motility in CRC (8). In the present study, it was demonstrated that TNS4 knockdown significantly decreased the migration and invasion of CRC cells. These results suggest that TNS4 exerts promoting effects on the metastasis of CRC cells. Furthermore, aerobic glycolysis is a crucial influencing factor for cancer metastasis (38,39), and in the present study, it was demonstrated that treatment with DASA-58, an activator of glycolysis, significantly reversed the effects of TNS4 silencing on the migration and invasion of CRC cells. These results suggest that the TNS4-induced promotion of the migration and invasion of CRC cells is glycolysis-dependent.

The activation of the  $\beta$ -catenin/c-Myc pathway has been reported to be associated with cancer (29,30). For example, CD36 has been reported to inhibit the glycolysis and tumorigenesis of CRC through the glypican 4- $\beta$ -catenin-c-Myc signaling axis (40). Liang *et al* (41) indicated that butyrate controlled gastric cancer cell proliferation, migration, invasion and aerobic glycolysis by downregulating Wnt/ $\beta$ -catenin/c-MYC signaling. In human hepatocellular carcinoma, Gankyrin has been reported to upregulate glycolysis and glutaminolysis to promote tumorigenesis, metastasis and drug resistance via upregulating c-Myc through the activation of  $\beta$ -catenin signaling (26). In the present study, it was revealed that TNS4 knockdown significantly reduced the protein expression of  $\beta$ -catenin and c-Myc in CRC cells. Consistent with these findings, it has been reported that TNS4 can activate the  $\beta$ -catenin pathway to regulate tumorigenicity and tumor angiogenesis (5,16,42). Moreover, SKL2001, an activator of  $\beta$ -catenin signaling, significantly reversed the effects of TNS4 knockdown on glucose consumption, lactate production, and the migration and invasion of CRC cells. Taken together, these results suggest that TNS4 promotes the glycolysis and metastasis of CRC cells through the  $\beta$ -catenin/c-Myc pathway.

In conclusion, the results of the present study demonstrated that TNS4 was highly expressed in CRC tissues, and that its high expression was positively associated with the TNM stages of patients with CRC. Moreover, the silencing of TNS4 inhibited the glycolysis, migration and invasion of CRC cells via the  $\beta$ -catenin/c-Myc pathway. This suggests that TNS4 may be an efficient target for CRC diagnosis and therapy.

## Acknowledgements

Not applicable.

## Funding

No funding was received.

## Availability of data and materials

The data generated in the present study may be requested from the corresponding author.

## Authors' contributions

YW and CX designed the experiments. YW and YL performed the experiments and analyzed the data. YW wrote

the manuscript. CX and YL made suggestions during the writing. YW and YL confirm the authenticity of all the raw data. All authors have read and approved the final manuscript.

## Ethics approval and consent to participate

The present study was approved by the Ethics Review Board of the First Affiliated Hospital of Soochow University (Suzhou, China; approval no. 2021-327). Written informed consent was obtained from the patients prior to obtaining the samples.

## Patient consent for publication

Not applicable.

## Competing interests

The authors declare that they have no competing interests.

## References

1. Biller LH and Schrag D: Diagnosis and treatment of metastatic colorectal cancer: A review. *JAMA* 325: 669-685, 2021.
2. Piawah S and Venook AP: Targeted therapy for colorectal cancer metastases: A review of current methods of molecularly targeted therapy and the use of tumor biomarkers in the treatment of metastatic colorectal cancer. *Cancer* 125: 4139-4147, 2019.
3. Holladay L, Luu J, Balendra V and Kmetz K: Current and potential treatment of colorectal cancer metastasis to bone. *Cancer Treat Res Commun* 37: 100763, 2023.
4. Lo SH: C-terminal tensin-like (CTEN): A promising biomarker and target for cancer. *Int J Biochem Cell Biol* 51: 150-154, 2014.
5. Lu X, Zhou B, Cao M, Shao Q, Pan Y and Zhao T: CTEN inhibits tumor angiogenesis and growth by targeting VEGFA through down-regulation of  $\beta$ -catenin in breast cancer. *Technol Cancer Res Treat* 20: 15330338211045506, 2021.
6. Al-Ghamdi S, Cachat J, Albasri A, Ahmed M, Jackson D, Zaitoun A, Guppy N, Otto WR, Alison MR, Kindle KB and Ilyas M: C-terminal tensin-like gene functions as an oncogene and promotes cell motility in pancreatic cancer. *Pancreas* 42: 135-140, 2013.
7. Sasaki H, Moriyama S, Mizuno K, Yukiue H, Konishi A, Yano M, Kaji M, Fukai I, Kiriyaama M, Yamakawa Y and Fujii Y: Cten mRNA expression was correlated with tumor progression in lung cancers. *Lung Cancer* 40: 151-155, 2003.
8. Albasri A, Al-Ghamdi S, Fadhil W, Aleskandarany M, Liao YC, Jackson D, Lobo DN, Lo SH, Kumari R, Durrant L, *et al*: Cten signals through integrin-linked kinase (ILK) and may promote metastasis in colorectal cancer. *Oncogene* 30: 2997-3002, 2011.
9. Asiri A, Toss MS, Raposo TP, Akhlaq M, Thorpe H, Alfahed A, Asiri A and Ilyas M: Cten promotes epithelial-mesenchymal transition (EMT) in colorectal cancer through stabilisation of Src. *Pathol Int* 69: 381-391, 2019.
10. Thorpe H, Asiri A, Akhlaq M and Ilyas M: Cten promotes epithelial-mesenchymal transition through the post-transcriptional stabilization of Snail. *Mol Carcinog* 56: 2601-2609, 2017.
11. Lu X, Gao J, Zhang Y, Zhao T, Cai H and Zhang T: CTEN induces epithelial-mesenchymal transition (EMT) and metastasis in non small cell lung cancer cells. *PLoS One* 13: e0198823, 2018.
12. Albasri A, Seth R, Jackson D, Benhasouna A, Crook S, Nateri AS, Chapman R and Ilyas M: C-terminal Tensin-like (CTEN) is an oncogene which alters cell motility possibly through repression of E-cadherin in colorectal cancer. *J Pathol* 218: 57-65, 2019.
13. Wang YX, Huang CY, Chiu HJ, Huang PH, Chien HT, Jwo SH and Liao YC: Nuclear-localized CTEN is a novel transcriptional regulator and promotes cancer cell migration through its downstream target CDC27. *J Physiol Biochem* 79: 163-174, 2023.
14. Lu X, Zhang Y, Pan Y, Cao M, Zhou X and Zhang T: Overexpression of CTEN is associated with gefitinib resistance in non-small cell lung cancer. *Oncol Lett* 21: 40, 2021.
15. Li Y, Mizokami A, Izumi K, Narimoto K, Shima T, Zhang J, Dai J, Keller ET and Namiki M: CTEN/tensin 4 expression induces sensitivity to paclitaxel in prostate cancer. *Prostate* 70: 48-60, 2010.

16. Liao YC, Chen NT, Shih YP, Dong Y and Lo SH: Up-regulation of C-terminal tensin-like molecule promotes the tumorigenicity of colon cancer through beta-catenin. *Cancer Res* 69: 4563-4566, 2009.
17. Schwartz L, Supuran CT and Alfarouk KO: The Warburg effect and the hallmarks of cancer. *Anticancer Agents Med Chem* 17: 164-170, 2017.
18. Zhong X, He X, Wang Y, Hu Z, Huang H, Zhao S, Wei P and Li D: Warburg effect in colorectal cancer: The emerging roles in tumor microenvironment and therapeutic implications. *J Hematol Oncol* 15: 160, 2022.
19. Liu J, Liu ZX, Wu QN, Lu YX, Wong CW, Miao L, Wang Y, Wang Z, Jin Y, He MM, *et al*: Long noncoding RNA AGPG regulates PFKFB3-mediated tumor glycolytic reprogramming. *Nat Commun* 11: 1507, 2020.
20. Yu S, Zang W, Qiu Y, Liao L and Zheng X: Deubiquitinase OTUB2 exacerbates the progression of colorectal cancer by promoting PKM2 activity and glycolysis. *Oncogene* 41: 46-56, 2022.
21. Shi T, Ma Y, Cao L, Zhan S, Xu Y, Fu F, Liu C, Zhang G, Wang Z, Wang R, *et al*: B7-H3 promotes aerobic glycolysis and chemoresistance in colorectal cancer cells by regulating HK2. *Cell Death Dis* 10: 308, 2019.
22. Liu M, Huang F, Zhang D, Ju J, Wu XB, Wang Y, Wang Y, Wu Y, Nie M, Li Z, *et al*: Heterochromatin protein HP1gamma promotes colorectal cancer progression and is regulated by miR-30a. *Cancer Res* 75: 4593-4604, 2015.
23. Chen Z, Hu Z, Sui Q, Huang Y, Zhao M, Li M, Liang J, Lu T, Zhan C, Lin Z, *et al*: LncRNA FAM83A-AS1 facilitates tumor proliferation and the migration via the HIF-1 $\alpha$ /glycolysis axis in lung adenocarcinoma. *Int J Biol Sci* 18: 522-535, 2022.
24. Lin S, Li Y, Wang D, Huang C, Marino D, Bollt O, Wu C, Taylor MD, Li W, DeNicola GM, *et al*: Fascin promotes lung cancer growth and metastasis by enhancing glycolysis and PFKFB3 expression. *Cancer Lett* 518: 230-242, 2021.
25. Rao J, Wang H, Ni M, Wang Z, Wang Z, Wei S, Liu M, Wang P, Qiu J, Zhang L, *et al*: FSTL1 promotes liver fibrosis by reprogramming macrophage function through modulating the intracellular function of PKM2. *Gut* 71: 2539-2550, 2022.
26. Liu R, Li Y, Tian L, Shi H, Wang J, Liang Y, Sun B, Wang S, Zhou M, Wu L, *et al*: Gankyrin drives metabolic reprogramming to promote tumorigenesis, metastasis and drug resistance through activating  $\beta$ -catenin/c-Myc signaling in human hepatocellular carcinoma. *Cancer Lett* 443: 34-46, 2019.
27. Dai M, Song J, Wang L, Zhou K and Shu L: HOXC13 promotes cervical cancer proliferation, invasion and Warburg effect through  $\beta$ -catenin/c-Myc signaling pathway. *J Bioenerg Biomembr* 53: 597-608, 2021.
28. Liu Y, Huang Y, Zhang J, Pei C, Hu J, Lyu J and Shen Y: TIMMDC1 knockdown inhibits growth and metastasis of gastric cancer cells through metabolic inhibition and AKT/GSK3 $\beta$ / $\beta$ -catenin signaling pathway. *Int J Biol Sci* 14: 1256-1267, 2018.
29. Rennoll S and Yochum G: Regulation of MYC gene expression by aberrant Wnt/ $\beta$ -catenin signaling in colorectal cancer. *World J Biol Chem* 6: 290-300, 2015.
30. Ashihara E, Takada T and Maekawa T: Targeting the canonical Wnt/ $\beta$ -catenin pathway in hematological malignancies. *Cancer Sci* 106: 665-671, 2015.
31. Wu YB, Li SY, Liu JY, Xue JJ, Xu JF, Chen T, Cao TY, Zhou H, Wu TT, Dong CL, *et al*: Long non-coding RNA NRSN2-AS1 promotes ovarian cancer progression through targeting PTK2/ $\beta$ -catenin pathway. *Cell Death Dis* 14: 696, 2023.
32. Liao YC and Lo SH: Tensins-emerging insights into their domain functions, biological roles and disease relevance. *J Cell Sci* 134: jcs254029, 2021.
33. Zafari N, Velayati M, Damavandi S, Pourali G, Mobarhan MG, Nassiri M, Hassanian SM, Khazaei M, Ferns GA and Avan A: Metabolic pathways regulating colorectal cancer: A potential therapeutic approach. *Curr Pharm Des* 28: 2995-3009, 2022.
34. Nenkov M, Ma Y, Gaßler N and Chen Y: Metabolic reprogramming of colorectal cancer cells and the microenvironment: Implication for therapy. *Int J Mol Sci* 22: 6262, 2021.
35. Zhao S, Guan B, Mi Y, Shi D, Wei P, Gu Y, Cai S, Xu Y, Li X, Yan D, *et al*: LncRNA MIR17HG promotes colorectal cancer liver metastasis by mediating a glycolysis-associated positive feedback circuit. *Oncogene* 40: 4709-4724, 2021.
36. Asiri A, Raposo TP, Alfahed A and Ilyas M: TGF $\beta$ 1-induced cell motility but not cell proliferation is mediated through Cten in colorectal cancer. *Int J Exp Pathol* 99: 323-330, 2018.
37. Fleming JC, Woo J, Moutasim K, Hanley CJ, Frampton SJ, Wood O, Ward M, Woelk CH, Ottensmeier CH, Hafizi S, *et al*: CTEN induces tumour cell invasion and survival and is prognostic in radiotherapy-treated head and neck cancer. *Cancers (Basel)* 12: 2963, 2020.
38. Yang J, Ren B, Yang G, Wang H, Chen G, You L, Zhang T and Zhao Y: The enhancement of glycolysis regulates pancreatic cancer metastasis. *Cell Mol Life Sci* 77: 305-321, 2020.
39. Feng J, Li J, Wu L, Yu Q, Ji J, Wu J, Dai W and Guo C: Emerging roles and the regulation of aerobic glycolysis in hepatocellular carcinoma. *J Exp Clin Cancer Res* 39: 126, 2020.
40. Fang Y, Shen ZY, Zhan YZ, Feng XC, Chen KL, Li YS, Deng HJ, Pan SM, Wu DH and Ding Y: CD36 inhibits  $\beta$ -catenin/c-myc-mediated glycolysis through ubiquitination of GPC4 to repress colorectal tumorigenesis. *Nat Commun* 10: 3981, 2019.
41. Liang Y, Rao Z, Du D, Wang Y and Fang T: Butyrate prevents the migration and invasion, and aerobic glycolysis in gastric cancer via inhibiting Wnt/ $\beta$ -catenin/c-Myc signaling. *Drug Dev Res* 84: 532-541, 2023.
42. Raposo TP, Alfahed A, Nateri AS and Ilyas M: Tensin4 (TNS4) is upregulated by Wnt signalling in adenomas in multiple intestinal neoplasia (Min) mice. *Int J Exp Pathol* 101: 80-86, 2020.



Copyright © 2024 Wang et al. This work is licensed under a Creative Commons Attribution-NonCommercial-NoDerivatives 4.0 International (CC BY-NC-ND 4.0) License.