GM130 regulates epithelial-to-mesenchymal transition and invasion of gastric cancer cells via snail

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Abstract: Gastric cancer is one of the most common causes of digestive tract tumor. Despite of recent advances in surgical techniques and development of adjuvant therapy, the underlying mechanisms of gastric cancer remain poorly understood and relevant insight into novel treatment strategies using gene target remains incomplete. Recently, several studies report that epithelial to mesenchymal transition (EMT) is a crucial process for the invasion and metastasis of epithelial tumors; however, the molecular mechanisms underlying this transition are unknown. As a cis-Golgi matrix protein, GM130 plays an important role in cell cycle progression and transport of protein in the secretory pathway. In this study, we found that GM130 expression has a positive correlation with the pathological differentiation and tumor node metastasis (TNM) stage of gastric cancer. High GM130 expression levels also predict shorter overall survival of gastric cancer patients. RNA interference-mediated knockdown of GM130 expression increased epithelial marker (E-cadherin) and decreased mesenchymal marker (N-cadherin and vimentin) expression in gastric cancer cells, suppressing cell invasion, and tumor formation. Furthermore, we found that GM130 upregulated expression of the key EMT regulator Snail (SNAI1), which mediated EMT activation and cell invasion by GM130. Taken together, our study indicates GM130 may be a promising therapeutic biomarker for gastric cancer.

Keywords: GM130, gastric cancer, EMT, invasion, migration

Introduction

Gastric cancer is the most common type of tumor in humans. Despite of recent advances in surgical techniques and development of chemotherapy/radiotherapy, the mortality of gastric cancer remains high and 5-year survival rate lower than 30% [1, 2]. A recently elucidated secretory pathway is attracting considerable interest as a promising anticancer target. To provide insight that will enable the development of new therapeutic strategies, it is crucial to elucidate the molecular mechanisms that promote the migration and invasion properties of gastric cancer. Recent studies have shown that a morphologic conversion, known as epithelial-to-mesenchymal transition (EMT), is associated with the acquisition of malignant characteristics in gastric cancer cells [3, 4].

GM130, a cis-Golgi Matrix Protein, is a peripheral membrane protein strongly anchored to the Golgi member which is isolated from the detergent and salt resistant Golgi matrix. Besides maintain of the Golgi structure and stacking of Golgi cisternae, it also is involved in the regulation of ER-to-Golgi transport and glycosylation, cell cycle progression, and the physiology of multicellular organisms. However, emerging evidence has indicated that GM130 has unexpected roles in the control of higher order cell functions such as cell division, polarization and directed cell migration and various disease. It has been reported that GM130 is overexpressed in cervical cancer cell lines [5, 6], lung cancer cell lines, and prostate cancer cell lines. Knockdown of GM130 inhibits the lateral fusion of Golgi stacks and disturbs the uniform distribution of Golgi enzymes affecting proper glycosylation of membrane and secretory proteins and also issues in delay of ER-to-Golgi transport or a partial inhibition. In addition, suppression of cells proliferation, invasion, angiogenesis and increase of autophagy have been reported in A549 cells which lack GM130 [7, 8].
GM130 promotes invasion of gastric cancer

A number of possible reasons may explain why this novel role for GM130 and in the poor prognosis of oncogenic gastric cancer has not been previously described. In this study, we report that the expression of GM130 was significantly up regulated in gastric cancer tissues compared with their noncancerous counterparts as shown by immunohistochemical staining and RT-PCR. In addition, siRNA-mediated knockdown GM130 expression inhibited cell proliferation and decreased cancer cell migration and invasion in vitro. We show that GM130 decreases the expression of the epithelial markers E-cadherin but increases the expression of mesenchymal markers fibronectin and N-cadherin. The results indicate GM130 induces EMT and cell invasion in gastric cancer cells by promoting Snail expression in gastric cancer.

Materials and methods

Cell lines and tissue samples

Three human gastric cancer cell line MKN-45, SGC-7901 and MKN-28 were purchased from the cell bank of Chinese Academy of Sciences (Shanghai, China). All the cell lines were maintained in RPMI-1640 supplemented with 10% FBS and 1% antibiotics. A total of 84 paraffin-embedded gastric cancer samples of which 42 were with paired adjacent tissues were obtained through the Bayannaoer City Hospital, Inner Mongolia. All gastric cancer patients who had not received any pre-operative chemotherapy, radiotherapy or immunotherapy were diagnosed by at least two experience pathologists independently. This study was approved by the Ethics Committee of The Bayannaoer City Hospital, Inner Mongolia.

Immunohistochemistry

All tissue sections (4 µm) were routinely dewaxed, rehydrated, and then was subjected to heat with citrate buffer in a microwave oven for antigen retrieval. Endogenous peroxidase activity was blocked for 15 min in 3% hydrogen peroxide in deionized water. The slides were then added blocking normal goat serum and followed by incubation at 4°C overnight with GM130 rabbit polyclonal antibody (1:100). Sections were incubated with biontinated second antibody and the streptavidin-biotin peroxidase complex (ZSGB-BIO, Beijing, China). In addition, tissue sections were stained with diaminobenzidine (ZSGB-BIO). Ultimately, all the treated sections were lightly counterstained with hematoxylin, dehydrated and mounted. Expression of GM130 was evaluated as the percentage of positive cells in a specimen, and by staining intensity as described previously [9]. The percentage of positive cells was evaluated quantitatively and scored as (0 = 0%, 1 = 1-25%, 2 = 26-50%, 3 = 51-75%, 4 = 76-100%) and the intensity was graded as follows (0 = none, 1 = weak, 2 = moderate, 3 = strong). The final result of GM130 expression was the product of scores of percentage of positively stained cells and the staining intensity (negative, 0-1; weakly positive, 1-2; moderately positive, 2-3; strongly positive, ≥ 3).

Quantitative real-time PCR

Total RNA was extracted from cultured cells using EZNA Total RNA Kit (OMEGA Bio-tek, USA), and cDNA was generated using PrimeScript RT reagent Kit with gDNA Eraser (TaKaRa, Otsu, Japan). Quantitative real-time PCR was performed using the SYBR Premix ExTaq II (TliRNaseH Plus) (TaKaRa, Otsu, Japan) with a CFX96 Real-Time PCR Detection System (Bio-Rad, Hercules, CA, USA).

Western blot analysis

After measuring the protein concentration of the lysates using the BCA method (Beyotime, China), equal amounts (35 µg) of protein were separated by SDS-PAGE, blotted onto PVDF membranes (Millipore, Billerica, MA). The membranes were blocked with TBST containing 5% skim milk for 1 hour at room temperature, incubated with the corresponding primary antibodies overnight at 4°C and then with secondary antibodies conjugated to horseradish peroxide for 2 h at room temperature. The protein was visualized using enhanced chemiluminescence (GE Healthcare Biosciences).

SiRNA transfection

MKN-45 cell line was transfected with three kinds of siRNAs targeting GM130 (GenePharma, Shanghai, China). The sequences of the siRNAs used to suppress GM130 expression was: sense, 5’-CUGGACUCAGCUAGUGUAATT-3’, and antisense, 5’-UUACAUA GCUGGAGUCCAGTT-3’. The scrambled siRNA as negative control was: sense, 5’-UUCU CCGAACGUGUCAGUGUAATT-3’, and antisense, 5’-ACGUAGACGUUCGGAGAATT-3’. Cell transfection was carried out using Lipo-
GM130 promotes invasion of gastric cancer

Fectamine 2000 (Invitrogen) according to manufacturer's instructions.

Cell migration and invasion assays

For transwell migration assays, $2 \times 10^4$ cells were seeded in the top chamber with the non-coated membranes (24 well insert, 8.0-mm, BD Bioscience). Cells were seeded in a serum-free medium and then migrated toward complete growth medium. For the invasion assay, cells were serum-starved overnight and $2 \times 10^4$ cells were seeded in a Matrigel-coated chamber and cultured for 48 h. The invaded or migrated cells were fixed with 70% methanol and stained 0.5% crystal violet. Cells invaded or migrated the lower surface of filters were counted in five randomly selected fields. All experiments were carried out in triplicate.

Luciferase reporter assay

The previously reported promoter region for Snail (-1, 558/+92) was amplified and cloned into the pGL4 vector (Promega) [10]. 293T cells were transfected with the pGL4-Snail/promoter together with pQCXIP-GFP-GM130 and pRTK-Luc to normalize transfection efficiency. Forty-eight hours later, the activities of firefly luciferase and Renilla luciferase were measured.

Figure 1. Expression of GM130 in human gastric cancer patient specimens and cell lines. (A) Expression of GM130 in normal gastric tissue and different stage gastric cancer (400×). (B) Over-expression of GM130 at mRNA level in gastric cancer tissues. Expression of GM130 in normal and gastric cancer tissue was analyzed by qRT-PCR (C) Kaplan-Meier plots of GM130 expression in 20 cases of gastric cancer patients. Overall survival rate was performed by log-rank test. P < 0.05 indicate significant differences between two groups. (D, E) Expression of GM130 in breast cancer cell lines (P < 0.001).
GM130 promotes invasion of gastric cancer

using the Dual Luciferase Reporter Assay System (Promega). Luciferase activity was measured in triplicate, and 3 independent experiments were carried out. To measure luciferase activity in the absence of GM130 expression, MKN-45 cells were transfected with control or GM130 siRNA together with pRTK-Luc and pGL4-Snail/promoter. Seventy-two hours later, luciferase activities were measured.

Statistical analysis

Statistical analysis for cell invasion, colony formation, and tumor formation in mice was conducted using the Student t test. P < 0.05 was considered statistically significant. The association between GM130 expression and tumor stage was determined by test.

Results

Upregulation of GM130 expression in human gastric cancer

To determine the clinicopathological significance of GM130, GM130 expression was evaluated by immunohistochemistry in 84 pairs of human gastric and adjacent non-cancerous counterparts. We found that strong expression of was observed in some cancer tissues, and the expression was correlated with tumor stage (Figure 1A). Increased GM130 expression was also found to correlate with shorter overall survival (P < 0.001) of patients (Figure 1B). Of the 84 samples, we observed that GM130 expression was more frequently in carcinomas (88.1%) than adjacent non-cancerous tissues cells (52.4%). To explore the role of GM130 in human gastric cancer progression, we first evaluated GM130 expression levels in various gastric cancer cell lines. Interestingly, we found that GM130 expression was related with the malignant characteristics of gastric cancer cells. MKN-45 was more invasive than other gastric cancer cell lines and was able to grow under anchorage-independent conditions. These findings show that GM130 is overexpressed in primary human gastric cancer and metastatic gastric cancer cells.

Downregulation of GM130 reduces migration and invasion of MKN-45 cells

To determine the effect of GM130 on cell migration and invasion. Migration was assessed using a short-term transwell assay (Figure 2A and 2B). And then, we assessed cell invasiveness, we used Matrigel-coated Boyden chambers. GM130 knockdown reduced cell migration and invasion of MKN-45 cells (Figure 2C and 2D).

GM130 regulates EMT in gastric cancer cells

Epithelial-to-mesenchymal transition is associated with malignant properties, such as invasion and anchorage-independent growth. We hypothesized that the upregulation of GM130
GM130 promotes invasion of gastric cancer

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may promote EMT in gastric cancer cells. High metastatic gastric cancer cell line (MKN-45) demonstrated elevated GM130 expression, were spindle shaped and exhibited reduced cell contact. GM130 knockdown resulted in a dramatic shift from spindle shaped to a cobblestone-like morphology. To test this hypothesis, the expression levels of epithelial and mesenchymal markers were examined. By western blot and immunofluorescence, we found that depletion GM130 expression caused an increase in epithelial marker (E-cadherin), but a decrease in mesenchymal markers (N-cadherin and vimentin) in MKN-45 cells (Figure 3A). These results indicate that GM130 has an important role in EMT regulation in gastric cancer cells.

GM130 directly regulates Snail expression

Aberrant expression of EMT transcription factors contributes to the appearance of an invasive phenotype by suppressing E-cadherin and inducing EMT in a wide variety of human cancers [11, 12]. We aimed to identify transcription factors whose expression is regulated by GM130. RNA was extracted from MKN-45 with GM130 siRNA, and the mRNA and protein expression of EMT-related transcription factors were examined by qRT-PCR and immunoblot. Among the transcription factors examined, Snail expression was decreased by the knockdown of GM130 in MKN-45 cells (Figure 4A and 4B).

We next conducted luciferase assays to determine whether Snail promoter activity was regulated by GM130 expression. 293T cells were transiently cotransfected with full-length GM130, and a reporter construct in which the human Snail promoter region was cloned upstream of firefly luciferase (pGL4-Snail/promoter). Exogenous expression of GM130 increased Snail promoter activity approximately 6-fold (Figure 4C). In addition, the transfection of GM130 siRNA into MKN-45 cells decreased promoter activity 60% to 80% compared with that of luciferase siRNA (Figure 4D). These results demonstrate that GM130 induces the transcription of the Snail gene.

Discussion

Gastric cancer is a very common disease worldwide and the second most frequent cause of cancer death. During gastric cancer carcinogenesis, genomic damage accumulate, affecting cellular functions such as self-sufficiency in growth signals, escaping antigrowth signals, apoptosis resistance, Sustained replicative potential, angiogenesis induction, and invasive or metastatic potential [13-15]. In addition, a cag pathogenicity island methylator phenotype (CIMP) has been a third pattern of genomic instability [16-18]. However, the exact mechanism which implicated in carcinogenesis and progression has not been fully elucidated.
GM130 promotes invasion of gastric cancer

Recently, novel pathway involving the ER, Golgi apparatus, and lysosomes are being recognized as potential targets for therapeutic interventions. GM130 plays an important role in the maintenance of Golgi structure and transport of proteins and lipids in the secretory pathway. Recent reports support the view that the down-regulation of GM130 inhibits the protein transportation between ER and Golgi and induces autophagy. While, autophagy contributes to tumor suppression and defects of autophagy are associated with tumorgenesis [18-20]. In this study, we first examined the expression of GM130 in human gastric cancer tissues and cell lines to explore the molecular mechanisms of gastric cancer tumorgenesis, and then to reveal new predictive markers. Our results showed that GM130 protein was primarily expressed in the cytoplasm of gastric cancer cells and GM130 expression was evidently increased in gastric cancer specimens compared with adjacent normal tissues. Furthermore, also immunohistochemistry results demonstrated that the overexpression of GM130 was associated with differentiation grade and TNM stage in gastric cancer. Recent evidence supports aerosol delivery of GTP-SPE/shGM130 can suppress tumor progression to lung adenocarcinoma. In addition, in support of a role for GM130 in tumor progression, knockdown of GM130 in a Caco-2 cells lead to a loss of E-cadherin expression, a phenotype that is often considered to be linked to tumor progression and a loss of epithelial identity and is in line with the notion that loss of polarity is a typical phenomenon in epithelial-mesenchymal transition [5, 6, 21, 22]. We also observed that GM130 expression, at both the mRNA and protein levels, in the human gastric cancer lines MKN-28, SGC-7901, MKN-45. Our results are consistent with GM130 might be involved in tumorgenesis and the progression of cancer.

To further explore the effects of GM130 on proliferation, invasion and migration in carcinoma
GM130 promotes invasion of gastric cancer

cell lines. We transfected the GM130 sequence-specific siRNA with Lipofectamine 2000 into the MKN-45 cells, which had high invasive and metastatic potential and expressed high levels of GM130 proteins. We found that the cell proliferation and growth of GM130-siRNA cells was markedly inhibited and the levels of invasion and migration capacity was significantly lower in comparison with the cells in the negative control group, indicating that downregulation of GM130 in MKN-45 gastric cancer lines suppressed proliferation and inhibited migration and invasion. In addition, we also found that downregulation of GM130 decreased the protein level of MMPs. It has been shown that small interfering RNA against GOLGA2/GM130 results in inhibition of cell proliferation, angiogenesis and invasion in A549 cells.

Recent studies reported that EMT-associated transcription factors confer cancer cells with malignant characteristics, such as invasion, metastasis, and resistance to chemotherapy. Interestingly, silencing GM130 can induce E-cadherin downregulation in Caco-2 cells. The loss of E-cadherin expression not only disrupts cell-cell adhesion but also activates multiple pathways that induce cellular migration, invasion, and metastasis. Divergent findings have been made concerning the role of GM130 in cancer, which seems to depend on the type of cancer investigated. For instance, in lung cancer, the downregulation of GM130 induces autophagy and suppresses angiogenesis, invasion. GM130 expression is elevated in lung cancer but downregulated in colorectal and breast cancer [6, 21, 22]. Future experiments are necessary to clarify the proposed mechanism to the effect of GM130 on tumor cells. These studies are currently under way. And then, this is the first report show that variation in GM130 level participate in tumorigenesis and the progression of gastric cancer, as well as we have identified GM130 as a novel regulator of EMT and cell invasion. In addition, the induction of Snail expression was shown to be critical for the GM130-mediated promotion of EMT and cell invasion. Thus, GM130 is a critical regulator of cancer progression and may be an important target for human gastric cancer treatment.

Disclosure of conflict of interest

None.

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GM130 promotes invasion of gastric cancer


