Quantitative proteomic profiling of tumor-associated vascular endothelial cells in colorectal cancer

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ABSTRACT
To investigate the global proteomic profiles of vascular endothelial cells (VECs) in the tumor microenvironment and antiangiogenic therapy for colorectal cancer (CRC), matched pairs of normal (NVECs) and tumor-associated VECs (TVECs) were purified from CRC tissues by laser capture microdissection and subjected to iTRAQ-based quantitative proteomics analysis. Here, 216 differentially expressed proteins (DEPs) were identified and used for bioinformatics analysis. Interestingly, these proteins were implicated in epithelial mesenchymal transition (EMT), ECM-receptor interaction, focal adhesion, PI3K-Akt signaling pathway, angiogenesis and HIF-1 signaling pathway, which may play important roles in CRC angiogenesis. Among these DEPs we found that Tenascin-C (TNC) was upregulated in TVECs of CRC and correlated with CRC multistage carcinogenesis and metastasis. Furthermore, the reduction of tumor-derived TNC could attenuate human umbilical vein endothelial cells (HUVECs) proliferation, migration and tube formation through ITGB3/FAK/Akt signaling pathway. Based on the present work, we provided a large-scale proteomic profiling of VECs in CRC with quantitative information, a certain number of potential antiangiogenic targets and a novel vision in the angiogenesis bio-mechanism of CRC.

KEY WORDS: Colorectal cancer, Quantitative proteomics, Angiogenesis, Tenascin-C

INTRODUCTION
Tumor angiogenesis plays a vital role in creating the tumor microenvironment and is necessary for tumor growth and metastasis. Currently, angiogenesis inhibitors have become important drugs in the treatment of solid tumors, including colorectal cancer (CRC) (Lopez et al., 2019; Riechmann and Grothey, 2017). Vascular endothelial cells (VECs), which interact with tumor cells, extracellular matrix and immune killer cells, form the major components of tumor microenvironment. Recently antiangiogenic strategies have largely focused on targeting VECs (Liu et al., 2016, 2015). Tumor-associated VECs (TVECs) undergo phenotypic and epigenetic changes during tumor initiation and progression (Xiong et al., 2009). Increasing evidence indicates that proteins are primary targets of therapeutic drugs, and that proteins located in TVECs are potential therapeutic targets against tumor angiogenesis and widely used in screening antiangiogenic drugs (Kalèn et al., 2009; Sonveaux, 2008).

Proteomics has introduced an effective cancer research method that provides new opportunities for discovering the therapeutic targets of CRC and for revealing the underlying molecular mechanism of this disease. The proteomics of CRC have been extensively studied (Álvarez-Chaver et al., 2018; Peng et al., 2016), but relatively few studies have focused on tumor microenvironment, especially the proteome of VECs in CRC. However, the VEC comprises only a relatively small percentage of the total tumor volume. Consequently, an altered VEC expression signature may be easily masked in whole-tumor studies. To overcome this limitation, laser capture microdissection (LCM) was exploited to isolate a relatively pure VEC from heterogeneous frozen tumor tissue samples in this study.

Tenascin-C (TNC), a large extracellular matrix protein, is highly expressed in pathological contexts, especially in cancer (Midwood et al., 2011). The high expression of TNC correlates with poor prognosis in several cancer types, including breast cancer, glioblastoma and colorectal cancer (Midwood et al., 2016). It has been clearly demonstrated that TNC promotes multiple events in cancer progression, such as tumor cell survival, proliferation, invasion and lung metastasis (Sun et al., 2018). Moreover, high TNC levels are also correlated with angiogenic switch, higher tumor vessel density and vessel leakiness (Midwood et al., 2016), suggesting that TNC plays multiple roles in angiogenesis.

We compared the protein expression level between CRC tumor VECs and matched adjacent nonmalignant colorectal (ANC) tissue VECs from the same patient. This approach has the advantage of eliminating some of the inherent heterogeneity between individual patients and between different cell types presented in the samples (Johann et al., 2010; Unwin et al., 2003). The resulting peptides of the enzymatically digested proteins were measured by iTRAQ-based quantitative proteomics. Compared to conventional proteomic technology, this approach possesses many advantages, such as high throughput, high accuracy, high repeatability and high sensitivity, and it can be used for various types of biological samples (Pierce et al., 2008). A total of 216 differentially expressed proteins (DEPs) were identified. TNC was further validated by immunohistochemical analysis, and the effects of TNC on human umbilical vein endothelial cells (HUVECs) proliferation, migration and tube formation were determined. Based on our data, we presented the proteome of VECs in CRC using LCM technology and quantitative proteomics analysis. These TVECs-related proteins...
may serve as potential therapeutic targets and increase our understanding of the of CRC angiogenesis mechanisms.

RESULTS
Proteomic profiling of VECs in CRC
We compared the global proteomic profiles of paired TVECs and NVECs in ten patients with CRC to identify proteins and pathways that may be associated with the CRC angiogenesis. As a result, A total of 2058 non-redundant proteins were repeatedly identified and quantified at a minimum confidence level of 95% (unused ProtScore >1.3) by triplicate iTRAQ labeling and 2D LC-MS/MS analyses (Table S1). A protein density plot was subsequently generated to determine the thresholds for clustering DEPs using the ratios of those quantified proteins (Chakraborty et al., 2015). Using 10%, 90% and in-between quantile-based thresholds, averaged ratio-fold changes >1.2129 or <0.7963 between two cohorts of proteins were categorized as upregulated and downregulated proteins, respectively (Fig. 1A).

A total of 216 proteins were found to be differentially expressed, including 119 upregulated proteins and 97 downregulated proteins in TVECs relative to NVECs (Table S2). Compared to previous literature, 45 of the top 100 DEPs (top 50 up- and top 50 downregulated proteins) have been proved as angiogenesis-related proteins, and 17 of the 45 proteins have been reported or predicted as potential targets for cancer antiangiogenic treatment. These proteins are presented in a heatmap format in Fig. 1D. In addition, the top 50 upregulated proteins that are either secreted or localized in the membrane are highlighted in gray in the heatmap as potential therapeutic targets in TVECs (gene names of the respective proteins are: FBLN5, PRG2, YBX1, CEACAM6, S100A9, THBS1, PCBP2, CANX, P4HB, RPL35A, S100A8, MANF, LRRCS9, FTL, TOMM22, ATP6V1D, TNC, BASP1 and RNASE3) (Fig. 1D). To some degree, these results indicate that our findings were consistent with previous studies and also make new discoveries.

Bioinformatics analysis
To obtain a biological view of the identified proteins, a total of 2058 proteins were classified according to cellular compartment levels using the PANTHER GO classification system. Variable cellular compartments cell part (38.8%), organelle (26.4%) and membrane (5.1%) are shown in (Fig. 1B). Metascape enrichment analysis, accounting for all the DEPs, demonstrated that processes related EMT, ECM-receptor interaction, focal adhesion, PI3K-Akt signaling pathway, angiogenesis and HIF-1 signaling pathway were over-represented (Fig. 1C, Table S3). The top 50 upregulated and downregulated proteins mapping to each Metascape enrichment term group are presented in a heatmap format in Fig. 1D. The subcellular localization of these proteins is also presented in the heatmap. Moreover, Metascape enrichment analysis mapped all the identified proteins to 66 signaling pathways (Table S4). Interestingly, the results demonstrated that TVECs exhibited an increased dependence on processes related to focal adhesion, PI3K-Akt signaling pathway, HIF-1 signaling pathway and EMT (Fig. 2A,B; Fig. S1, Table S5), which might play an important role in CRC angiogenesis.

Validation of the expression of TNC in VECs using immunohistochemistry
In our study, we found TNC was involved in focal adhesion, PI3K/Akt signaling pathway and EMT. To confirm the expression and location of TNC in CRC tissues, we detected the expression of TNC using immunohistochemistry in 30 cases of non-neoplastic colonic mucosa (NCM), 30 cases of adenomatous colorectal polyps (AD), 30 cases of colorectal carcinoma in situ (CIS) and 50 cases of invasive colorectal carcinoma (ICC). As shown in Fig. 3 and Table 1, the expression levels of TNC were progressively increased during the CRC carcinogenic process from early stage, AD, to late stage, ICC. (P<0.05). Strong TNC immunostaining was readily detected in the stoma and VECs of the CIS and ICC (Fig. 3A,B), whereas there was weak staining in AD and negative staining in NCM (Fig. 3C,D). In addition, the expression levels of TNC in VECs of lymph nodes with metastasis were stronger compared to those in lymph nodes without metastasis (Fig. 3E,F). Immunostaining of NCM and ICC for the EC marker CD34 was used as a positive control for VECs (Fig. 3G,H).

Furthermore, we examined the relationship between the expression levels of TNC and clinico-pathological characteristics in the 80 cases of CRC tissues above (30 cases of CIS and 50 cases of ICC). The results showed that TNC expression levels in VECs of CRC were closely correlated with lymph nodes metastasis (P<0.05) and distant metastasis (P<0.01) but did not correlate with age or gender (P>0.05; Table 2).

Reduction of tumor-derived TNC inactivates ITGB3/FAK/Akt signaling
Tumor-derived TNC promotes or enhances the process of angiogenesis in different tumor models (Hirata et al., 2009; Kawamura et al., 2018; Pezzolo et al., 2011; Rupp et al., 2016), but the mechanistic insight has not been fully elucidated. In our previous study, the TNC level is much higher in high metastatic potential CRC cell line SW620 compared to the other four CRC cell lines, particularly in the conditioned medium (Li et al., 2016). In the current study, we generated CRC cell line SW620 with knockdown of TNC (Fig. 4A,B), then collected conditioned media from SW620, SW620/Vector and SW620/shTNC cells and subsequently cultured HUVECs with the conditioned media for 24 h. The results indicated that conditioned media from SW620/shTNC cells reduced the expression of ITGB3, Phospho-FAK Tyr397, Phospho-Akt Ser473 (Fig. 4C); however, it did not alter the expression of TNC (Fig. 4C). These results demonstrated that tumor-derived TNC has a positive influence on ITGB3/FAK/Akt signaling pathway.

Reduction of tumor-derived TNC impairs tubulogenesis, proliferation and migration of HUVECs in vitro
To further confirm whether TNC has an impact on tubulogenesis activity of HUVECs, we plated HUVECs on matrigel together with conditioned media from SW620, SW620/Vector and SW620/shTNC cells, respectively. As shown in Fig. 5A and B, when TNC knockdown, the incubation of HUVECs with SW620/shTNC conditioned media resulted in a 60% decrease in the formation of capillary-like structures compared with the tubules formed by HUVECs incubated with SW620 conditioned media, and similar results were obtained in cell proliferation and migration of HUVECs using CCK8, wound-healing and transwell chamber assays (Fig. 5C–G). Our results suggested that TNC has a positive influence on the proliferation, migration and tubulogenesis of HUVECs.

DISCUSSION
Tumor angiogenesis is a complex process leading to abnormalities in vascular structure and function (Sasaki et al., 1991). Similar to vasculature in individual organs, VECs are tissue-specific, which mostly depends on the tumor microenvironment (Jin et al., 2016). In this study, we compared global proteome profiles of VECs derived
Fig. 1. See next page for legend.
PI3K/AKT pathway in tumor cells can also increase the secretion of HIF-1α and VEGF. However, the activation of the PI3K/AKT pathway in angiogenesis and tumor progression were well documented (Shen et al., 2018). Many molecules in the focal adhesion complex are implicated in downstream signaling pathways, such as the Akt1 (Higuchi et al., 2013), MAPK/ERK pathway (Ye et al., 2017) and Wnt signaling (Yu et al., 2012). In this way, pathways impacted by the focal adhesion complex are as varied as apoptosis (Bouchard et al., 2008), cell proliferation (Luo et al., 2018), cell migration and angiogenesis (Zhao and Guan, 2011). In our findings, the focal adhesion-related proteins were significantly increased in TVECs as compared to controls (Table S5). Furthermore, the focal adhesion, PI3K-Akt signaling pathway, HIF-1 signaling pathway and EMT were identified as significantly and consistently proangiogenic categories in CRC as these pathway-related proteins were significantly upregulated in TVECs compared to controls.

Focal adhesion is a subcellular structure which acts as a scaffold for many signaling pathways involving integrin or the mechanical force exerted on cells (Shen et al., 2018). Many molecules in the focal adhesion complex are implicated in downstream signaling pathways, such as the Akt1 (Higuchi et al., 2013), MAPK/ERK pathway (Ye et al., 2017) and Wnt signaling (Yu et al., 2012). In this way, pathways impacted by the focal adhesion complex are as varied as apoptosis (Bouchard et al., 2008), cell proliferation (Luo et al., 2018), cell migration and angiogenesis (Zhao and Guan, 2011). In our findings, the focal adhesion-related proteins were significantly increased in TVECs as compared to controls (Fig. 2A), which indicated that the focal adhesion pathway may play a crucial role in CRC angiogenesis. Therefore, uncovering the molecular processes underlying focal adhesion hub signaling will foster the development of reasonable and feasible multimodal treatment options towards CRC.

The role of PI3K/Akt signaling pathway and HIF-1 signaling pathway in angiogenesis and tumor progression were well documented (Karar and Maiti, 2011). Hypoxia leads to the stabilization of HIF-1α and is a major stimulus for tumor cells to increase the expression of VEGF. However, the activation of the PI3K/AKT pathway in tumor cells can also increase the secretion of VEGF. Moreover, PTEN/PI3K/AKT regulates the proangiogenic factors, including HIF-1α under hypoxic conditions and controls tumor-induced angiogenesis and metastasis (Joshi et al., 2014). The HIF-1α signaling pathway and PI3K signaling pathway have been exploited for the development of new cancer therapies (Post et al., 2004; Tanaka et al., 2015; Thorpe et al., 2015). In the current study, HIF-1 and PI3K/AKT signaling pathway-related proteins were significantly upregulated in TVECs compared with controls (Fig. 2B; Fig. S1). Those findings indicate that PI3K and HIF-1 inhibitors are excellent candidates for the treatment of CRC.

EMT underlies the progression and metastasis of malignant tumors by enabling cancer cells to depart from their primary tumors, invade surrounding tissues and spread to distant organs. Endothelial-to-mesenchymal transition (EndMT) is often categorized as a specialized form of EMT. Recent studies suggested that EndMT may play a role in angiogenic sprouting by enabling the so-called tip cells, which lead an emerging vascular plexus, to migrate into adjacent tissue (Potenta et al., 2008; Welch-Reardon et al., 2015).

In our study, EMT-related proteins were significantly increased in TVECs as compared to controls (Table S5). For example, FBLN5 (fold change=−1.81) initiates EMT and enhances EMT induced by TGF-beta in mammary epithelial cells via a MMP-dependent mechanism (Lee et al., 2008). Based on the clinical application of anti-angiogenesis therapy in metastatic CRC, the EMT-angiogenesis and EMT stemness links in the CRC cells, newly synthesized drugs with antiangiogenic/anti-EMT properties could be one of the medications used in the future in the targeted therapy of patients with CRC (Gurzu et al., 2016).

In our findings, TNC was implicated in focal adhesion, PI3K/Akt signaling pathway and EMT. Several researchers have demonstrated that TNC promotes neoplastic angiogenesis in various cancers including breast cancer, glioblastoma and oral and pharyngeal squamous cell carcinoma (Aulta et al., 2003; Mai et al., 2002; Rupp et al., 2016). However, few reports have investigated the expression of TNC in the VECs of CRC and the mechanism for promoting CRC angiogenesis. In our study, we found that the expression of TNC in VECs was correlated with CRC multistage carcinogenesis. This finding suggested that TNC may have a positive effect on CRC angiogenesis and progression. As a following step, we highlighted the role of TNC in triggering ITGB3/FAK/Akt-473 signaling pathway, which promoted HUVEC migration and angiogenesis. Moreover, TNC can be secreted in large amounts by SW620 cells (Li et al., 2016), indicating that TNC is more likely to end up in the blood or other body fluids in a ‘measurable’ concentration in CRC patients, which can be used for blood-based diagnostics. Therefore, we demonstrated the potential regulation mechanism of TNC and indicated its possible use as a diagnostic and therapeutic target for CRC.

In conclusion, our study built the first proteome of VECs in CRC and investigated the role of vascular proteins in the process of colorectal angiogenesis. In addition, we demonstrated the function of TNC in regulating ITGB3/FAK/Akt signaling and promoting angiogenesis in human CRC. These findings not only provide the proteomic profiling of VECs in CRC, but also highlight some angiogenesis-related proteins, which may be the potential antiangiogenic targets and suggest new insights into the mechanism of angiogenesis in CRC.

**MATERIALS AND METHODS**

**Sample collection**

Ten fresh-frozen samples of CRC and ten matched ANC tissues were taken from the Department of Surgery, Xiangya Hospital, Central South University, China, and used for iTRAQ labeling. The patients received neither radiotherapy nor chemotherapy before curative surgery and provided a written informed consent form for the study, which was approved by the local ethical committee. All specimens were obtained from surgical resection and stored at −80°C until further investigation.

An independent set of formalin-fixed paraffin-embedded tissue specimens, namely, 30 cases of NCM, 30 cases of AD, 30 cases of CIS and 50 cases of ICC, were obtained from the Department of Pathology of the Xiangya Hospital at Central South University and used for immunohistochemical analysis.
Laser capture microdissection
The frozen samples were prepared, stained and diagnosed by pathological examination and a rapid immunohistochemical staining technique was performed in less than 13 min with the DAKO rapid Envision™-HRP system as described (Wu et al., 2010). To identify and visualize the VECs in human colorectal tissues for sequential LCM, the specific VEC marker CD31 was used for immunostaining. After the blood vessels were apparent, the glass slides were replaced with LCM-specific glass slides. The stained areas that had the characteristic morphology of VECs were captured by LCM. LCM was performed with a Leica AS LMD system to purify the cells of interest from each type of tissue as previously described (Cheng et al., 2008; Li et al., 2012). Each captured cell population was over 95% homogeneous as determined by direct microscopic visualization.

Protein extraction
LCM-enriched VECs were dissolved with an addition of lysis buffer (7 mol/l urea, 2 mol/l thiourea, 65 mmol/l dithiothreitol, 0.1 mmol/l phenylmethylsulfonyl fluoride), and followed by centrifugation (12,000 rpm, 30 min, 4°C). The protein content was determined using 2D Quantification Kit (GE Healthcare). To diminish the effects of biological variation on the proteomic results, equal amounts of proteins from ten cases of microdissected samples of CRC and ANC were pooled respectively.

Trypsin digestion and labeling with iTRAQ reagents
Trypsin digestion and iTRAQ labelling were performed according to the manufacturer’s protocol (Applied Biosystems). Briefly, 100 μg protein of each pooled sample was reduced, alkylated and then digested overnight at
The instrument was run with peptide recognition mode enabled. Collision energy was 30 eV and the underfill ratio was defined as 0.1%. The maximum IT was 50 ms and isolation window was 2 m/z. The normalized 17,500 at m/z 200. Full MS automatic gain control (AGC) target was 1e6, and resolution for higher-energy collisional dissociation spectra was set to 70,000 at m/z 200 value was determined based on the predictive automatic gain control technique. Survey scans were acquired at a resolution of 70,000 at m/z 200. The labeled digests were then mixed and dried.

Off-line 2D Liquid chromatography (LC)-electrospray ionization (ESI) tandem MS (MS/MS) analysis

The mixed peptides were first separated on a strong cation exchange column into ten fractions according to the procedure described in our previous study (Liu et al., 2017). Each fraction was dried, dissolved in buffer C (0.1% formic acid, 5% acetonitrile) and analyzed on a Q-Exactive mass spectrometer (Thermo Fisher Scientific, Waltham, MA, USA) in information-dependent mode. Briefly, peptides were separated on reverse-phase columns (EASY column, 10 cm, ID 75 μm, 3 μm, C18-A2) with an EASY-nLC™ system. Peptides were separated by a linear gradient mobile phase A (5% acetonitrile, 0.1% formic acid) and mobile phase B (84% acetonitrile, 0.1% formic acid) from 5% to 40% of mobile phase B in 60 min at a flow rate of 300 nl/min. MS data were acquired from 300-1800 m/z for fragmentation by higher-energy collisional dissociation, with up to 20 precursors selected for MS/MS and dynamic exclusion for 60 s. The target value was determined based on the predictive automatic gain control technique. Survey scans were acquired at a resolution of 70,000 at m/z 200 and resolution for higher-energy collisional dissociation spectra was set to 17,500 at m/z 200. Full MS automatic gain control (AGC) target was 1e6, maximum IT was 50 ms and isolation window was 2 m/z. The normalized collision energy was 30 eV and the underfill ratio was defined as 0.1%. The instrument was run with peptide recognition mode enabled.

Data analysis

MS/MS spectra were searched using the MASCOT engine (Matrix Science, London, UK; version 2.2) embedded into Proteome Discoverer 1.4�

Table 1. TNC expression in multistage of colorectal epithelial carcinogenic vessels

| Multistage | Number | Low (0-2) | Medium (3-4) | High (5-6) | P-value |
|------------|--------|-----------|--------------|------------|---------|
| NCM        | 30     | 23        | 6            | 1          | P<0.0001|
| AD         | 30     | 15        | 13           | 2          |         |
| CIS        | 30     | 4         | 17           | 9          |         |
| ICC        | 50     | 7         | 16           | 27         |         |

Notes: P<0.05 by linear trend chi square test, NCM versus AD versus CIS versus IC.

37°C with trypsin (mass spectrometry grade; Promega). The peptide samples were collected and then labeled with iTRAQ reagents as follows: CRC, labeled with iTRAQ 116, 117 and 118; ANC, labeled with iTRAQ 113, 114 and 115. The labeled digestes were then mixed and dried.

Fig. 3. Representative results of immunohistochemistry show the expression of TNC in vessels. Original magnification: ×200. Top panel: TNC immunostaining of NCM (A), ACP (B), CIS (C) and ICC (D). Negative staining was observed in VECs of NCM and ACP, moderate staining in CIS, and strong staining in ICC. Bottom panel: strong staining was observed in VECs of lymph nodes with metastasis (E). Negative staining was observed in VECs of lymph nodes without metastasis (F). Endothelial cell marker (anti-CD34) immunostaining of NCM (G) and ICC (H). The arrows indicate the vessels.

Table 2. Correlation between expression of TNC protein and clinico-pathological characteristics of patients with colorectal carcinoma

| Item          | Number | Low (0-2) | Medium (3-4) | High (5-6) | P-value |
|---------------|--------|-----------|--------------|------------|---------|
| Gender        |        |           |              |            |         |
| Male          | 46     | 6         | 18           | 22         | P=0.840 |
| Female        | 34     | 5         | 15           | 14         |         |
| Age           |        |           |              |            |         |
| ≥65           | 43     | 7         | 16           | 20         | P=0.655 |
| <65           | 37     | 4         | 17           | 16         |         |
| TNM stages    |        |           |              |            | P=0.022 |
| CIS           | 30     | 4         | 17           | 9          |         |
| III–IV        | 19     | 4         | 9            | 6          |         |
| Matched lymph nodes |   | 31       | 3            | 7          | 21       |
| Metastasis    | 31     | 3         | 12           | 16         | P=0.001 |
| Non-metastasis| 49     | 21        | 19           | 9          |         |

Notes: P-value determined by linear trend chi square test between TNM stages; P-value determined by Mann–Whitney U-test, male versus female, age ≥65 versus <65, metastatic lymph nodes versus non-metastatic lymph nodes.
control for protein loading and was detected using a mouse anti-ACTB antibody (1:2000; Proteintech, 20536-1-AP). Membranes were incubated with the corresponding secondary antibodies, including anti-rabbit and anti-mouse peroxidase (HRP)-linked IgG (1:1000; KPL) for 1 h.

**Cell culture, cell transfection and conditioned media preparation**

Human colon cancer cell lines SW620 and HUVECs were maintained in RPMI-1640 or DMEM medium supplemented with 10% FBS in a humidified chamber with 5% CO₂ at 37°C. SW620 cells and HUVECs were purchased from the Cell Bank of Type Culture Collection of the Chinese Academy of Sciences (Shanghai, China).

SW620 cells were cultured in six-well plates until reaching 70% of confluence, the cells were transfected with lentiviral shRNA-TNC (GGAGTACTTTATCCGTGTATT) and lentiviral control (Cyagen Bioscience Inc., Guangzhou, China) at an MOI of 20 in the presence of 5 µg/ml polybrene for 24 h and treated with 3 µg/ml puromycin for three days. The generated cell clones were tested for shRNA-TNC stable expression.

The untransfected (SW620), empty vector (SW620/Vector) and TNC shRNA plasmid-transfected SW620 (SW620/shTNC) cells were chosen for conditioned media preparation. Conditioned media were collected as previously described (Li et al., 2016). Briefly, approximately 3×10⁶ cells were grown to 70% confluency, and the medium was exchanged with serum-free RPMI-1640 medium. Conditioned media were collected 24 h after the change of media and stored at −20°C for use. For subsequent experiments, the conditioned media were filtered using a 0.22 µm filter (Millipore) and concentrated using a Millipore centrifugal filter (3 kDa). The protein concentration was determined by a standard Bradford protein assay (Thermo Fisher Scientific). After conditioned media were removed, the cell monolayer was washed, scraped and lysed in the presence of protease inhibitors. Protein concentration was determined using a Pierce BCA Protein Assay Kit (Thermo Fisher Scientific).

**HUVEC proliferation assay**

The HUVECs were plated at 2×10⁵ cells per well in 96-well tissue culture plates and cultured in a 1:1 mixture of 10% FBS complete DMEM and conditioned media (total 200 µl). The cells were cultured for 6 days. Every 24 h, 20 µl of CCK8 (5 mg/ml; Beyotime) was added to the wells, and cells were further incubated for 2 h. The absorbance of each well was read with a BioTek Instruments EL310 Microplate Autoreader at 450 nm. The CCK8 assay was performed three times in triplicate.

**Wound healing assay**

Cell migration was determined by scratch wound healing assay. Briefly, cells were grown in a 1:1 mixture of 10% FBS complete DMEM and conditioned media overnight to confluence in a six-well plate. Monolayers of cells were wounded by dragging a pipette tip. Cells were washed to remove cellular debris and allowed to migrate for 12–24 h. Images were taken at 0 h and 12 h after wounding under the inverted microscope.

**HUVEC migration assay**

Migration activity was measured by Transwell assay (Corning, 3422). Approximately 5×10⁴ HUVECs were added to the upper chamber in 200 µl of 1% FBS DMEM medium. The lower chamber contained 250 µl of 10% FBS complete DMEM and 250 µl of conditioned media. The plates were incubated for 24 h at 37°C in 5% CO₂. After incubation at 37°C for 48 h, cells were fixed with 4% paraformaldehyde and stained with 0.5% Crystal Violet. Each clone was plated in triplicate for each experiment, and each experiment was repeated at least three times.

**HUVEC angiogenesis assay**

Matrigel (BD Biosciences, 543248) was melted at 4°C, added to 48-multiwell plates (Corning) at 100 µl/well, and then incubated at 37°C for 30 min. HUVECs (4×10³ cells) were resuspended in a 1:1 mixture of 250 µl of 20% FBS complete DMEM and 250 µl of conditioned media (total 500 µl with 10% FBS). The cells were added to the wells, and after 7 h incubation at 37°C, HUVEC tube formation was assessed by microscopy. Each well was imaged under a light microscope. The numbers of branches were calculated and quantified using MacBiophotonics ImageJ software (NIH).
All statistical analyses were performed using the SPSS software package (version 13.0; SPSS, Inc.) and Prism 5.0 software (GraphPad). Data are shown as the mean±s.e.m. The difference in TNC protein expression between the two different kinds of tissue (CRC versus ANC) was analyzed using a linear trend chi-square test. The relationship between TNC expression and the clinico-pathological characteristics in patients with CRC was analyzed using the Mann–Whitney U-test and linear trend chi-square test. For in vitro assays, the data are reported as biological replicates, with technical replicates indicated in the figure legends. Student’s t-tests were used to compare the differences between groups.

Statistical analysis
All statistical analyses were performed using the SPSS software package (version 13.0; SPSS, Inc.) and Prism 5.0 software (GraphPad). Data are shown as the mean±s.e.m. The difference in TNC protein expression between the two different kinds of tissue (CRC versus ANC) was analyzed using a linear trend chi-square test. The relationship between TNC expression and the clinico-pathological characteristics in patients with CRC was analyzed using the Mann–Whitney U-test and linear trend chi-square test. For in vitro assays, the data are reported as biological replicates, with technical replicates indicated in the figure legends. Student’s t-tests were used to compare the differences between groups.
(unpaired, two-tailed) were performed to determine whether a difference between two values was statistically significant, with P<0.05 considered significant. In *vitro* assays were performed in triplicate unless otherwise stated. P-values <0.05 were considered statistically significant (**P<0.05; ***P<0.001).

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**Competing interests**

The authors declare no competing or financial interests.

**Author contributions**

Methodology: Z. Xie; Software: X.L.; Validation: H.H.; Formal analysis: Y.F., Z. Xiao; Investigation: G.W., Q.Y., M.Z.; Resources: Y. Cai; Data curation: Y.H.; Writing - original draft: G.W.; Writing - review & editing: M.L., Y.Z.; Supervision: Z.C.; Project administration: Y. Chen; Funding acquisition: F.P.

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**Supplementary information**

Supplementary information available online at http://bio.biologists.orglookup doi:10.1242/bio.042838.supplemental

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