Glaucocalyxin A Inhibits the Malignancies of Gastric Cancer Cells by Downregulating MDM2 and RNF6 via MiR-3658 and the SMG1-UPF mRNA Decay Pathway

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Gastric cancer (GC) ranks as the most common gastrointestinal cancer and is among the leading causes of cancer death worldwide. Glaucocalyxin A (GLA), an entkauranoid diterpene isolated from Rab-dosia japonica var., possesses various bioactivities. To date, the data on the effect of GLA on GC are still minimal, and the molecular mechanisms remain largely unknown. Herein, we found that GLA could significantly inhibit the proliferation, cell adhesion, and invasion of HGT-1, SNU-1, SNU-6, and NCI-N87 GC cells in a dose-dependent manner. GLA enhanced the apoptosis of the GC cells as evidenced by the increased caspase-3 activity and the elevated levels of cleaved caspase-3 and cleaved PARP in GC cells in the presence of GLA. We then showed that the downregulation of Murine Double Minute Clone 2 (MDM2) and Ring Finger Protein 6 (RNF6) by GLA was implicated in the GLA-induced inhibition of the GC cells. Furthermore, MDM2 and RNF6 were identified as the targets of miR-3658 that was downregulated in the GC cells and upregulated by GLA. Moreover, it was shown that miR-3658 was hypermethylated in the GC cells, and GLA could rescue the expression of miR-3658 via demethylation by abrogating EZH2-mediated epigenetic silencing. In addition to the miR-3658-MDM2/RNF6 regulatory axis, activation of the SMG1-UPF mRNA decay pathway contributed to the downregulation of MDM2 and RNF6 by GLA in the GC cells. The inhibitory effect of GLA on gastric cancer and the expression of MDM2 and RNF6 was also validated in in vivo study. Our findings suggest that has the therapeutic potential for GC by downregulating oncogenes via posttranscriptional regulation.

Keywords: gastric cancer, Glaucocalyxin A, malignancy, MDM2, RNF6, miR-3658, mRNA decay

INTRODUCTION

GC is the most common gastrointestinal cancer and a leading cause of cancer death worldwide (1). Despite the tremendous efforts and progress made in recent decades in treating GC, the overall survival rate for patients remains poor. Therefore, it is urgent to discover and develop new therapeutic approaches or agents for GC. Glaucocalyxin A (GLA), an entkauranoid diterpene...
isolated from Rabdosia japonica var., exhibits multiple pharmacological effects such as anticancer effect, induction of osteogenesis, suppression of inflammation, and prevention of myocardial infarction (2–9). MDM2 and RNF6 are extensively-studied oncproteins in various malignancies, including GC (10–13). Mdm2 is an oncogene mapped to chromosome 12q13-14, and the most critical function of MDM2 in cancers is to target the tumor suppressor P53 protein via posttranslational degradation or blockade of activity (10). MDM2 protein is so far the best studied oncoprotein in various malignancies, including GC (10–16, NCI-N87, normal gastric epithelial GES-1 cells, and Hs 738. St/Int cells were from ATCC, Virginia, USA. RPMI-1640 medium, Dulbecco’s Modified Eagle’s Medium (DMEM), fetal bovine serum (FBS), 100 U/ml penicillin and streptomycin, miR-3658 inhibitor, Lipofectamine LTX transfection reagent, 4% paraformaldehyde, SMG1 siRNA and control, Pierce BCA protein assay kit, and IL-1β and TNF-α ELISA kits were obtained from Thermo Fisher Scientific, Massachusetts, USA. Human HGT-1 cells, Matrigel, fibronectin, 5-Aza-CdR, miR-3658 mimic, lysis buffer, PVDF membrane, ECL detection reagent, EZ-Magna CHIP kit, actinomycin D, and N-Methyl-N-nitrosourea (MNU) were obtained from MilliporeSigma, Massachusetts, USA. Matrigel-coated Transwell inserts (8 μm: pore size) and cell culture plates were obtained from Corning, New York, USA. Primary antibodies used in the present study include rabbit anti-cleaved caspase-3 antibody, rabbit anti-cleaved PARP antibody, Rabbit anti-β-integrin antibody (Cell Signaling Technology, Massachusetts, USA), rabbit anti-MDM2 antibody, rabbit anti-RNF6 antibody, rabbit anti-GAPDH antibody, rabbit anti-EZH2 antibody, and rabbit anti-H3K27me3 antibody (Abcam, Massachusetts, USA). Goat-anti rabbit IgG, rabbit normal IgG, and WST-8/CCK8 kit were obtained from Abcam. Caspase-3 colorimetric assay kit was obtained from BioVision (California, USA). Human MDM2 and RNF6 lentivirus particles were from Creative Biogene, New York, USA. Trizol and SuperScript IV first-stand Synthesis System were from Invitrogen, Massachusetts, USA. TaqMan MicroRNA Reverse Transcription Kit, U6 snRNA, and PowerUp SYBR Green Master Mix PCR kit were from Applied Biosystems, Massachusetts, USA. Myeloperoxidase (MPO) assay kit was the product from HyCult Biotechnology, Uden, The Netherlands.

MicroRNAs (miRNAs) are small single-stranded non-coding RNA molecules containing 19-23 nucleotides that have emerged as posttranscriptional regulators of the target genes (16, 17). MiRNAs target downstream genes by interacting with the specific binding site in the three prime untranslated regions (3′-UTR) in mRNA. MiRNAs have been documented to be involved in GC initiation and progression, and the pattern of deregulated miRNAs represents the features related to diagnosis, prognosis, and therapeutic resistance in GC (16–19). Recent data have revealed that miR-3658 acts as tumor suppressor miRNA in colorectal cancer by targeting octamer-binding transcription factor 4 (Oct4) (17). In addition to deregulated mRNA silencing by miRNAs that occurred at the posttranscriptional level in cancer cells, the impairment of the nonsense-mediated mRNA decay (NMD) mechanism plays a critical role in driving the onset and progression of cancer via posttranscriptional regulation (20).

To date, the data on the effect of GLA on GC are still minimal, with molecular mechanisms yet to be extensively revealed. A recent study conducted by Zhou et al. showed that GLA could prevent hypoxia-induced epithelial-mesenchymal transition in MGC-803 GC cells via the PI3K/Akt signaling pathway (21). Herein, we found that GLA can inhibit the malignant behaviors of GC cells by downregulating the expression of MDM2 and RNF6 via epigenetic abrogation of EZH2-mediated miR-3658 silencing. Moreover, GLA can also suppress the expression of MDM2 and RNF6 by enhancing mRNA decay through activation of the SMG1-UPF mechanism.

**MATERIALS AND METHODS**

**Materials**

GLA (Figure 1A) is a natural compound obtained from BOC Sciences, New York, USA, with white powder appearance and >98% purity. The molecular formula of GLA is C20H28O4, and the molecular weight of 332.43 Da. GLA was dissolved in dimethyl sulfoxide (DMSO). Human SNU-1, SNU-16, NCI-N87, normal gastric epithelialGES-1 cells, and Hs 738.St/Int cells were from ATCC, Virginia, USA. RPMI-1640 medium, Dulbecco’s Modified Eagle’s Medium (DMEM), fetal bovine serum (FBS), 100 U/ml penicillin and streptomycin, miR-3658 inhibitor, Lipofectamine LTX transfection reagent, 4% paraformaldehyde, SMG1 siRNA and control, Pierce BCA protein assay kit, and IL-1β and TNF-α ELISA kits were obtained from Thermo Fisher Scientific, Massachusetts, USA. Human HGT-1 cells, Matrigel, fibronectin, 5-Aza-CdR, miR-3658 mimic, lysis buffer, PVDF membrane, ECL detection reagent, EZ-Magna CHIP kit, actinomycin D, and N-Methyl-N-nitrosourea (MNU) were obtained from MilliporeSigma, Massachusetts, USA. Matrigel-coated Transwell inserts (8 μm: pore size) and cell culture plates were obtained from Corning, New York, USA. Primary antibodies used in the present study include rabbit anti-cleaved caspase-3 antibody, rabbit anti-cleaved PARP antibody, Rabbit anti-β-integrin antibody (Cell Signaling Technology, Massachusetts, USA), rabbit anti-MDM2 antibody, rabbit anti-RNF6 antibody, rabbit anti-GAPDH antibody, rabbit anti-EZH2 antibody, and rabbit anti-H3K27me3 antibody (Abcam, Massachusetts, USA). Goat-anti rabbit IgG, rabbit normal IgG, and WST-8/CCK8 kit were obtained from Abcam. Caspase-3 colorimetric assay kit was obtained from BioVision (California, USA). Human MDM2 and RNF6 lentivirus particles were from Creative Biogene, New York, USA. Trizol and SuperScript IV first-stand Synthesis System were from Invitrogen, Massachusetts, USA. TaqMan MicroRNA Reverse Transcription Kit, U6 snRNA, and PowerUp SYBR Green Master Mix PCR kit were from Applied Biosystems, Massachusetts, USA. Myeloperoxidase (MPO) assay kit was the product from HyCult Biotechnology, Uden, The Netherlands.

**Cell Culture**

Human GC cell lines HGT-1, SNU-1, SNU-16, and NCI-N87 were maintained in RPMI-1640 medium supplemented with 10% FBS and 100 U/ml penicillin and streptomycin at 37°C in a humidified incubator with 5% CO2. Normal human gastric epithelialGES-1 cells and Human Hs 738.St/Int cells were cultured in DMEM supplemented with 10% and FBS and 100 U/ml antibiotics.

**Cell Counting Kit 8 (WST-8/CCK8) Assay**

Cell viability was examined using the WST-8/CCK8 kit according to the manufacturer’s instruction. Briefly, the GC cells were seeded at 5×10^3 cells/well in 96-well culture plates. After 3 hours of recovery, the cells were treated with serial concentrations of GLA for 24 h, 48 h, or 72 h. Subsequently, WST-8/CCK8 solution was added to each well and incubated for 4 hours at 37°C. WST-8/CCK8 tetrazolium salt was reduced by cellular dehydrogenases to a formazan product soluble in culture medium. The amount of formazan produced was proportional to the number of living cells and was measured by absorbance at 460 nm using a BioTek microplate reader.

**Western Blot Analysis**

GC cells with different treatment were lysed in lysis Buffer. Gastric tumor tissues from mice were lysed with lysis buffer
using a Miltenyi gentleMACS Dissociator. Protein concentrations were determined using BCA protein assay kit. 30-50 µg of total protein was subjected to 4-10% SDS-PAGE gels, transferred onto PVDF membranes, and blocked in 5% non-fat milk for 1 h at room temperature. The membranes were then incubated with specific primary antibodies overnight at 4°C. The membranes were subsequently incubated with secondary antibodies conjugated with horseradish peroxidase for 1 h at room temperature. The immunoblots were subsequently visualized using ECL detection reagents. The densitometry of the bands was analyzed using Image J.

**Cell Adhesion Assay**
For the cell adhesion assay, a 96-well plate was pre-coated with 300 µg/mL of Matrigel and 10 µg/mL fibronectin at 4°C overnight and blocked with 1% (w/v) albumin. GC cells were centrifuged, resuspended in complete medium, and seeded on the 96-well plate at a density of 1 × 104 per well. After adhering for 90 min, the cells in the wells were washed with phosphate-buffered saline (PBS), fixed with 4% paraformaldehyde for 10 min, and stained with 0.5% crystal violet for 10 min. The stained attached cells were lysed using 30% glacial acetic acid for 10 min. OD values at 570 nm were then measured.

**Cell Invasion Assay**
For the cell invasion assay, 2 × 10^4 GC cells (100 µl per well) were seeded in a Matrigel-coated Transwell insert (8 µm pore size) in a 24-well culture plate and maintained in serum-free medium. After incubation for 12 h at 37°C, cells on the top surface of the insert were removed with a cotton swab, and cells that invaded to the bottom surface of the insert were fixed with 4% paraformaldehyde for 30 minutes, stained Transwell inserts in DAPI for 10 minutes then subjected to microscopic inspection.

**Transfection**
HGT and SNU-1 cells reaching 70% confluence in a six-well culture were transfected with 10 pM of miR-3658 inhibitor, mimic, or the negative controls, and 2 µg of PGL3 luciferase reporter vectors containing either wild-type 3’ UTR or muted 3’UTR of MDM2 or RNF6 using Lipofectamine LTX reagent according to the manufacturer’s instructions. The culture medium was replaced with complete medium at 6 h after transfection. The cells were harvested for subsequent assays after 48 h transfection.

**RNA Extraction and Reverse Transcription Quantity (RT- PCR)**
Total RNA was extracted from the GC cells and gastric tumor tissues from mice using Trizol. 5 ng of purified RNA were retrotranscribed using TaqMan MicroRNA Reverse Transcription Kit. The levels of miR-3658 were measured using Real-Time quantitative PCR according to the TaqMan MicroRNA Assay protocol, and U6 snRNA served as an internal control. The PCR reaction was carried out in a 96-well PCR microplate at 95°C for 10 min followed by 40 cycles of 95°C for 15 s and 60°C for 1 min. The primers for miR-3658 are forward 5’-GTTGGGGGTITTA AGA AAA A CAC CAT-3’ and reverse 5’-GTGCA GG GTCCGAGGT-3’. To analyze the mRNA levels of MDM2 and RNF6, 1 µg/ml of total RNA was reversely transcribed into cDNA using a SuperScript IV first-st in synthesis System. Real-Time PCR was then performed using a PowerUp SYBR Green Master Mix PCR kit. The PCR reactions were performed at 95°C for 15 min, followed by 40 cycles of 95°C for 30 s and 58°C for 30 s. The primers for human MDM2, RNF6, and the internal control 18S were as follows: MDM2 (human) forward 5’-CACCTCAGAGTCCAGCTT-3’ and reverse 5’- CGCCAAACAAAT CCTCTAGA-3; MDM2 (mouse) forward 5’-ATGAGGTCATCGGGTACAGT-3’ and reverse primer 5’-CACATCCAAG CCTTCT TCTG-3; RNF6 (human) forward 5’-AGAAAGATGCCAGCAAGGCGGACACTCAGCATC-3; 18S (human) forward 5’- AGCCAGCGCGGTAT ACCAGCTT-3’ and reverse 5’- CGGGCACACTCAGGCTAGCATC-3; 18S (mouse) forward 5’- AGTCCCTGCA CTTTGTACA-3’ and reverse 5’- CGATCGAGGGGCTCACA-3’. Primers for mouse RNF6 were the products of Ambion, Cambridge, USA. The PCR results were calculated using the 2-ΔΔCT method.

**Chromatin Immunoprecipitation (CHIP) Assay**
CHIP assays were performed using an EZ-Magna CHIP kit according to the manufacturer’s instructions. After being fixed with 1% formaldehyde, nucleoprotein complexes were sheared to 200–500 bp with sonication and then immunoprecipitated with anti-EZH2 or anti-H3K27me3 antibodies overnight at 4°C. The enrichment of the DNA fragment of miR-3658 upstream coding region in the immunoprecipitated complexes was then detected using PCR.

**Statistical Analysis**
All the data are shown as mean ± standard deviation (S.D). Statistical significance was determined by unpaired Student’s two-tailed t-test or one-way ANOVA analysis, with p < 0.05 was considered to be significant (*p < 0.05, **p < 0.01, ***p < 0.001). Incidences in the in vivo study were assessed by Chi-square test, followed by pairwise analysis. Each experiment was performed a minimum of three times.

**Animal Experiments**
Animal experiments were performed according to the guidelines provided by the experimental animal center of Hangzhou Huqingyu Hall Pharmaceutical Co., Ltd, Hangzhou, China, and approved by the Animal Care and Use Committee of Hangzhou Huqingyu Hall Pharmaceutical Co., Ltd. 24 six-week-old C57BL/6 mice were divided into 4 groups with 6 mice in each group: vehicle control, H. pylori + MNU, GLA only, H. pylori + MNU+ GLA. H. pylori and MNU were used to induce gastric cancer in the mice. The mice were inoculated with a total dose of 1 × 108 colony-forming units of H. pylori (Sydney strain 1) per mouse in 200 µl of trypticase broth by oral gavage 6 times for 2 weeks before MNU
administration. The mice were given MNU freshly dissolved in distilled drinking water at the concentration of 240 ppm in a light-shielded bottle on alternate weeks for 10 weeks. The mice were then kept being housed with free access to food and water, waiting for the development of gastric tumors for 18 weeks. 10 mg/kg GLA were administrated by oral gavage twice a week for 12 weeks after 6 weeks post MNU treatment. The mice were sacrificed by CO₂ asphyxiation and stomach tissue was taken for the subsequent process at 18 weeks after completion of MNU induction. To examine the pro-inflammatory status in the gastric tissues by measuring the levels of mucosal myeloperoxidase (MPO), a marker of inflammation, and pro-inflammatory cytokines, 20 mg of scraped gastric mucosa from mice was homogenized in 500 µL of lysis buffer on ice and centrifuged at 13,000 rpm for 15 minutes. The supernatant was collected for MPO assay and the ELISA for IL-1β and TNF-α according to the manufacturers’ instructions.

RESULTS

GLA Inhibits the Malignant Behavior of GC Cells

We sought to investigate the effect of GLA on the malignant behaviors of GC cells. HGT-1, SNU-1, SNU-6, and NCI-N87 GC cell lines were treated with 0, 0.1, 0.5, 2.5, 5, 10, 20, and 40 µM of GLA for 24 h, 48 h, or 72h. We found that GLA exerted an antiproliferative effect on all four GC cell lines in a dose-dependent fashion (Figure 1B) but didn’t significantly affect the cell viability of normal human gastric epithelial GES-1 cells and human Hs 738.St/Int cells at the concentrations from 0.1 µM to 10 µM (Supplementary Figure 1). Thus, 10 µM was set as the highest concentration of GLA in our mechanistic assays. We examined the effect of GLA on the apoptotic status of the GC cells. The caspase-3 activity and the expression of cleaved caspase-3 and cleaved PARP were significantly increased by GLA treatment (Figure 1C). Collectively, the GC cells exhibited a greatly reduced ability to proliferate, accompanied by dramatically improved apoptosis in the presence of GLA. Cell adhesion is among the essential steps for cancer cells to invade. We further examined the impact of GLA on the cell adhesion process in the GC cells. The expression of β1-integrin, a critical protein associated with cell adhesion, was downregulated in the presence of GLA (Figure 1D). As shown in Figure 1E, GLA significantly blocked the adhesion of GC cells to fibronectin and Matrigel in a dose-dependent manner. Consistently, the invasive capability of the GC cells was also hampered by GLA (Figure 1F).

The Inhibitory Effect of GLA on GC Cells Is Mediated by Downregulating MDM2 and RNF6

Mdm2 and RNF6 are classified as oncogenes in various types of cancer, including gastric cancer. Our previous screening of the genes that are regulated by GLA in gastric cancer cells revealed that Mdm2 and RNF6 were the most downregulated genes, among others, in response to GLA treatment in gastric cancer cells. Thus, we selected Mdm2 and RNF6 in the present study to explore the molecular mechanism underlying the inhibitory effect of GLA in gastric cancer cells. We explored the role played by GLA-MDM2/RNF6 in controlling the malignant behavior of GC cells. As shown in Figure 2A, the expression of MDM2 and RNF6 were dramatically upregulated in all the four GC cells at mRNA and protein levels, and GLA significantly inhibited the expression of MDM2 and RNF6. We then explored the relationship between MDM2 and RNF6 expression and GLA effect. Given the consistent results in the four gastric cell lines in the presence of GLA, we then chose and focused on two cell lines (HGT-1 and SNU-1) among them for further mechanistic studies. We found that restoration of MDM2 and RNF6 by ectopic overexpression could significantly abrogate the inhibitory effect of GLA on proliferation (Figure 2B), adhesion (Figure 2C), and invasion (Figure 2D) of the HGT-1 and SNU-1 cells.

MDM2 and RNF6 Are the Targets of MiR-3658 That Is Downregulated in GC Cells and Upregulated by GLA Treatment

It has been reported that microRNAs play important roles in regulating the expression of oncogenes. To confirm whether microRNAs are involved in inhibiting the expression of MDM2 and RNF6 by GLA, TargetScan and miRDB prediction algorithms were utilized to predict the microRNAs targeting MDM2 and RNF6. MDM2 and RNF6 were both predicted targets of miR-3658 (Figure 3A). Compared to the levels of miR-3658 expressed in GES-1 normal gastric epithelial cells and Hs 738.St/Int cells, miR-3658 was significantly downregulated in GC cells (Figure 3B). MiR-3658 was upregulated in GC cells in response to GLA treatment (Figure 3B). MiR-3658 mimic could dramatically downregulate the expression of MDM2 and RNF6 at mRNA and protein levels in the GC cells (Figure 3C). Thus, we deduced that miR-3658-MDM2/RNF6 might be implicated in the inhibition of GC cells by GLA. It was found that overexpression of miR-3658 by transfection of its mimic enhanced the effect of GLA, whereas the inhibitor of miR-3658 mitigated the effect of GLA in GC cells (Figure 3D). To further validate whether miR-3658 directly targets MDM2 and RNF6, a PGL3 reporter vector containing either wild-type 3′ UTR or mutated 3′UTR of MDM2 or RNF6 was performed. As shown in Figure 3E, the fluorescent reporter intensity was decreased in the cells with transfection of wild-type 3′UTR of MDM2 or RNF6 but remained unaffected in the cells transfected with mutant 3′UTRs in the GC cells transfected with miR-3658 mimic. These findings suggest that miR-3658 silences MDM2 and RNF6 by directly targeting.

MiR-3658 Is Hypermethylated in GC Cells, and GLA Can Restore MiR-3658 by Abrogating EZH2-Mediated Epigenetic Silencing

Since miR-3658 could act as a tumor suppressor microRNA and is downregulated in gastric cancer, we sought to explore the mechanism underlying the restoration of miR-3658 by GLA in
GC cells. Cancer-related aberrant methylation is commonly observed in miRNA genes, and the hypermethylation leads to the downregulation of the expression of tumor-suppressive miRNAs in cancer cells (22–25). To investigate whether DNA methylation is involved in the downregulation of miR-3658 in GC cells, we treated GC cells with 5-Aza-CdR, an S-phase-specific demethylating agent, to induce demethylation. As shown in Figure 4A, the levels of miR-3658 were significantly elevated in the cells treated with 5-Aza-CdR or GLA, suggesting that miR-3658 is hypermethylated in GC cells, and GLA may...
increase miR-3658 expression via demethylation. EZH2 is the enzymatic subunit of Polycomb Repressive Complex 2 (PRC2) that functions to silence the transcription of target genes by hyper-methylating histone H3 lysine 27. As shown in Figure 4B, EZH2 could directly bind to the upstream region of miR-3658, and the occupancy of histone methylation H3K27me3 was also detected in the CHIP precipitate. In contrast, GLA treatment blocked the association of EZH2 and H3K27me3 with the upstream region of miR-3658. These findings indicate that GLA could restore miR-3658 expression by abrogating EZH2-mediated epigenetic silencing through hypermethylation.

GLA Activated the SMG1-UPF mRNA Decay Mechanism, Which Mediated mRNA Decay of MDM2 and RNF6

SMG1-UPF mechanism functions to downregulate the expression of various essential factors that facilitate tumorigenesis by inducing their mRNA decay (26, 27). Our analysis of pathway-based sequencing data integration and visualization conducted and generated using RNA from GC cells treated with GLA performed by Arraystar Inc. showed that GLA could activate the SMG1-UPF3B pathway by phosphorylation (Figure 5A). To investigate the effect of GLA on the mRNA stability of MDM2 and RNF6 in gastric tumor cells, we...
used actinomycin D, a transcription inhibitor widely used to inhibit the synthesis of new mRNA in the assessment of mRNA decay. We found that GLA significantly enhanced mRNA decay of both MDM2 and RNF6 in HGT-1 and SNU-1 cells (Figure 5B). Given that the generation of new mRNAs of MDM2 and RNF6 was blocked by actinomycin D, the decrease in the levels of the mRNAs in GLA-treated cells reflected an accelerated mRNA degradation compared to that in control cells in the absence of GLA. Furthermore, silencing the expression of SMG1 by siRNA led to the reduction in the mRNA decay enhancement triggered by GLA treatment (Figure 5B). A significant difference between control siRNA and SMG1 siRNA control was not observed in our study, which is consistent with the fact that the activation or function of SMG1-UPF3B by phosphorylation is closely involved in GLA-accelerated decay of MDM2 and RNF6 mRNAs and the SMG1-UPF3B pathway is inactivated in gastric cancer cells in the absence of GLA. Thus, our
results indicate that GLA could inhibit the expression of MDM2 and RNF6 in GC cells via not only upregulation of miR-3658 but also activation of the SMG1-UPF mRNA decay mechanism.

GLA Alleviates Gastric Cancer and Downregulates the Expression of MDM2 and RNF6 In Vivo

We then sought to determine whether GLA could inhibit gastric cancer induced by a combination of *H. pylori* and MNU in mice. The effective number of mice and the incidences of gastric tumors are shown in Table 1 and Figure 6A. The incidences and tumor weight were significantly decreased in GLA treated group. No gastric tumors were formed in the negative control group and GLA only group. The levels of mucosal MPO and the expression of pro-inflammatory cytokines, IL-1β and TNF-α in gastric mucosal were examined to evaluate the pro-inflammatory status. As shown in Figure 6B, the gastric mucosal MPO levels were significantly increased in *H. pylori* and MNU group compared to that of the negative control group and GLA only group, which was reduced by GLA treatment. The changes in the levels of IL-1β and TNF-α were consistent with the MPO result. Furthermore, the expression of MDM2 and RNF6 was increased in the tumor tissues in *H. pylori* and MNU group, and the effect was abrogated by GLA treatment (Figure 6C). There was no detectable MDM2 and RNF6 expression in the control group and GLA only group (data not shown).

**DISCUSSION**

Despite the advancements in the treatment of GC, the 5-year overall survival of GC remains low. The discovery and development of new molecular compounds for treating GC and exploring the underlying molecular mechanisms are essential for developing novel therapeutic regimens that improve GC prognosis. In the present study, we revealed the GLA-MDM2/RNF6 inhibitory mechanism underlying the suppressive effect of GLA on GC. The studies on the role of GLA and the mechanism in inhibiting GC remain minimal. In the present study, we found that GLA significantly inhibited the malignant behavior of GC cells, and the inhibitory effect of GLA on GC cells was mediated by downregulating MDM2 and RNF6. The downregulation of MDM2 and RNF6 by GLA was at least by restoring miR-3658 and inducing SMG1-UPF-MDM2/RNF6 mRNA decay machinery.

MDM2 inhibits the transactivation of tumor suppressor P53. Being a P53-specific E3 ubiquitin ligase, MDM2 is a principal factor in P53 destruction (28). MDM2 has proved to play an essential role in cancer progression by inducing cell proliferation, invasion, and therapeutic resistance and inhibiting the apoptotic process, serving as a useful predictive marker for poor prognosis in various human cancers such as GC, breast cancer, hepatocellular carcinoma, and lung cancer (11, 29–31). MDM2 expression parameters are closely correlated with *H. pylori* infection and higher stages GC (29). We found that GLA treatment dramatically inhibited the expression of MDM2, suggesting that the reduction of MDM2 levels in the presence of GLA is involved in hampering the malignant

| Table 1: Incidence of gastric tumors in the in vivo study. |
|-----------------------------------------------|
| **Group** | **Effective number** | **Incidence (%)** | **Total tumor** |
| Negative control | 0 | 0 | |
| *H. pylori* + MNU | 6 | 100 | |
| GLA | 0 | 0 | |
| *H. pylori* + MNU + GLA | 3 | 50* | |

Incidence were assessed by Chi-square test, followed by pairwise analysis.

*P<0.05 vs H. pylori + MNU group.
behavior of GC cells. RNF6, a RING-type E3 ubiquitin ligase, has been found to play an oncogenic role in cancer (15, 32, 33). RNF6 belongs to the RNF family, which induces proteasomal degradation of its downstream proteins by ubiquitination and tagging (34–36). Similar to MDM2, RNF6 is also downregulated by GLA in GC cells. Given the E3 ubiquitin ligase property of both MDM2 and RNF6 in cancer metabolism (37), GLA might exert its antitumor effect on GC cells by blocking MDM2 and RNF6-mediated deregulated ubiquitination processes.

We then sought to investigate how GLA downregulates MDM2 and RNF6 expression in GC cells. The roles played by miRNAs, either as a tumor suppressor or as tumor enhancer, in regulating oncogene expression in cancers have been widely reported (38–40). MiR-3658 was then introduced as a bona fide upstream negative regulator for both MDM2 and RNF6 by prediction algorithms, which was validated by the inverse correlation in expression between miR-3658 and MDM2/RNF6 and the direct interaction in the in vitro assay. Hosseini et al. have shown that miR-3658 acts as a tumor suppressor in colorectal cancer cells. MiR-3658 was downregulated and the forced overexpression of miR-3658 in colorectal cancer cells brought about proliferation inhibition, cell cycle arrest, and retarded mobility (17). Consistent with their findings, we found that miR-3658 was downregulated in GCs and functioned to correct the malignant behaviors of GC cells. Furthermore, the levels of miR-3658 were elevated in response to GLA treatment in GC cells, suggesting that restoration of the function of miR-3658-MDM2/RNF6 axis by GLA contributed to its antitumor effect on GC cells. These findings indicate that miR-3658 plays a tumor-suppressor role and participates in mediating the downregulation of MDM2/RNF6 by GLA. Given the critical role of miR-3658 in the GLA-MDM2/RNF6 mechanism, we then sought to examine how GLA affected the expression of miR-3658. Studies have demonstrated that DNA methylation plays a crucial role in silencing tumor-suppressive miRNAs in tumors (41–43). To determine whether suppression of DNA hypermethylation was involved in the upregulation of miR-3658 by GLA in the GC cells, 5-Aza-CdR, a widely used demethylation inducer, was utilized to treat GC cells. The levels of miR-3658 were significantly increased in the presence of 5-Aza-CdR. Thus, we deduced that suppression of miR-3658 in GC cells was mediated by hypermethylation that was then validated by anti-EZH2 and anti-H3K27me3 CHIP assays. These results suggest that GLA can “switch on” the transcription of miR-3658 via demethylation. Further studies are needed to examine whether GLA could play a general role of demethylation inducer in cancer cells by epigenetically “turning on” the transcription of hypermethylated tumor suppressor genes.

In addition to the miR-3658 pathway, our pathway analysis data in GC cells treated with GLA showed that GLA could activate the SMG1-UPF pathway that functions to downregulate the expression of various essential factors facilitating tumorigenesis by inducing mRNA decay (26, 27). The SMG1-UPF pathway is a core part of NMD, a quality-control mechanism that rapidly degrades aberrant mRNAs (22, 23). Impairment of NMD-mediated mRNA degradation against oncogenesis is involved in various pathophysiological processes of cancer (44–46). Tumor suppressor genes and factors are prone to induce NMD that can downregulate the expression of many
oncogenes and factors implicated in signaling pathways that lead to cancer (44, 47). We found that GLA enhanced the mRNA decay of MDM2 and RNF6 via SMG1 pathway, thus to further decrease their expression at the posttranscriptional level. Based on the findings in our study, the SMG1-UPF mRNA decay pathway and miR-3658 silencing are two independent posttranscriptional regulatory mechanisms underlying the inhibitory effect of GLA on gastric cancer cells, and they have an additive effect of enabling the desired anti-cancer effect of GLA. However, more studies are needed to further validate the relationship between miR-3658 demethylation and SMG1-UPF mRNA decay induced by GLA, such as functional analysis, p53 degradation or protein ubiquitination.

Our results reveal that the activation of both miR-3658-mediated RNA silencing and the SMG1 mRNA decay mechanism that hampers the expression of oncogenes, Mdm2 and RNF6, contributes to the inhibitory effect of GLA on GC cells. Being a natural compound from Rab-dosia japonica var, GLA could greatly benefit treating GC by targeting the expression of oncogenes represented by MDM2 and RNF6 in a posttranscriptional dependent manner.

DATA AVAILABILITY STATEMENT

The original contributions presented in the study are included in the article/Supplementary Material. Further inquiries can be directed to the corresponding author.

ETHICS STATEMENT

Animal experiments were performed according to the guidelines provided by the experimental animal center of Hangzhou Huqingyu Hall Pharmaceutical Co., Ltd, Hangzhou, China, and approved by the Animal Care and Use Committee of Hangzhou Huqingyu Hall Pharmaceutical Co., Ltd.

AUTHOR CONTRIBUTIONS

YL, PC, DQ, and LC conceived the project. DQ, YL, PC, and LC performed the study. PC, YL, DQ, and LC wrote the manuscript. PC supervised the project. All authors contributed to the article and approved the submitted version.

SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fonc.2022.871169/full#supplementary-material

Supplementary Figure 1 | Cell viability of GES-1 and Hs 738.St/Int cells treated with 0, 0.1, 0.5, 2.5, 5, and 10 µM of GLA at 24 h, 48 h, or 72h (% of control). The data represented the mean ± SD (n=3).
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