Augmented O-GlcNAcylation of AMP-activated kinase promotes the proliferation of LoVo cells, a colon cancer cell line

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Colon cancer is a major health problem because of high prevalence and high mortality rates in Western countries, such as North America, Australia, and Europe. In 2015, the American Cancer Society reported approximately 1.6 million people diagnosed with colon cancer and an annual mortality of 590,000 patients. Thus, colon cancer is the third most prevalent and most common cause of cancer-related deaths in both men and women.¹

Diabetes has been predicted to be a significant risk factor for various cancers, including colon cancer, in recent years.² Many previous studies have suggested the association between type 2 diabetes and the prevalence of various cancers, including colon, liver, breast, pancreas, and endometrial cancers.³–⁷ Although insulin or insulin-like growth factor-1 has been associated with increased colon cancer risk in diabetic patients,⁸–⁹ further studies are required to elucidate the mechanisms by which a diabetic condition increases the occurrence of colon cancer.

Protein O-GlcNAcylation is a critical post-translational modification in which an O-GlcNAc residue is enzymatically transferred to serine/threonine residues of certain proteins.¹⁰–¹² Like phosphorylation, protein O-GlcNAcylation has been observed in various proteins, including transcriptional factors, cytoskeletal proteins, nuclear pore complex proteins, and tumor promoter proteins. Both protein modifications target the serine/threonine residues and have often been reported to participate in competitive interplay with each other.

Protein O-GlcNAcylation is regulated by two enzymes, namely, OGT, which transfers GlcNAc from the donor substrate, UDP-GlcNAc, to serine/threonine residues on certain proteins, and OGA, which in turn cleaves GlcNAc from the proteins.¹³ The production of UDP-GlcNAc, as well as the activities of OGT and OGA, is one of the rate-limiting steps for O-GlcNAcylation. Uridine diphospho-GlcNAc is synthesized from glucose through the hexosamine pathway, and GFAT is the rate-limiting enzyme in the pathway. Thus, the rate of protein O-GlcNAcylation is dependent on glucose concentrations and the activities of OGT, OGA, and GFAT.¹⁰ Uridine diphospho-GlcNAc has been reported to be elevated under high blood sugar conditions by increased
Corresponding to the amino-terminal sequence of human was produced by immunizing animals with a synthetic peptide. Antibodies were purchased from Cell Signaling Technology for AMPK silencing in LoVo cells was carried out by phosphorylating CREB-regulated transcription coactivator 2. The mTOR pathway, which accelerates cellular growth, is also regulated by AMPK. AMP-activated kinase activation suppresses the mTOR pathway, which in turn inhibits the synthesis of new proteins. Intriguingly, in some cases, AMPK could also act as a tumor suppressor through induction of G1/S arrest in coordination with the p53 tumor suppressor. Thus, AMPK activation is expected to negatively regulate tumor growth and can therefore serve as a promising therapeutic strategy against tumors.

Recently, AMPK was reported to be modified through O-GlcNacylation under high-glucose conditions. Here, we examined whether protein O-GlcNacylation promotes in vitro and in vivo growth of a human colon cancer cell line, LoVo cells, and determined the AMPK-mediated mechanisms by which protein O-GlcNacylation promotes tumor growth.

Materials and Methods

Antibodies. Anti-phospho-AMPK (T172) (pAMPK), AMPK, phospho-ACC (S79), ACC, phospho-mTOR (S2448), mTOR, p70 phospho-ribosomal kinase (S424) (pS6K), and p70 S6K (S6K) antibodies were purchased from Cell Signaling Technologies (Danvers, MA, USA). AMP-activated kinase antibody was produced by immunizing animals with a synthetic peptide corresponding to the amino-terminal sequence of human AMPKz. Anti-O-GlcNac (RL-2) and anti-OGT antibodies were obtained from Novus Biologicals (Littleton, CO, USA) and Santa Cruz Biotechnology (Dallas, TX, USA), respectively. Anti-β-actin and α-tubulin antibodies were obtained from Sigma-Aldrich (St. Louis, MO, USA) and MBL (Nagoya, Japan).

Chemicals. Thiamet G was purchased from Enzo Life Science (Farmingdale, NY, USA). Both AICAR and BML-275 were obtained from Wako Pure Chemical Industries (Osaka, Japan). Cell Counting Kit-8 was purchased from Dojindo (Kumamoto, Japan).

Cell culture. LoVo cells were obtained from Riken Biorepository Center (Riken BRC, Tsukuba, Japan). Cells were maintained in DMEM/10% FBS containing 100 μg/mL streptomycin and 100 U/mL ampicillin.

Transfection. Transfection of pCAGGS/Ogt for OGT overexpression or siRNA against AMPK (Santa Cruz Biotechnology) for AMPK silencing in LoVo cells was carried out using TransIT-X2 system (Mirus Bio, Madison, WI, USA) according to the manufacturer’s instructions.

Cell growth assays. WST-8 assay. Anchorage-dependent cell growth was measured spectrophotometrically using CCK-8 (Wako Pure Chemical Industries) according to the manufacturer’s instructions. Briefly, LoVo cells were plated onto a 96-well plate at 2 × 10^3 cells/well and incubated overnight to allow cell attachment. The cells were then incubated with or without inhibitors for varying time periods (24 or 48 h) either in DMEM containing 10% FBS or in serum-free DMEM. To measure cell proliferation, cells were incubated with a WST-8 reagent for 1 h and absorbance was measured at a wavelength of 450 nm using a microplate reader (Model 680; Bio-Rad, Hercules, CA, USA).

Colony formation assay. Anchorage-dependent cell growth on 6-well plates was evaluated by colony formation assay as described previously. First, the bottom layer was prepared using 2.5 mL of 0.5% low-melting agarose (Agarose SFR: Amresco, Solon, OH, USA) dissolved in growth medium as described earlier. LoVo cells were prepared at a density of 1 × 10^5 cells/mL. Approximately 50 μL of cell suspension and 1.4 mL of 0.3% low-melting agarose in the growth medium were overlaid on the bottom of each well. To prevent moisture loss, an additional 0.5–1 mL growth media was overlaid, and PBS was added in between spaces. After 4 weeks, colonies were stained with 0.05% crystal violet, counted, and then imaged.

In vivo assay of LoVo cells. Transplantation of LoVo cells into BALB/c-nu/nu mice. BALB/c- nu/nu mice (male, 5 weeks old) were purchased from Japan SLC (Shizuoka, Japan). The mice was maintained in pathogen-free conditions, fed with standard chow (Clea Japan, Tokyo, Japan) and kept in comfortable conditions at 20°C in appropriate, humidity, and with 12:12-h light : dark cycles, in the Division of Research Animal Laboratory, Osaka Medical College (Takatsuki, Japan) during the experiment. Painful procedures were carried out under appropriate anesthesia as described below. LoVo cells at a density of 5 × 10^6 cells in 100 μL PBS were s.c. grafted onto the backs of BALB/c- nu/nu mice (5 weeks old, n = 6 for PBS group; n = 6 for TMG group; total, n = 12) under anesthesia with tribromoethanol (300–400 mg/kg), i.p. Each mouse was grafted at two places (right and left side of the back). Thiamet G was given at 10 mg/kg prior to grafting, and subsequently every other day until the mice were killed. Tumor volumes were measured once a week and calculated using the formula: V = length × width^2 / 2, length > width. After 6 weeks, mice were killed with excess pentobarbital sodium (150–200 mg/kg), i.p. Finally, 6 of 12 transplanted cells in each group formed tumors and were subjected to subsequent analyses. The developed tumors were excised, weighed, and then fixed with 4% paraformaldehyde in PBS for histological analyses. The Osaka Medical College Animal Experiment Ethical Committee approved the animal experiment (permission number: 28033) and all the procedures were carried out in accordance with The Regulations of Animal Experiments of Osaka Medical College (http://www.osaka-med.ac.jp/dps/eca/info.html?rule).

Western blot analysis. Preparation of cell lysates. Cells were lysed with a lysis buffer (1% Triton X-100, 0.25% deoxycholic acid, and 0.1 M NaCl, pH 7.4) containing a protease inhibitor cocktail (Nacalai Tesque, Kyoto, Japan). Excised tumors were homogenized with a Potter-type homogenizer (Model LR-41C; Yamato Scientific, Tokyo, Japan). Protein concentrations of the lysates were determined using a BCA protein assay kit (Pierce, Rockford, IL, USA). To detect AMPK O-GlcNAcylation, lysates were subjected to co-immunoprecipitation with anti-
AMPK antibody conjugated with magnetic beads (Bio-Rad). Precipitates were subjected to Western blot analysis using the anti-\(\text{O-}\text{GlcNAc}\) antibody (RL-2) as described below.

Sodium dodecyl sulfate–PAGE and Western blot analysis. Proteins were subjected to SDS-PAGE. Separated proteins were transferred onto a PVDF membrane (Millipore, Billerica, MA, USA) using a Trans-blot SD semi-dry electrophoretic transfer cell (Bio-Rad). The membrane was blocked with 5% skim milk, incubated with the appropriate primary antibody, followed by the appropriate HRP-conjugated secondary antibody, and then developed using a Luminata Forte Western HRP substrate (Millipore). Images were captured using a Chemidoc system (Bio-Rad).

Statistical analysis. All experiments were carried out at least twice. Representative results were subjected to statistical analyses. All statistical analyses were undertaken using MEPHAS (http://www.gen-info.osaka-u.ac.jp/testdocs/tomocom/).

Results

Thiamet G, an OGA inhibitor, promoted growth of LoVo cells. The association between diabetes and cancer has received...
increasing research attention in recent studies. High blood sugar levels are known to increase O-GlcNAcylation.\(^{(30,31)}\) O-GlcNAcylation has been reported to be elevated in various types of cancer, including colon cancer, and is often accompanied by increased OGT.\(^{(32)}\) To elucidate the effects of increased O-GlcNAcylation on tumor growth of colon cancer in diabetic patients, we examined the growth of LoVo cells in the presence or absence of TMG, which increases O-GlcNAcylation. The growth assay was carried out in serum-free medium. As expected, the growth of LoVo cells was significantly higher both at 24 and 48 h after TMG treatment in a concentration-dependent manner (Fig. 1a). Given that AMPK is recognized as a target for the treatment of metabolic syndrome and type 2 diabetes,\(^{(33)}\) and has been reported to be a

![Figure 2](image)

**Fig. 2.** Ogt overexpression accelerated cell growth, and inactivated or activated the AMP-activated kinase (AMPK) signaling pathway or mTOR pathway, respectively, in LoVo cells. Ogt cDNA or mock DNA was transfected into LoVo cells. Twenty-four hours later, the cells were plated into 6-well or 96-well plates. (a) Cells in 6-well plates were collected after 48 h, lysed, and subjected to Western blot analysis to examine O-GlcNAcylation levels using RL-2. (b) Cells in 96-well plates were used to measure cell growth. The measurement was started the next day (0 h) using CCK-8. Growth was measured at 0, 24, and 48 h. **0.01 < \(P\) < 0.05, Student’s t-test. (c–f) Lysates in (A) were also analyzed for the phosphorylation levels of acetyl CoA carboxylase (ACC) (c), AMPK (d), mTOR (e), and S6 kinase (S6K) (f) with Western blot analysis. The phosphorylation and expression levels of these proteins were analyzed using respective anti-phospho- or total protein antibodies. The experiment was carried out several times and representative data are shown. Band densities were measured and ratios of phosphorylated to total proteins were analyzed statistically with Student’s t-test. *0.01 < \(P\) < 0.05. Molecular weight markers are indicated.

![Figure 3](image)

**Fig. 3.** AMP-activated kinase (AMPK) inhibition accelerated cell growth and inactivated the mTOR pathway in LoVo cells. (a) LoVo cells were cultured in serum-free DMEM with the indicated concentrations of BML-275 for 24 h. Cells were then lysed and subjected to Western blot analysis using anti-phospho-AMPK (pAMPK) and phospho-mTOR (pmTOR) antibodies. The total expression levels of AMPK and mTOR were used as controls. Experiments were carried out several times and representative data are shown. Ratios of phosphorylated to total AMPK or mTOR were analyzed statistically with Dunnett’s test. *\(P\) < 0.05. Molecular weight markers are shown. (b) LoVo cells were cultured with the indicated concentrations of BML-275, an AMPK inhibitor, for 24 h. Growth was measured using CCK-8. Absorbance rates of the cells with the indicated concentrations of BML-275 relative to that of ones without BML-275 were analyzed statistically with Steel’s test. *\(P\) < 0.05.
metabolic tumor suppressor (Ogt) we hypothesized that increased O-GlcNAcylation with TMG treatment can affect AMPK activity by competing with O-GlcNAcylation at Thr172. Results showed that TMG treatment for 24 h led to the decreased phosphorylation and increased O-GlcNAcylation of AMPK (Fig. 1b,c). We also examined the effect of O-GlcNAcylation on anchorage-independent growth. Results revealed that the numbers of colonies were higher in TMG-treated cells compared to controls (Fig. 1d). Collectively, elevated O-GlcNAcylation combined with TMG treatment enhanced the growth of LoVo cells in both anchorage-dependent and -independent manners and simultaneously reduced AMPK phosphorylation, which is potentially mediated by increased O-GlcNAcylation.

**Ogt overexpression promoted growth of LoVo cells.** O-GlcNAcylation is regulated by OGT and OGA. Chemical inhibition of OGA with TMG treatment increased O-GlcNAcylation. To further examine the influence of O-GlcNAcylation on the growth of LoVo cells, we alternatively overexpressed Ogt in LoVo cells, which genetically increased O-GlcNAcylation. As expected, O-GlcNAcylation was increased with the overexpression of Ogt in the cells (Fig. 2a). The growth of Ogt-overexpressing LoVo cells was also significantly higher both at 24 and 48 h compared to that of the mock-transfected cells (Mock), which was consistent with the TMG treatment data (Fig. 2b). We next examined the phosphorylation of AMPK and the main downstream substrate, ACC, in Ogt-overexpressing LoVo cells. As expected, the phosphorylation of both molecules was significantly compromised in the Ogt-overexpressing cells (Fig. 2c,d). We also examined the phosphorylation of mTOR, which is strongly enhanced in various cancers, and its main downstream substrate, S6K. Intriguingly, the phosphorylation of mTOR as well as S6K was significantly increased in the Ogt-overexpressing cells (Fig. 2e,f). Collectively, increased O-GlcNAcylation by Ogt-overexpression as well as TMG significantly accelerated the growth of LoVo cells, which was accompanied with a compromised AMPK pathway and reciprocally increased mTOR pathway.

**Treatment with BML-275, a specific AMPK inhibitor, accelerated growth of LoVo cells.** Treatment with TMG led to a dose-dependent increase in AMPK O-GlcNAcylation in parallel with AMPK dephosphorylation (Fig. 1b,c). To verify whether TMG-induced growth promotion of LoVo cells was mediated by AMPK activity, we treated LoVo cells using an AMPK-specific inhibitor, BML-275, and measured cell growth. BML-275 effectively inhibited AMPK phosphorylation in a dose-dependent manner (Fig. 3a). Phosphorylation of mTOR was increased following BML-275 treatment (Fig. 3a), consistent with previous results (Fig. 2e). The growth rates of LoVo cells increased in proportion to mTOR phosphorylation (Fig. 3b), which further supported the idea that inhibition of AMPK phosphorylation by O-GlcNAcylation can enhance cell growth by activating the mTOR pathway.

**Silencing of AMPK accelerated growth, but abolished TMG-induced growth acceleration, in LoVo cells.** To further verify that increased O-GlcNAcylation by either TMG or Ogt overexpression accelerates the growth of LoVo cells through the AMPK signaling pathway, we silenced AMPK expression in LoVo cells using siRNA. The AMPK expression using an siRNA
against AMPK (siAMPK) was silenced by approximately 50% compared to the control (Fig. 4a). We examined the growth of the siRNA-treated cells at the basal state or with various concentrations of TMG treatment. The LoVo cells with siAMPK (siAMPK cells) grew significantly faster than ones with siCTR (siCTR cells) at the basal state (Fig. 4b). Growth acceleration in proportion to higher concentrations of TMG in siAMPK cells was significantly lower compared to that in siCTR cells, which directly indicated that TMG accelerated the growth of LoVo cells through the AMPK signaling pathway. O-GlcNAcylation level at any concentration of TMG treatment was not significantly different between the cells (Fig. 4d). In contrast, phosphorylation of mTOR and S6K was significantly higher in siAMPK cells at the basal state (Fig. 4e,f). In siCTR cells, only high-dose (1 μM) TMG treatment significantly upregulated both mTOR and S6K phosphorylation (Fig. 4e,f), which is consistent with the data that AMPK phosphorylation was downregulated by approximately 50% with 1 μM TMG treatment (Fig. 1b). These results strongly suggest that O-GlcNAcylation regulated the growth of LoVo cells by control of the AMPK signaling pathway. The AMPK signaling pathway could be one of the potent effectors in O-GlcNAcylation-induced tumor growth.

Treatment with AICAR abolished TMG-induced growth acceleration in LoVo cells. To further validate whether AMPK directly mediated TMG-induced growth acceleration, LoVo cells were treated with AICAR, a specific AMPK activator. As shown in Figure 5(a), AICAR significantly increased AMPK phosphorylation and simultaneously reduced mTOR phosphorylation in a dose-dependent manner. Next, we examined the effects of AICAR on the growth of LoVo cells. LoVo cells were cultured in DMEM containing 10% FBS, treated with AICAR at the indicated concentrations, and incubated for 24 h, after which cell growth was measured. Thiamet G treatment was found to inhibit the growth of LoVo cells in a dose-dependent manner (Fig. 5b). Although 1 mM AICAR significantly suppressed cancer cell growth, lower AICAR concentrations did not produce significant effects (Fig. 1a). We next examined
whether treatment with varying concentrations of AICAR influenced the growth-promoting effects of TMG. Based on growth assay using 10% FBS containing medium, 1 μM TMG still significantly enhanced the growth of LoVo cells (Fig. 5c). Moreover, co-administration of AICAR with TMG completely abolished the TMG-induced cell growth (Fig. 5c), implying that TMG-induced growth of LoVo cells was mediated, at least partly, by the AMPK pathway.

LoVo cell-derived tumors transplanted into TMG-treated BALB/c-nu/nu (nude) mice were larger than those transplanted into control mice. So far, we have shown that increased protein O-GlcNAcylation accelerated the in vitro growth of LoVo cells, which could be due to inhibition of the AMPK activity and reciprocal activation of the mTOR pathway. We then used a TMG-treated mouse model to determine whether the same results are observed in vivo. Daily i.p. injection of TMG at 10 mg/kg was reported to increase protein O-GlcNAcylation in mice.(37) LoVo cells were transplanted onto the backs of nude mice at 5 × 10⁶ cells per spot. Thiamet G was given following transplantation and then treated continuously every other day. Tumor sizes were measured weekly, and mice were killed at 6 weeks after transplantation. Tumors excised from TMG-treated mice were observed to be larger than those excised from control mice (Fig. 6a). As shown in Figure 6(b), the sizes of tumors from mice with TMG were varied, but the average tumor volume was significantly larger in TMG-treated mice than in control mice (Fig. 6c).

Increased O-GlcNAcylation of AMPK in LoVo cell-derived tumor tissues excised from TMG-treated mice reduced AMPK phosphorylation and activated the mTOR pathway. We examined O-GlcNAcylation and phosphorylation of proteins excised from LoVo cell-derived tumor tissues. Tumors from nude mice were surgically removed and homogenized using a potter-type homogenizer as described in “Materials and Methods”. O-GlcNAcylation was higher in tumors derived from TMG-treated mice than in tumors derived from control mice (Fig. 7a). Accordingly, AMPK phosphorylation was significantly lower in tumors derived from TMG-treated mice than in tumors from control mice (Fig. 7b). Consistent with lower phosphorylation, O-GlcNAcylation of AMPK was significantly higher in tumors of TMG-treated mice (Fig. 7c). We examined the phosphorylation levels of AMPK, mTOR, and S6K, a downstream mTOR target, in tumors derived from control and TMG-treated mice (Fig. 7d). Consistent with in vitro data derived from LoVo cells (Figs. 3a,4e,f), TMG treatment showed augmented O-GlcNAcylation, which in turn led to AMPK inactivation and subsequent activation of the mTOR/S6K pathway (Fig. 7d). These results could be attributed to larger tumor volumes in TMG-treated mice than in untreated mice.

Discussion
Here, we showed that protein O-GlcNAcylation significantly enhanced the growth of colon cancer cells through AMPK inhibition and reciprocal activation of the mTOR/S6K pathway. The LKB1/AMPK pathway has increasingly gained attention as a connecting mechanism between diabetes and occurrence of cancers, including colon cancer. Although metformin, a biguanides class anti-type II diabetic drug, is also known as an AMPK activator, it has been reported to inhibit breast cancer growth by activating the AMPK pathway. These findings imply that AMPK plays a pivotal role in

Fig. 6. Transplanted LoVo cell-derived tumor formation was accelerated in Thiamet G (TMG)-treated BALB/c-nu/nu mice. LoVo cells were s.c. transplanted onto the backs of Balb/c-nu/nu nude mice (n = 6), with or without daily mouse treatment. Tumor sizes were measured weekly starting 1 week after transplantation. Images of the removed tumors are shown (top row, TMG treatment; bottom row, control). (b) The size of each tumor was measured weekly starting 1 week after transplantation. Tumor size was calculated using the formula (width × depth²)/2; width > depth). (c) Average tumor volume (n = 5) was calculated. Tumors with sizes that deviated from the mean ± SD were excluded in each group. **P < 0.01, Student's t-test.
regulating colon cancer physiology and can thus potentially act as a promising therapeutic target for the treatment of colon cancer. In this study, we showed that O-GlcNAcylation of AMPK, which has been reported to be increased under diabetic conditions, was observed in a colon cancer cell line, LoVo, and was found to be correlated with cancer cell growth (Figs 1, 2). Treatment with BML-275, a specific AMPK inhibitor (Fig. 3a, b) as well as AMPK silencing (Fig. 4b), enhanced the growth of LoVo cells in a dose-dependent manner. The mTOR activity in control LoVo cells was accelerated with
TMG treatment, whereas the activity in AMPK-silencing cells was not changed, indicating that the activation of the mTOR signaling pathway by TMG treatment is through AMPK, probably through the O-GlcNAcylation of AMPK (Fig. 4e,f). In contrast, treatment with AICAR, a specific AMPK activator, disrupted O-GlcNAcylation-dependent growth of LoVo cells (Fig. 5c). Although O-GlcNAcylation occurs on thousands of intracellular proteins and affects many signaling pathways that are critical for cell proliferation, our data strongly suggest that AMPK could be one of the major effectors for O-GlcNAcylation-mediated tumor growth. Importantly, suppression of AMPK by augmentation of O-GlcNAcylation or specific inhibition of AMPK led to the activation of the mTOR/S6K pathway. Consistent with the above findings, inverse results were obtained through the specific activation of AMPK. Conceivably, activation of the mTOR pathway by augmented AMPK O-GlcNAcylation is associated with increased cancer risk in diabetic patients.

O-linked N-acetylglucosamine signaling is strongly associated with cellular metabolism, given that the donor sugar for O-GlcNAcylation, UDP-GlcNAc, is synthesized from glucose, glutamine, and uridine triphosphate through the hexosamine biosynthetic pathway. Glucose influx is increased in diabetes and cancer and results in enhanced protein O-GlcNAcylation. To examine whether increased O-GlcNAcylation affects colon cancer development in vivo, we explanted LoVo cells onto TMG-treated nude mice. As shown in Figure 6, TMG-treated nude mice developed larger tumors compared to control nude mice. Consistent with in vitro results (Figs 2,4), AMPK O-GlcNAcylation was also increased, along with reciprocal activation of the mTOR pathway (Fig. 7). Taken together, augmented O-GlcNAcylation, which was observed in diabetic patients, would be directly associated with colon cancer development through inactivation of the antitumorigenic AMPK pathway.

AMP-activated kinase is a therapeutic target for metabolic syndrome and type 2 diabetes, and metformin activates AMPK. AMP-activated kinase is also activated by non-steroidal anti-inflammatory drugs and other anti-inflammatory agents, indicating that AMPK activation can potentially alleviate inflammation and pancreatic cancer. Once activated, AMPK counteracts the effects of mTOR, a molecular target in various cancers. In addition, p53, a tumor suppressor, and some transcription factors have been shown to be regulated by AMPK. Furthermore, O-GlcNAcylation, AMPK inhibitor, or siAMPK-mediated suppression of AMPK accelerated tumor growth (Figs 1–4), whereas AMPK activation suppressed tumor growth (Fig. 5), strongly suggesting that AMPK acts as a tumor suppressor.

The EGFR signaling pathway is upregulated in many primary and metastatic colon cancers. Anti-EGFR mAbs, such as cetuximab and panitumumab, play important roles in chemotherapy against colon cancer and act by inhibiting the activation of the EGFR and its downstream pathways, namely, the RAS-RAF-MAPK the PI3K-PTEN-AKT pathways. The response rate to anti-EGFR mAbs ranges from 10% to 20% in colon cancer patients, and several types of non-responders have been widely accepted, such as patients who harbor mutations in the KRAS, Braf, PIK3CA, and PTEN genes.

Recently, activation of the autophagy pathway through AMPK activation has been reported to be associated with chemoresistance to BRAF-V600E. Several studies have reported that autophagy induction in response to therapeutics could serve as a pro-survival role that contributes to chemoresistance. Although AMPK can activate autophagy under certain conditions, AMPK activation with AICAR treatment reduced the growth of LoVo cells (Fig. 5c) despite autophagy activation in vitro (Fig. S1A). Surprisingly, autophagy was not detected in tumors excised from LoVo cell-transplanted TMG-treated nude mice, in which AMPK was activated (Fig. S1B). Therefore, it is likely that the accelerated growth of LoVo cells observed both in vitro and in vivo was not due to autophagy activation.

Taken together, our findings suggested that augmented O-GlcNAcylation promotes the proliferation of LoVo cells through AMPK activation. We propose AMPK as a promising therapeutic target for the treatment of colon cancer, especially for patients who show resistance to chemotherapeutic drugs, such as those that target the EGFR pathway.

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Disclosure Statement

The authors have no conflict of interest.

Abbreviations

| Abbreviation | Description |
|--------------|-------------|
| ACC          | acetyl CoA carboxylase |
| AICAR        | 5-aminoimidazole-4-carboxamide ribonucleotide |
| AMPK         | AMP-activated kinase |
| EGFR         | epidermal growth factor receptor |
| G FAT        | glucose-6-phosphate/fructose aminotransferase |
| GlcNAc       | N-acetylgalactosamine |
| LKB1         | liver kinase B1 |
| OGA          | O-GlcNAcase |
| O-GlcNAc     | O-linked N-acetylgalactosamine |
| OGT          | O-GlcNAc transferase |
| S6K          | S6 kinase |
| TMG          | Thiamet G |
| UDP-GlcNAc   | uridine diphospho-N-acetylgalactosamine |

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Autophagy was activated in 5-aminoimidazole-4-carboxamide ribonucleotide (AICAR)-treated LoVo cells, but not in cells transplanted in Fig. S1.

Supporting Information

Original Article

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Supporting Information

Additional Supporting Information may be found online in the supporting information tab for this article:

Fig. S1. Autophagy was activated in 5-aminimidazole-4-carboxamide ribonucleotide (AICAR)-treated LoVo cells, but not in cells transplanted in Thimet G (TMG)-treated BALB/c-nu/nu mice.